



**Rawalpindi Medical University University Residency Program 2021 MS OPHTHALMOLOGY** 

# C U L U M



# **RAWALPINDI MEDICAL UNIVERSITY**

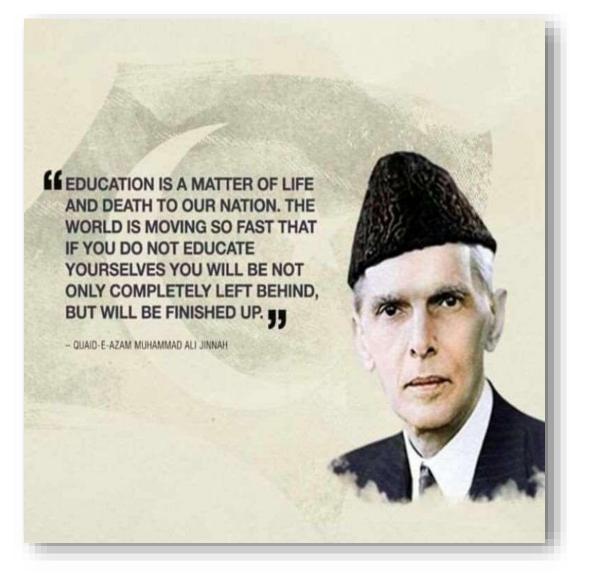
# **CURRICULUM & REGULATIONS**

# FOR

# **MS OPHTHALMOLOGY**

# (Revised February, 2025)









# PREFACE

The horizons of Medical Education are widening & there has been a steady rise of global interest in Post Graduate Medical Education, an increased awareness of the necessity for experience in education skills for all healthcare professionals and the need for some formal recognition of postgraduate training in Internal Medicine.

We are seeing a rise in the uptake of places on postgraduate courses in medical education, more frequent issues of medical education journals and the further development of e-journals and other new online resources. There is therefore a need to provide active support in Post Graduate Medical Education for a larger, national group of colleagues in all specialties and at all stages of their personal professional development. If we were to formulate a statement of intent to explain the purpose of this log book, we might simply say that our aim is to help clinical colleagues to teach and to help students to learn in a better and advanced way. This book is a state-of-the-art log book with representation of all activities of the MS Ophthalmology program at RMU.A summary of the curriculum is incorporated in the logbook for convenience of supervisors and residents. MD curriculum is based on six Core Competencies of ACGME (Accreditation Council for Graduate Medical Education) including

Patient Care, Medical Knowledge, System Based Practice, Practice Based Learning, Professionalism, Interpersonal and Communication Skills. A perfect monitoring system of a training program including monitoring of teaching and learning strategies, assessment and Research Activities cannot be denied so we at RMU have incorporated evaluation by Quality

Assurance Cell and its comments in the logbook in addition to evaluation by University Training Monitoring Cell (URTMC). Reflection of the supervisor in each and every section of the logbook has been made sure to ensure transparency in the training program. The mission of Rawalpindi Medical University is to improve the health of the communities and we serve through education, biomedical research and health care. As an integral part of this mission, importance of research culture and establishment of a comprehensive research structure and research curriculum for the residents has been formulated and a separate journal for research publications of residents is available.

Prof. Muhammad Umar (Sitara-e-Imtiaz) (MBBS, MCPS, FCPS, FACG, FRCP (Lon), FRCP (Glasg), AGAF) Vice Chancellor Rawalpindi Medical University & Allied Hospitals



# **RMU Motto**



# **Mission and Vision of Rawalpindi Medical University**



Vision

To impart evidence-based research-oriented health professional education in order to provide best possible patient care and inculcate the values of mutual respect, ethical practice of healthcare and social accountability.

# Mission

Highly recognized and accredited center of excellence in Medical Education, using evidence-based training techniques for development of highly competent health professionals, who are lifelong experiential learner and are socially accountable



## CONTRIBUTIONS

SR. NO.	NAME & DESIGNATION		CONTRIBUTIONS IN FORMULATION & REVISION OF CURRICULUM OF OPHTHALMOLOGY
1.		<b>Prof. Dr. Fuad Ahmad</b> <b>Khan Niazi</b> Head Of ophthalmology Department RMU & Allied Hospitals, Rawalpindi	Over all synthesis, structuring & over all write up of Curriculum of MS Ophthalmology, Log Book of MS Ophthalmology & Allied and also Log Book for MS Ophthalmology rotations under guidance of Prof. Muhammad Umar Vice Chancellor, Rawalpindi Medical University, Rawalpindi. Also, Proof reading & synthesis of final print version of Log Books of MS Ophthalmology and Rotation Log Book
2.		<b>Dr. Ambreen Gull</b> Associate Professor Ophthalmology Department Benazir Bhutto Hospital, Rawalpindi	Assistance of Professor Dr. Fuad Ahmad Khan Niazi in revising & updating the MS & MD curriculum, adding frameworks, Calgary model, EPA's, formulating the log books and computer work under his direct guidance & supervision.
3.		<b>Dr. Fatima Sidra</b> Senior Registrar Ophthalmology Department Holy Family Hospital, Rawalpindi	Assistance of Professor Dr. Fuad Ahmad Khan Niazi in revising & updating the MS & MD curriculum under his direct guidance & supervision.



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4.	<b>Dr. Saira Bano Satti</b> Senior Registrar Ophthalmology Department Holy Family Hospital, Rawalpindi	Assistance of Professor Dr. Fuad Ahmad Khan Niazi in formulating the log books & computer work under his direct guidance & supervision.
5.	<b>Dr. Bilal Humayun</b> <b>Mirza</b> Ex-Senior Registrar Ophthalmology Department DHQ Hospital, Rawalpindi	Assistance of Professor Dr. Fuad Ahmad Khan Niazi in formulating the log books & computer work under his direct guidance & supervision.



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# **Table of Contents**





SECTION I: PREAMBLE	12
Introduction to curriculum	14
ACGME Competency Model	14
Overview of Program	
Admission criteria	15
Training sites	15
Program Leadership	16
Rules and Regulations	16
Statutes	
Program Structure	
Framework of the Program	
Training Pathway MS ophthalmology	19
Clinical and Surgical Training Resources	
RMU Postgraduate Competency Model	
RMU Postgraduate Competency Framework of Ophthalmology	22
Aim of The Program	
Outcomes of the Program	23
SECTION II: COURSE CONTENT	
Content Overview	27
Bloom's Taxonomy Domains	27
Credit Hours of MS Ophthalmology Program	
Core Content Clinical Ophthalmology	
Core Content Optics and Refraction	
SECTION III: SPECIFIC LEARNING OUTCOMES	
Specific Learning Outcomes Optics and Refraction	
Specific Learning Outcomes Clinical Ophthalmology	45
SECTION IV: TEACHING AND LEARNING STRATEGIES	
Overview	
Outline of Different Teaching and Learning Strategies	
Teaching Schedule	
SECTION V: WORKSHOPS	



Framework of Workshops in MS Ophthalmology Program	179
Learning Objectives of Workshops	180
SECTION VI: ROTATIONS (ELECTIVES)	191
Overview	192
Rotation Framework	
Rotation planner	194
Minimum Log Book Entries for Rotations	194
SECTION VII: RESEARCH	
Resident research pathway	197
Outline of research curriculum	
Model of Research at Rawalpindi Medical University	
Research milestones	200
SECTION VIII: TRAINING MILESTONES	202
Charting the Road to Competence: Developmental Milestones	
SECTION IX: ASSESSMENT AND EVALUATION	
Framework of Assessment for MS Ophthalmology Residency Program	224
Assessment Hours Allocation	
Assessment framework based on Miller's Pyramid and ACGME core competend MS Ophthalmology program	
Formative Assessment	
Summative Assessment	
Table Of Specifications	232
Research Assessment	
SECTION X: ENTRUSTABLE PROFESSIONAL ACTIVITY (EPA)	
Overview	
EPA of MS Ophthalmology Training Program	
SECTION XI: LOGBOOK	269
Introduction to Logbook	270
Logbook Entry Map	271
SECTION XII: PORTFOLIO	272
SECTION XIII: REFERENCES	277
Teaching Methods	278
Assessment methods	278
Milestones	
SECTION XIV: APPENDICES	

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# **SECTION I: PREAMBLE**





# **Mission of Training Program**

To cultivate dedication in trainees, develop expertise in diagnostic and therapeutic ophthalmology, advancing clinical and research knowledge, and delivering high standards of patient-centered, evidence-based care. To promote ocular health locally and globally, embedding lifelong commitment to program values.





We are pleased to introduce this updated edition of the MS Ophthalmology curriculum, designed to provide a comprehensive, structured, and competency-driven educational experience for our residents. This revision reflects our commitment to excellence in ophthalmic education, integrating feedback from faculty and residents, advancements in medical knowledge, and best practices in training.

# **Introduction to curriculum**

A curriculum is a structured framework outlining the essential knowledge, skills, attitudes, and competencies that learners need to acquire over a specified training period. It includes a comprehensive plan for teaching, assessment, and learning activities tailored to achieve the desired educational outcomes. A curriculum aligns learning objectives with educational content, teaching methodologies, and assessment strategies to ensure that learners develop both theoretical knowledge and practical skills needed for their professional roles.

# **ACGME Competency Model**

At Rawalpindi Medical University (RMU), we are committed to training our residents according to the Accreditation Council for Graduate Medical Education (ACGME) competencies, incorporating six core domains: Patient Care, Medical Knowledge, Practice-Based Learning and Improvement, Interpersonal and Communication Skills, Professionalism, and Systems-Based Practice. Additionally, we emphasize research as a fundamental component of our training. RMU proudly leads as the first public sector university in Pakistan to implement an ACGME-aligned postgraduate curriculum, reflecting our dedication to producing competent, compassionate, and research-oriented clinicians prepared to excel in today's dynamic healthcare landscape.

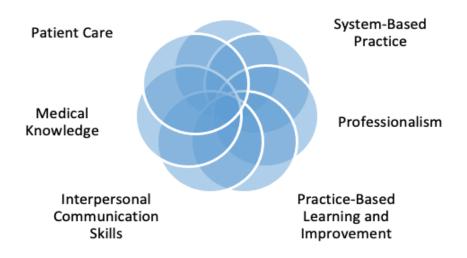


Figure 1: ACGME core competencies (Ref: <u>https://sl.bing.net/hO8ZQtFeM1I</u>)



# **Overview of Program**

The **MS Ophthalmology** program is a well-established and globally recognized medical degree designed to equip trainees with in-depth expertise in the field of ophthalmology. This program, conferred as a **Master of Surgery in Ophthalmology** (**MS Ophthalmology**), reflects decades of tradition and excellence in ophthalmic education and clinical training. The structured curriculum prepares residents with essential diagnostic, therapeutic, and surgical skills to advance ophthalmic healthcare and research, meeting international standards of medical education and patient care. The program of MS Ophthalmology of Rawalpindi Medical University is conducted with a goal to develop ophthalmologists who can provide quality eye care to meet the needs of patients both now and in the future, and who can contribute to the field of ophthalmology through participation in research.

## Admission criteria

#### 1. Resident Appointments

#### **Eligibility Criteria**

For admission in MS Ophthalmology course, the candidate shall be required to have:

- MBBS degree
- Completed one year House Job
- One year experience in Ophthalmology/General surgery/Allied surgical discipline in the given order of preference
- Registration with PMDC
- Passed Entry Test conducted by the University & aptitude interview by the Institute concerned
- Merit will be adhered to strictly for induction as per RMU rules.

**Exemptions:** A candidate holding FCPS ophthalmology/FRCS ophthalmology/Diplomat American Board shall be exempted from Entrance and Midterm Examinations and shall be directly admitted for Exit Examination, subject to fulfillment of requirements for the examination.

#### **Number of Residents**

The minimum number of residents in an accredited four-year program is eight or two per year.

#### **Training sites**

- 1. Holy Family Hospital, Rawalpindi Medical University
- 2. Benazir Bhutto Hospital, Rawalpindi Medical University

Monthly case presentations, journal club meetings and academic tests will be held for all the residents at each participating site alternatively.



# Program Leadership

#### **Program Director**

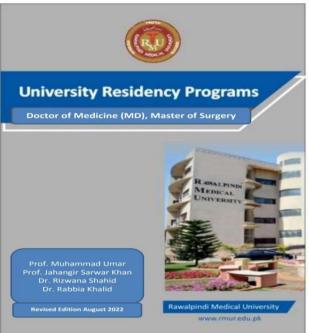
The Program Director, serving as the Chairman of the Ophthalmology Department at Rawalpindi Medical University, oversees the residency program.

#### Faculty

The ophthalmology teaching faculty includes the Professor, Associate Professor, Assistant Professor, and Senior Registrars, all appointed according to PMDC standards, and actively engaged in resident education and training.

#### **Additional Program Personnel**

The Qualified allied vision science professionals, including an Optometrist, Orthoptist, and Ophthalmic Technologist, support resident training in their specialized fields within the ophthalmology department.



## **Rules and Regulations**

Residents must maintain at least 80% attendance in clinical rotations, academic sessions, and case presentations, with full participation in exams, journal clubs, and assessments, all under faculty supervision. They must demonstrate professionalism, ethical conduct, and respect toward patients, staff, and peers. Competency in diagnostic and therapeutic ophthalmic procedures must be achieved progressively, alongside completion of coursework, case logs, and continuous assessments. Participation in research, presentations, conferences, and surgical audits is required, with regular feedback from faculty to address deficiencies. Duty hours are capped at 80 hours per week to ensure well-being, and any non-compliance with program rules may lead to disciplinary action or delay in certification.



# Statutes

The MS Ophthalmology Residency Program at Rawalpindi Medical University follows structured statutes that guide clinical training, academic performance, and ethical practice. These statutes define essential competencies, promote professionalism, require research participation, and provide regular assessments to support growth. Collaboration with hospitals and community health initiatives expands clinical exposure, aligning with the program's mission to develop skilled, ethical, and community-focused ophthalmologists.

## **Program Structure**

The duration of MS Ophthalmology course shall be four (4) years consisting of structured training in a recognized department under the guidance of an approved supervisor. The course is structured in two phases:

**Phase I** is structured for the 1<sup>st</sup> and 2<sup>nd</sup> calendar year. Doctors entering this will require closely supervised training in basic examination methods and techniques and should rapidly be introduced to the elements of surgery and the management of general outpatients and accident and emergency ophthalmic patients. In their second year, they will be expected to take a larger role in both theatre and outpatients, where they will benefit from special clinics. The training units should therefore provide a broad-based training in general ophthalmic medicine and surgery and exposure to the common subspecialties. The candidate shall undertake didactic and interactive training in Basic Ophthalmic Medical and Surgical Sciences, Optics & Refraction, Biostatistics & Research Methodology and Community Ophthalmology. At the end of 1<sup>st</sup> year, a theory and clinical based formative assessment will be conducted. At the end of 2<sup>nd</sup> year mid-term examination (MTA) will be held, comprising of MCQ based theory and Clinical OSCE.

**Phase II** is structured for 3<sup>rd</sup> and 4<sup>th</sup> calendar years in MS Ophthalmology. The trainee should see sufficient patients in a clinic to develop competency and fluency in managing patients in an outpatient setting but the number seen must not be excessive to the extent that training is impaired. The actual number of patients seen should be appropriate to the competency of the trainee and the complexity of the clinical condition of the patient. Surgical experience should develop as indicated by the learning outcomes in the curriculum. It is essential for the trainee to perform sufficient numbers of surgical cases (particularly cataract procedures) to experience a full range of clinical situations so that the trainee learns techniques to manage a range of cases and becomes competent in managing complications. At the end of 3<sup>rd</sup> year, a theory and clinical based formative assessment will be conducted. The candidate will have to achieve sufficient clinical and research capability during this phase so as to qualify his final term examination (FTA) for the award of degree.



# Framework of the Program

Component	Details	
Course Title	MS OPHTHALMOLOGY	
Training Center	Department of Ophthalmology, Rawalpindi	
	Medical University (RMU) & Allied Hospitals	
Duration of Course	4 years	
Credit Hours	132 hours	
Supervision	Structured training under the guidance of an	
	approved supervisor.	
	Induction Period	
Basic Training (Phase I)		
Duration	1 <sup>st</sup> and 2 <sup>nd</sup> year in the Department of	
Focus	Ophthalmology	
	Orientation and training in Ophthalmology and	
Rotations	mandatory workshops	
	1 month in Emergency Medicine	
Assessment (Part I)	Continuous internal assessment based on competency	
Mid Term Assessment	& Formative assessment: In-Training- Assessment	
(MTA)	Year-1 At the end of 2 years, candidates will take the Mid	
	Term Assessment (Summative)	
Advanced Training (Phase II)		
Duration	3 <sup>rd</sup> and 4 <sup>th</sup> year	
Focus	Ophthalmology, Research, and Thesis writing	
Rotations		
	16 weeks	
Assessment (Part II)	Competency-based continuous internal assessment & Formative assessment: In-Training Assessment Year-3	
Final Term Assessment	At the end of four years, candidates will take the Final	
(FTA)	term assessment (Summative)	
Research Component	Research component aligned with the Research Cycle,	
	including thesis writing and submission according to	
	RMU guidelines	



Phase	Year	Placement Departments and Duration	Research	Assessment
Phase 1	1 <sup>st</sup> year	<ul> <li>Ophthalmology (11 months)</li> <li>Emergency medicine (1 month)</li> </ul>	<ul> <li>1 Disease statistical review</li> <li>Synopsis topic submission</li> </ul>	<ul> <li>Formative</li> <li>In-Training (Theory &amp; Clinical)</li> </ul>
Flidse I	2 <sup>nd</sup> year	<ul> <li>Ophthalmology (12 months)</li> </ul>	<ul> <li>Synopsis submission</li> <li>DRB</li> <li>ERB/IRF</li> <li>BASR</li> </ul>	<ul> <li>Summative</li> <li>MTA (Mid training Assessment) (Theory &amp; Clinical)</li> </ul>
Phase 2	3 <sup>rd</sup> year	<ul> <li>Ophthalmology (20 months)</li> <li>Dermatology (2 weeks)</li> <li>Radiology (2</li> </ul>	<ul> <li>Data collection</li> <li>Data analysis</li> <li>Thesis writing</li> </ul>	<ul> <li>Formative</li> <li>In-Training (Theory &amp; Clinical)</li> </ul>
Phase 2	4 <sup>th</sup> year	<ul> <li>weeks)</li> <li>Pathology (2 weeks)</li> <li>Oncology (2 weeks)</li> <li>Neurology (2 weeks)</li> <li>Plastic surgery (2 weeks)</li> <li>Community Ophthalmology (1 month)</li> </ul>	<ul> <li>BASR- Thesis approval</li> <li>Thesis completion certificate (DME)</li> </ul>	<ul> <li>Summative</li> <li>FTA (Final term assessment) (Theory &amp; Clinical)</li> </ul>

# Training Pathway MS ophthalmology



#### Clinical and Surgical Training Resources Ambulatory and In-patient Department

The hospitals serve a diverse patient population with a wide range of ophthalmologic conditions, encompassing both adult and pediatric cases. This high patient volume enables residents to refine their diagnostic, therapeutic, and procedural skills and to evaluate treatment efficacy. Additionally, the inpatient department includes an exam room equipped with a slit-lamp for detailed evaluations.

#### **Operating rooms**

Each participating site provides fully equipped operating rooms for ophthalmic procedures, including anterior and posterior segment surgeries, strabismus correction, and oculoplastic. Training is enhanced by teaching aids attached to operating microscopes, along with audiovisual systems that broadcast live procedures to a separate teaching room for resident observation and learning.

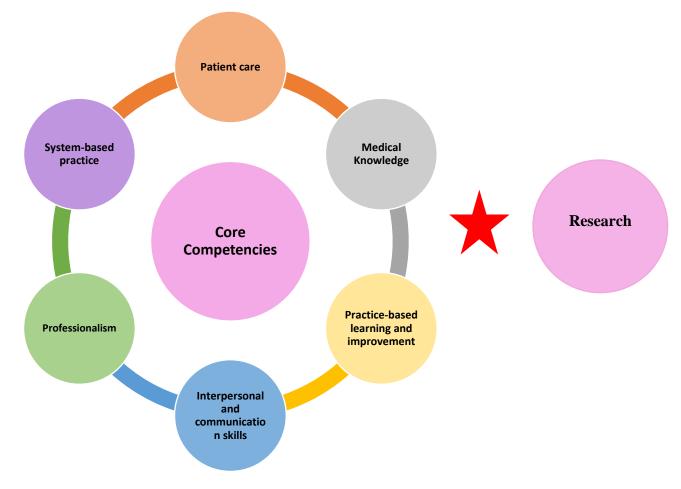
#### **Diagnostic Ophthalmology and Lasers**

Residents have access to essential diagnostic tools, including fundus photography, fluorescein angiography, automated perimetry, ultrasonography, keratometry, orthoptic assessment equipment, refraction tools, Nd-YAG and argon laser.



## **RMU Postgraduate Competency Model**

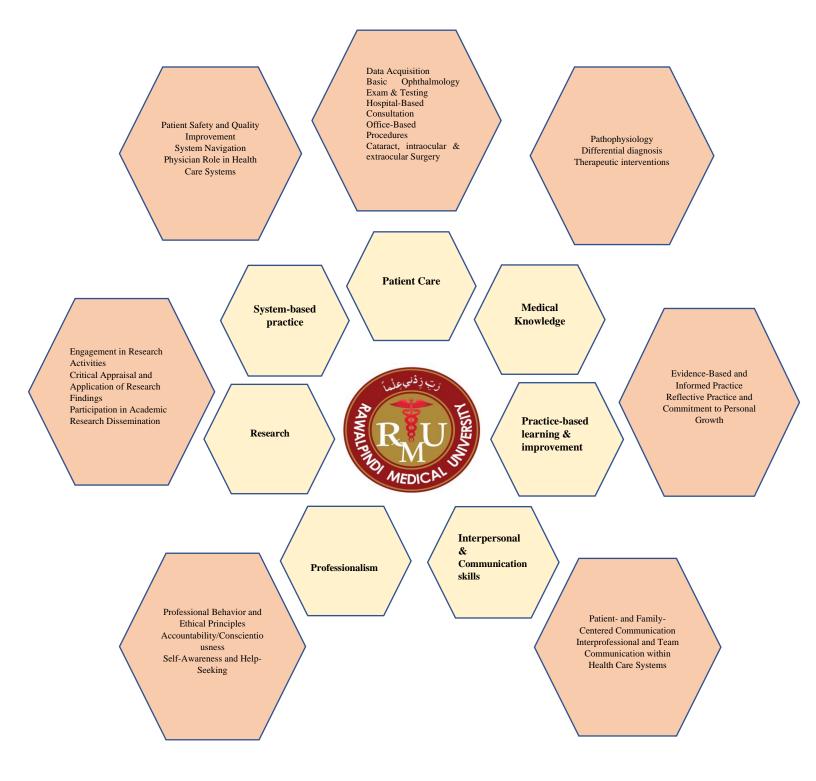
The Rawalpindi Medical University (RMU) postgraduate Competency Model is designed to equip residents for the dynamic demands of modern healthcare. Rooted in ACGME core principles with research as an additional core competency, this model defines essential competencies for all RMU graduates, aligning them with both Pakistan's healthcare priorities and global standards. This approach ensures RMU postgraduates emerge as skilled clinicians, ethical leaders, and compassionate problem-solvers. Complementing this, the RMU postgraduate Competency Model takes a holistic approach, mapping clear developmental pathways from foundational knowledge to advanced clinical practice. Through these competencies, RMU residents are prepared to thrive across diverse healthcare settings, adapt to rapid changes, and contribute meaningfully to society.



# **RMU Postgraduate Competency Model**



# **RMU Postgraduate Competency Framework of Ophthalmology**



# **RMU Ophthalmology Postgraduate Competency Framework**



## Aim of The Program

The aim of the MS Ophthalmology Residency Program is to develop skilled and compassionate ophthalmologists who excel in patient-centered care and integrate advanced medical knowledge. The program fosters professionalism, effective communication, and a commitment to lifelong learning, preparing residents to diagnose and treat diverse ocular conditions. Ultimately, it strives to produce leaders who advocate for optimal patient outcomes, collaborate within healthcare teams, and address the needs of their communities through research and education.

## **Outcomes of the Program**

#### 1. Patient Care

#### Data Acquisition – Basic Ophthalmology Exam and Testing

• To interview, examine, and use appropriate tests to assess a given condition independently.

#### **Hospital-Based Consultation**

- To triage and manage hospital-based consultation independently.
   Office-Based Procedures
- To perform common office-based procedures independently.
   Cataract Surgery Technical Skill
- To perform cataract surgery and manage complications independently. Extraocular Surgery (Plastics, Strabismus)
- To perform extraocular surgery and manage complications. Intraocular Surgery (Cornea, Retina, Glaucoma)
- To gain experience with surgery in these subspecialties.

#### 2. Medical Knowledge

#### Pathophysiology

• To demonstrate progressive understanding of the pathophysiology of common and complex ophthalmic conditions.

#### **Differential Diagnosis**

 To progress in knowledge from creating a broad differential to a problem-focused differential to guide accurate clinical evaluation and management, and avoid unnecessary testing and use of resources.

#### **Therapeutic Interventions**

• To obtain comprehensive understanding of medical and surgical therapeutic interventions.



#### 3. Practice-Based Learning and Improvement

#### Evidence-Based and Informed Practice

- To incorporate evidence and patient values into clinical practice. Reflective Practice and Commitment to Personal Growth
- To seek clinical performance information with the intent to improve care; reflects on all domains of practice, personal interactions, and behaviors, and their impact on colleagues and patients (reflective mindfulness); develop clear objectives and goals for improvement in some form of a learning plan.

#### 4. Interpersonal and Communication Skills

#### Patient- and Family-Centered Communication

 To deliberately use language and behaviors to form constructive relationships with patients, identify communication barriers including self-reflection on personal biases, and minimize them in the doctor-patient relationships; to organize and lead communication around shared decision making.

#### Interprofessional and Team Communication

• To effectively communicate with the health care team, including consultants, in both straightforward and complex situations

#### **Communication within Health Care Systems**

• To effectively communicate using a variety of methods

#### 5. Professionalism

#### **Professional Behavior and Ethical Principles**

 To recognize and address lapses in ethical and professional behavior, demonstrate ethical and professional behaviors, and use appropriate resources for managing ethical and professional dilemmas.

#### Accountability/Conscientiousness

• To take responsibility for one's own actions and the impact on patients and other members of the health care team.

#### Self-Awareness and Help-Seeking

• To identify, use, manage, improve, and seek help for personal and professional well-being for self and others.



#### 6. Systems-Based Practice

#### Patient Safety and Quality Improvement (QI)

 To engage in the analysis and management of patient safety events, including relevant communication with patients, families, and health care professionals; to conduct a QI project.

#### System Navigation for Patient-Centered Care

• To effectively navigate the health care system, including the interdisciplinary team and other care providers, to adapt care to a specific patient population to ensure high-quality patient outcomes.

#### Physician Role in Health Care Systems

• To evaluate and understand the physician's role in the complex health care system and how to optimize the system to improve patient care and the health system's performance.

#### 7. Research

#### **Engagement in Research Activities**

• Develop research skills by engaging in clinical or basic research activities, contributing to the body of knowledge in ophthalmology.

#### Critical Appraisal and Application of Research Findings

• Critically appraise and apply research findings to clinical practice, using evidence-based approaches to improve patient care.

#### Participation in Academic Research Dissemination

• Participate in presentations, conferences, or publications, actively contributing to the research mission of the Ophthalmology Department.



# **SECTION II: COURSE CONTENT**



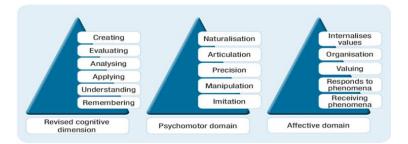


# **Content Overview**

The MS Ophthalmology program is structured to provide a comprehensive understanding of both **Clinical Ophthalmology** and **Optics & Refraction**, ensuring residents develop the knowledge and skills essential for diagnosing, treating, and managing a wide range of ocular conditions. A detailed syllabus is crucial for structuring the learning pathway, ensuring comprehensive coverage of essential topics and competencies in ophthalmology. By breaking down each content area, the curriculum enables focused study, promotes skill development in specific sub-specialties, and ensures that residents are prepared for both routine and complex cases in their practice. It also facilitates uniformity across training institutions, helps meet accreditation standards, and provides a benchmark for evaluation and assessment throughout the program.

## **Bloom's Taxonomy Domains**

To effectively guide learning and assessment in the MS Ophthalmology program, **Cognitive**, **Psychomotor**, and **Affective** domains are outlined below, based on Bloom's Taxonomy.



<b>Cognitive Domain</b>	Psychomotor Domain	Affective Domain
Remembering C1	Imitation/Perception P1	Receiving A1
Understanding C2	Manipulation/guided response P2	Responding A2
Applying C3	Precision/complex overt response P3	Valuing A3
Analyzing C4	Articulation P4	Organization A4
Evaluating C5	Naturalization P5	Characterization by value set A5
Creating C6		

The residents must demonstrate technical skills for effective patient care at minimum levels of competencies identified.

#### Key for Assessing Competencies:

- 1. Level 1: Observer Status.
- 2. Level 2: Assistant Status.
- 3. Level 3: Performed Under Direct Supervision.
- 4. Level 4: Performed Under Indirect Supervision.

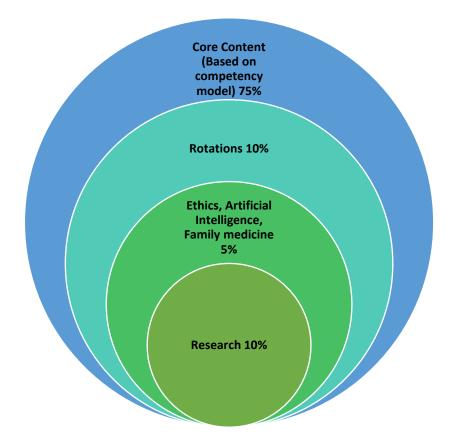


# **Credit Hours of MS Ophthalmology Program**

According to the HEC criteria, 16 teaching/learning hours equate to 1 credit hour:

- 1. For each training year:
  - 33 credit hours x 16 hours = **528 teaching/learning hours per year**.
- 2. For the 4-year program:
  - $\circ$  528 hours x 4 years = 2,112 total teaching/learning hours for the entire program.

Therefore, the **MS Ophthalmology training program comprises a total of 2,112 teaching/learning hours, equating to 132 credit hours** over the four years. This extensive allocation aligns with HEC criteria and supports a robust foundation in clinical and academic training for ophthalmology residents.



## **Core Content Clinical Ophthalmology**

#### Modules And Themes

The comprehensive content of ophthalmology is divided into four modules for postgraduate residents and relevant themes are assigned for each module.

#### **Module 1: Anterior Segment**

Theme: Diseases and Disorders of the Eyelids, Lacrimal System, Conjunctiva and Orbit

• Eyelids (Weeks 1-8)



- Introduction
- Non-neoplastic lesions
- Benign epidermal tumors
- Benign pigmented lesions
- Benign adnexal tumors
- Miscellaneous benign tumors
- Malignant tumors
- Disorders of the eyelashes
- Allergic disorders
- Bacterial infections
- Viral infections
- Blepharitis
- o Ptosis
- Ectropion
- Entropion
- Miscellaneous acquired disorders
- Cosmetic eyelid and periocular surgery
- Congenital malformations

#### • Lacrimal Drainage System (Weeks 9-10)

- Introduction
- Acquired obstruction
- Congenital obstruction
- Chronic canaliculitis
- Dacryocystitis

#### • Conjunctiva (Weeks 11-12)

- Introduction
- Bacterial conjunctivitis
- Viral conjunctivitis
- Allergic conjunctivitis
- Conjunctivitis in blistering mucocutaneous disease
- Miscellaneous conjunctivitides
- Degenerations
- Subconjunctival hemorrhage
- Orbit (Weeks 13-16)
  - Introduction
  - Thyroid eye disease
  - $\circ$  Infections
  - Non infective inflammatory disease
  - Vascular abnormalities
  - Cystic lesions
  - Tumors
  - Anophthalmic socket
  - Craniosynostosis

#### Module 2: Cornea, Sclera, and Refractive Surgery

#### **Theme: Corneal Diseases and Surgical Interventions**

- Cornea (Weeks 17-26)
  - $\circ$  Introduction



- Bacterial keratitis
- Fungal keratitis
- Herpes simplex keratitis
- Herpes zoster ophthalmicus
- Interstitial keratitis
- Protozoan keratitis
- Helminthic keratitis
- o Bacterial hypersensitivity-mediated corneal disease
- Rosacea
- Peripheral corneal ulceration/thinning
- Neurotrophic keratopathy
- Exposure keratopathy
- o Miscellaneous keratopathies
- Corneal ectasias
- Corneal dystrophies
- Corneal degenerations
- Metabolic keratopathies
- Contact lenses
- Congenital anomalies of the cornea and globe

#### • Corneal and Refractive Surgery (Weeks 27-30)

- Keratoplasty
- Keratoprostheses
- Refractive procedures

#### • Episclera and Sclera (Weeks 31-34)

- Anatomy
- Episcleritis
- Immune-mediated scleritis
- Infectious scleritis
- Scleral discoloration
- Blue sclera
- Miscellaneous conditions

#### Module 3: Glaucoma, Lens, and Uveitis

#### Theme: Intraocular Pressure, Cataract Management, and Inflammatory Eye Diseases

#### • Glaucoma (Weeks 35-44)

- Introduction
- Tonometry
- Gonioscopy
- Evaluation of the optic nerve head
- Imaging in glaucoma
- Perimetry
- Medical treatment of glaucoma
- Laser treatment of glaucoma
- Trabeculectomy
- Non-penetrating glaucoma surgery
- o Drainage shunts
- Ocular hypertension
- Primary open-angle glaucoma



- Normal-tension glaucoma
- Primary angle-closure glaucoma
- Classification of secondary glaucoma
- Pseudoexfoliation
- Pigment dispersion
- Neovascular glaucoma
- Inflammatory glaucoma
- Lens-related glaucoma
- Traumatic glaucoma
- o Iridocorneal endothelial syndrome
- Glaucoma associated with intraocular tumors
- o Glaucoma secondary to epithelial ingrowth
- $\circ$  Iridoschisis
- Primary congenital glaucoma
- Iridocorneal dysgenesis
- Glaucoma in phacomatoses

#### • Lens (Weeks 45-48)

- Acquired cataract
- Management of age-related cataract
- Congenital cataract
- Ectopia lentis
- Abnormalities of lens shape

#### Uveitis (Weeks 49-52)

- Classification
- Anterior uveitis
- Uveitis in spondyloarthropathies
- Fuchs uveitis syndrome
- Uveitis in juvenile idiopathic arthritis
- Uveitis in bowel disease
- Uveitis in renal disease
- Intermediate uveitis
- Vogt-Koyanagi-Harada (VKH) syndrome
- Sympathetic ophthalmitis
- Lens-induced uveitis
- Sarcoidosis
- Behçet disease
- Parasitic uveitis
- Viral uveitis
- Fungal uveitis
- Bacterial uveitis
- Miscellaneous idiopathic chorioretinopathies

#### **Module 4: Posterior Segment and Miscellaneous Topics**

#### Theme: Vitreoretinal Diseases, Ocular Oncology, and Systemic Effects

#### • Retinal Vascular Disease (Weeks 53-57)

- Retinal circulation
- Diabetic retinopathy
- Non-diabetic retinopathy



- Retinal venous occlusive disease
- Retinal arterial occlusive disease
- Ocular ischemic syndrome
- Hypertensive eye disease
- Sickle cell retinopathy
- Thalassemia retinopathy
- Retinopathy of prematurity
- Retinal artery macroaneurysm
- Primary retinal telangiectasia
- o Eales disease
- Radiation retinopathy
- Purtscher retinopathy
- Valsalva retinopathy
- Lipaemia retinalis
- Retinopathy in blood disorders

#### • Acquired Macular Disorders (Weeks 58-62)

- Introduction
- Clinical evaluation of macular disease
- Investigation of macular disease
- o Age-related macular degeneration
- Retinal angiomatous proliferation
- o Polypoidal choroidal vasculopathy
- Peripheral exudative hemorrhagic chorioretinopathy
- o Idiopathic choroidal neovascularization
- Vitreomacular interface disorders
- Central serous chorioretinopathy
- o Idiopathic macular telangiectasia
- o Cystoid macular edema
- o Microcystic macular edema
- Degenerative myopia
- Angioid streaks
- Choroidal folds
- Hypotony maculopathy
- Solar retinopathy
- Focal choroidal excavation

#### • Hereditary Fundus Dystrophies (Weeks 63-65)

- Introduction
- Investigation
- o Generalized photoreceptor dystrophies
- Macular dystrophies
- Generalized choroidal dystrophies
- Hereditary vitreoretinopathies
- $\circ$  Albinism
- Cherry-red spot at the macula
- Retinal Detachment and Vitreous Opacities (Weeks 66-70)
  - Introduction
  - Peripheral lesions predisposing to retinal detachment
  - Posterior vitreous detachment
  - o Retinal breaks
  - Rhegmatogenous retinal detachment



- o Tractional retinal detachment
- Exudative retinal detachment
- Pars plana vitrectomy
- Vitreous opacities

#### Ocular Tumors (Weeks 71-75)

- Benign epibulbar tumors
- Malignant and premalignant epibulbar tumors
- $\circ$  Iris tumors
- Iris cysts
- Ciliary body tumors
- Tumors of the choroid
- Neural retinal tumors
- Retinal vascular tumors
- Primary intraocular lymphoma
- Tumors of the retinal pigment epithelium
- Paraneoplastic syndromes

#### • Neuro-ophthalmology (Weeks 76-80)

- Neuroimaging
- Optic nerve
- o Pupils
- o Chiasm
- Retrochiasmal pathways
- Ocular motor nerves
- Supranuclear disorders of ocular motility
- o Nystagmus
- Ocular myopathies
- Miller Fisher syndrome
- Neurofibromatosis
- Migraine
- Neuralgias
- Facial spasm

#### • Trauma and Ocular Side Effects of Systemic Medication (Weeks 81-82)

- Eyelid trauma
- Orbital trauma
- Trauma to the globe
- Chemical injuries
- Ocular side effects of systemic medication

#### • Ocular Anesthesia & Surgeries (Weeks 83-84)

- Surface, infiltration, regional anesthesia
- Premedication, sedation for local anesthesia
- Premedication for general anesthesia
- o Akinesia & intraocular tension during anesthesia
- Cardio pulmonary complication with anesthesia
- Cardiac arrest & local anesthetic emergency
- Operative Surgeries

#### • Ocular Diagnostic & Operative Instrument (Weeks 85-86)

- Radiology in ophthalmologic diagnosis
- Ultrasonography A scan & B scan



- Fluorescein angiography.
- Pachymeter
- Auto perimeter
- Autorefractometer
- Applanation tonometry
- o Indirect ophthalmoscope
- Recent trends / advances in ophthalmology

#### • Artificial Intelligence (AI) in Imaging and Diagnostics

- AI applications in retinal imaging (e.g., fundus photography, OCT)
- AI in glaucoma screening and progression prediction
- AI for corneal disease detection (e.g., keratoconus screening)
- Role of AI in cataract grading and surgical planning
- AI in pediatric ophthalmology, especially for amblyopia and strabismus detection

#### • AI in Surgery and Robotics

- Overview of AI-guided robotic systems in ocular surgery
- Augmented reality (AR) and AI-based intraoperative assistance
- Virtual surgery planning and simulation tools

#### • Ethics in Ophthalmology

- Foundations of Medical Ethics
- Principles of autonomy, beneficence, non-maleficence, and justice.
- Ethics in informed consent and patient rights.

#### • Confidentiality and Patient Privacy in Eye Care

- Handling patient information securely.
- Addressing ethical concerns related to sensitive diagnoses (e.g., hereditary conditions).

#### • Ethical Dilemmas in Ophthalmology

- Common issues like treatment refusal, resource limitations, and cost considerations.
- Discussion of scenarios and ethical decision-making frameworks.
- Ethical Communication
- Maintaining respect, empathy, and cultural competence in patient interactions.
- Handling difficult conversations about prognosis and lifestyle impacts of eye conditions.

#### • Research Ethics and Integrity

- Guidelines for ethical research with human subjects.
- Informed consent and transparency in clinical trials and observational studies.



#### Implementation in the Academic Calendar

- Module 1: Anterior Segment (Weeks 1-16)
- Module 2: Cornea, Sclera, and Refractive Surgery (Weeks 17-34)
- Module 3: Glaucoma, Lens, and Uveitis (Weeks 35-52)
- Module 4: Posterior Segment and Miscellaneous Topics (Weeks 53-85)

Regular assessments and practical sessions will be integrated into the schedule to reinforce learning and provide hands-on experience.

## **Core Content Optics and Refraction**

- Properties of Light and Visual Function
- Reflection of Light
- Refraction of Light
- o Prisms
- Spherical Lenses
- Astigmatic Lenses
- Optical Prescriptions, Spectacle Lenses
- Aberrations of Optical systems Including the Eye
- Refraction by the Eye
- Optics of Ametropia
- o Presbyopia
- Contact Lenses
- o Optics of Low Vision Aids
- Instruments
- o Lasers
- Practical Clinical Refraction
- o Refractive Surgery

#### Modules and Themes

#### Module 1: Fundamentals of Light and Optics (Weeks 1-8

#### Themes:

- Properties of Light and Visual Function
- Reflection of Light
- Refraction of Light

#### Module 2: Optical Systems and Lenses (Weeks 9-16)

#### Themes:

- o Prisms
- Spherical Lenses
- Astigmatic Lenses
- Aberrations of Optical Systems, Including the Eye



#### Module 3: Clinical Application of Optics (Weeks 17-24)

#### Themes:

- Optical Prescriptions and Spectacle Lenses
- Refraction by the Eye
- Optics of Ametropia
- o Presbyopia

#### Module 4: Specialized Optics and Technology in Ophthalmology (Weeks 25-32)

#### Themes:

- Contact Lenses
- Optics of Low Vision Aids
- Instruments
- o Lasers
- Practical Clinical Refraction
- Refractive Surgery



# **SECTION III: SPECIFIC LEARNING OUTCOMES**





# Specific Learning Outcomes Optics and Refraction

### 1<sup>st</sup> Year Resident Outcomes

By the completion of their 1<sup>st</sup> year residency, the residents should be able to:

Physic	al Optics	Cognitive Domain
1.	Describe the wave and particle nature of light.	C2, C3
2.	Explain the phenomenon of diffraction.	
3.	Explain the concepts of interference and coherence.	
4.	Define optical resolution.	
5.	Explain polarization.	
6.	Explain light scattering.	
7.	Define and compare transmission and absorption.	
8.	Explain photometry.	
9.	Define illumination.	
10.	Describe image quality.	
11.	Differentiate brightness and radiance.	
12.	Define refractive index.	
Geome	etric Optics	
Reflect	tion (Mirrors)	
1.	List the laws of reflection.	
2.	Explain images and objects as light sources.	
3.	Define refractive index.	
Refrac	tion	
1.	Explain the law of refraction (Snell law), including:	
	a. Passage of light from one medium to another	
	b. Absolute index of refraction	
	c. Total internal reflection	
2.	Explain critical angle and total internal reflection.	
Prisms		
1.	Define a prism.	C2, C3
2.	Explain the notation of prisms (e.g., prism diopters).	
3.	Describe the use of prisms in ophthalmology (i.e., diagnostic and	
	therapeutic).	
4.	Explain prentice rule.	
5.	Describe Fresnel and similar prisms.	



<ul> <li>6. Explain the concept of thin prisms.</li> <li>7. Explain the prismatic effect of lenses.</li> <li>8. Define spherical decentration and prism power.</li> <li>C2, C3</li> <li>Spherical Lenses <ol> <li>Define a spherical lens.</li> <li>Describe the cardinal points.</li> <li>Recite the thin lens and thick lens formulas.</li> <li>Define vergence of light, including diopter, convergence, divergence, and vergence formula.</li> <li>Define the terms concave and convex.</li> <li>Define the terms concave and surfaces</li> <li>b. Cross cylindrical lenses, including: <ol> <li>Spherical decronic.</li> </ol> </li> <li>Astigmatic Lenses</li> <li>Describe toric lenses.</li> </ol> </li> <li>Clinical Optics <ol> <li>Captine ametropia.</li> <li>Define ametropia.</li> <li>Define antisekonia (including Knapp rule).</li> <li>Define antisekonia (including Knapp rule).</li> <li>Define aphakia.</li> <li>Explain optical parameters affecting retinal image size.</li> <li>Describe topical parameters affecting retinal image size.</li> <li>Describe topical acuity, including: <ol> <li>Distance and near acuity measurement</li> <li>Minimal acuity (i.e., visible, perceptible, separable, legible)</li> <li>Visual acuity charts</li> </ol> </li> <li>Describe higher-order aberrations of the eye.</li> <li>Explain how accommodation is affected by age.</li> <li>Explain how accommodation is affected by age.</li> <li>Explain how accommodative problems.</li> <li>Describe the publice ffect impacts visual acuity.</li> <li>Explain how the pinhole effect impacts visual acuity.</li> <li>Explain how the pinhole effect impacts visual acuity.</li> <li>Explain accommodative problems.</li> <li>Describe the epidemiology of refractive errors, including:</li> </ol> </li> <li>2. Outpace and near acuity measurement bio. Bais accommodative insufficiency or excess.</li> <li>Explain how accommodative problems.</li> <li>Describe</li></ul>			
<ul> <li>8. Define spherical decentration and prism power.</li> <li>C2, C3</li> <li>Spherical Lenses <ol> <li>Define a spherical lens.</li> <li>Describe the cardinal points.</li> <li>Recite the thin lens and thick lens formulas.</li> <li>Define the terms concave and convex.</li> <li>Define the terms concave and convex.</li> <li>Define the term magnification, including linear, angular, relative size, and electronic.</li> </ol> </li> <li>Astigmatic Lenses <ol> <li>Describe cylindrical lenses, including: <ol> <li>Spherocylindrical lenses and surfaces</li> <li>Cross cylinders (e.g., Jackson cross cylinder)</li> <li>Describe toric lenses.</li> </ol> </li> <li>Clinical Optics <ol> <li>Define ametropia.</li> <li>Define anteropia.</li> <li>Define anteropia.</li> <li>Define antisometropia.</li> <li>Define aniseikonia (including Knapp rule).</li> <li>Define aniseikonia (including Knapp rule).</li> <li>Define aphakia.</li> <li>Explain optical parameters affecting retinal image size.</li> <li>Describe the pupillary response and its effect on the resolution of the optical system (Stiles-Crawford effect).</li> <li>Define visual acuity, including: <ol> <li>Distance and near acuity measurement</li> <li>Minimal acuity (i.e., visible, perceptible, separable, legible)</li> <li>Visual acuity charts</li> </ol> </li> <li>Describe higher-order aberrations of the eye.</li> <li>Explain how accommodation is affected by age.</li> <li>Explain how the pinhole effect impacts visual acuity.</li> <li>Explain how the pinhole effect impacts visual ac</li></ol></li></ol></li></ul>	6.	Explain the concept of thin prisms.	
<ul> <li>Spherical Lenses <ol> <li>Define a spherical lens.</li> <li>Describe the cardinal points.</li> <li>Recite the thin lens and thick lens formulas.</li> <li>Define vergence of light, including diopter, convergence, divergence, and vergence formula.</li> <li>Define the terms concave and convex.</li> <li>Define the term magnification, including linear, angular, relative size, and electronic.</li> </ol> </li> <li>Astigmatic Lenses <ol> <li>Describe cylindrical lenses, including: <ol> <li>Spherocylindrical lenses and surfaces</li> <li>Cross cylinders (e.g., Jackson cross cylinder)</li> </ol> </li> <li>Describe toric lenses.</li> </ol> </li> <li>Clinical Optics <ol> <li>Define emmetropia.</li> <li>Define anteropia.</li> <li>Define anteropia.</li> <li>Define antigmatism.</li> <li>Define ansigmatism.</li> <li>Define anisometropia.</li> <li>Define anisometropia.</li> <li>Define anisometropia.</li> <li>Define anisometropia.</li> <li>Define aniseikonia (including Knapp rule).</li> <li>Define aphakia.</li> <li>Explain optical parameters affecting retinal image size.</li> <li>Describe the pupillary response and its effect on the resolution of the optical system (Stiles-Crawford effect).</li> <li>Define visual acuity, including: <ol> <li>Distance and near acuity measurement</li> <li>Minimal acuity (i.e., visible, perceptible, separable, legible)</li> <li>Visual acuity charts</li> </ol> </li> <li>Describe higher-order aberrations of the eye.</li> <li>Explain how accommodation is affected by age.</li> <li>Explain how the pinhole effect impacts visual acuity.</li> <li>Explain how the pinh</li></ol></li></ul>	7.	Explain the prismatic effect of lenses.	
<ol> <li>Define a spherical lens.</li> <li>Describe the cardinal points.</li> <li>Recite the thin lens and thick lens formulas.</li> <li>Define vergence of light, including diopter, convergence, divergence, and vergence formula.</li> <li>Define the terms concave and convex.</li> <li>Define the terms concave and convex.</li> <li>Define the term magnification, including linear, angular, relative size, and electronic.</li> <li>Astigmatic Lenses         <ol> <li>Describe cylindrical lenses, including:</li></ol></li></ol>	8.	Define spherical decentration and prism power.	C2, C3
<ol> <li>Describe the cardinal points.</li> <li>Recite the thin lens and thick lens formulas.</li> <li>Define vergence of light, including diopter, convergence, divergence, and vergence formula.</li> <li>Define the terms concave and convex.</li> <li>Define the term magnification, including linear, angular, relative size, and electronic.</li> <li>Astigmatic Lenses         <ol> <li>Describe toplindrical lenses, including:</li></ol></li></ol>	Spher	ical Lenses	
<ul> <li>3. Recite the thin lens and thick lens formulas.</li> <li>4. Define vergence of light, including diopter, convergence, divergence, and vergence formula.</li> <li>5. Define the term sconcave and convex.</li> <li>6. Define the term magnification, including linear, angular, relative size, and electronic.</li> <li>Astigmatic Lenses <ol> <li>Describe cylindrical lenses, including: <ul> <li>a. Spherocylindrical lenses and surfaces</li> <li>b. Cross cylinders (e.g., Jackson cross cylinder)</li> </ul> </li> <li>2. Describe toric lenses.</li> </ol></li></ul> <li>Clinical Optics <ul> <li>Define mmetropia.</li> <li>Define myopia.</li> <li>Define anteropia.</li> <li>Define antisometropia</li> <li>Define antisometropia.</li> <li>Define antiscikonia (including Knapp rule).</li> <li>Define antiscikonia (including Knapp rule).</li> <li>Define attration of the optical system (Stiles-Crawford effect).</li> </ul> 11. Define visual acuity, including: <ul> <li>a. Distance and near acuity measurement</li> <li>b. Minimal acuty (i.e., visible, perceptible, separable, legible)</li> <li>c. Visual acuty charts</li> </ul> 12. Describe higher-order aberrations of the eye. <ul> <li>13. Explain how accommodative problems.</li> <li>14. Explain how the pinhole effect impacts visual acuity.</li> <li>15. Explain accommodative problems.&lt;</li></ul></li>	1.	Define a spherical lens.	
<ul> <li>4. Define vergence of light, including diopter, convergence, divergence, and vergence formula.</li> <li>5. Define the term sconcave and convex.</li> <li>6. Define the term magnification, including linear, angular, relative size, and electronic.</li> <li>Astigmatic Lenses <ol> <li>Describe cylindrical lenses, including: <ol> <li>Spherocylindrical lenses and surfaces</li> <li>Cross cylinders (e.g., Jackson cross cylinder)</li> </ol> </li> <li>2. Describe toric lenses.</li> <li>Clinical Optics</li> <li>Clinical Optics</li> <li>Define emmetropia.</li> <li>Define mometropia.</li> <li>Define anteropia.</li> <li>Define antisometropia.</li> <li>Define astigmatism.</li> <li>Define astigmatism.</li> <li>Define anisometropia.</li> <li>Define anisometropia.</li> <li>Define aphakia.</li> <li>Explain optical parameters affecting retinal image size.</li> <li>Describe the pupillary response and its effect on the resolution of the optical system (Stiles-Crawford effect).</li> <li>Define visual acuity, including: <ol> <li>Distance and near acuity measurement</li> <li>Minimal acuity (i.e., visible, perceptible, separable, legible)</li> <li>Visual acuity charts</li> </ol> </li> <li>Describe higher-order aberrations of the eye.</li> <li>Explain how the pinhole effect impacts visual acuity.</li> <li>Explain how the pinhole effect impacts visual acuity.</li> <li>Explain how the pinhole effect impacts visual acuity.</li> <li>Explain accommodative problems.</li> </ol> </li> <li>C2, C3</li> </ul>	2.	Describe the cardinal points.	
<ul> <li>divergence, and vergence formula.</li> <li>5. Define the terms concave and convex.</li> <li>6. Define the term magnification, including linear, angular, relative size, and electronic.</li> <li>Astigmatic Lenses <ol> <li>Describe cylindrical lenses, including: <ol> <li>Spherocylindrical lenses and surfaces</li> <li>Cross cylinders (e.g., Jackson cross cylinder)</li> </ol> </li> <li>2. Describe toric lenses.</li> </ol></li></ul> <li>Clinical Optics <ul> <li>C2, C3</li> </ul> </li> <li>2. Define emmetropia.</li> <li>Define myopia.</li> <li>Define myopia.</li> <li>Define ansigmatism.</li> <li>Define anisometropia.</li> <li>Describe the pupillary response and its effect on the resolution of the optical system (Stiles-Crawford effect).</li> <li>11. Define visual acuity, including: <ul> <li>Distance and near acuity measurement</li> <li>Minimal acuity (i.e., visible, perceptible, separable, legible)</li> <li>Visual acuity charts</li> </ul> </li> <li>12. Describe higher-order aberrations of the eye.</li> <li>13. Explain how accommodation is affected by age.</li> <li>14. Explain how accommodation is affected by age.</li> <li>15. Explain how the pinhole effect impacts visual acuity.</li> <li>15. Explain accommodative problems.</li> <li>16. Describe convergence or accommodativ</li>	3.	Recite the thin lens and thick lens formulas.	
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<ul> <li>6. Define the term magnification, including linear, angular, relative size, and electronic.</li> <li>Astigmatic Lenses <ol> <li>Describe cylindrical lenses, including: <ol> <li>Spherocylindrical lenses and surfaces</li> <li>Cross cylinders (e.g., Jackson cross cylinder)</li> </ol> </li> <li>2. Describe toric lenses.</li> </ol></li></ul> <li>Clinical Optics <ul> <li>Define emmetropia.</li> <li>Define myopia.</li> <li>Define ansizementropia.</li> <li>Define aphakia.</li> <li>Explain optical parameters affecting retinal image size.</li> <li>Describe the pupillary response and its effect on the resolution of the optical system (Stilles-Crawford effect).</li> </ul> </li> <li>11. Define visual acuity, including: <ul> <li>Distance and near acuity measurement</li> <li>Minimal acuity (i.e., visible, perceptible, separable, legible)</li> <li>Visual acuity charts</li> </ul> </li> <li>12. Describe higher-order aberrations of the eye.</li> <li>13. Explain how accommodation is affected by age.</li> <li>14. Explain how the pinhole effect impacts visual acuity.</li> <li>15. Explain accommodative problems.</li> <li>16. Describe convergence or accommodative insufficiency or excess.</li> <li>17. Define accommodative-convergence over accommodation (AC/A) ratio.</li>		divergence, and vergence formula.	
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Astigmatic Lenses       1. Describe cylindrical lenses, including:       a. Spherocylindrical lenses and surfaces       b. Cross cylinders (e.g., Jackson cross cylinder)         2. Describe toric lenses.       Clinical Optics       C2, C3         1. Define emmetropia.       Define emmetropia.       C2, C3         2. Define ametropia.       Define myopia.       C2, C3         3. Define myopia.       Define hypermetropia (hyperopia).       C2, C3         5. Define astigmatism.       Define aniseikonia (including Knapp rule).       C2, C3         8. Define aniseikonia (including Knapp rule).       Define aphakia.       C2, C3         9. Explain optical parameters affecting retinal image size.       C2, C3         10. Describe the pupillary response and its effect on the resolution of the optical system (Stiles-Crawford effect).       C3         11. Define visual acuity, including: <ul> <li>a. Distance and near acuity measurement</li> <li>b. Minimal acuity (i.e., visible, perceptible, separable, legible)</li> <li>c. Visual acuity charts</li> </ul> C2, C3         12. Describe higher-order aberrations of the eye.       C3         13. Explain how the pinhole effect impacts visual acuity.       C3         15. Explain how the pinhole effect impacts visual acuity.       C2, C3         15. Explain accommodative problems.       C2, C3         16. Describe convergence or accommodati	6.	Define the term magnification, including linear, angular, relative	
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ratio.	16	. Describe convergence or accommodative insufficiency or excess.	C2, C3
	17	7. Define accommodative-convergence over accommodation (AC/A)	
18. Describe the epidemiology of refractive errors, including:		ratio.	
	18	. Describe the epidemiology of refractive errors, including:	



a.	Prevalence	
b.	Inheritance	
с.	Changes with age	
d.	Surgical considerations	
19. Descri	be the potential problems with aphakic spectacles.	C2, C3
20. Descri	be the effect of spectacles and contact lens correction on	
accom	modation and convergence (i.e., amplitude, near point, far	
point)		
21. Explai	n the principles of contrast sensitivity measurements.	
22. Descri	be the correction of ametropia, including:	
a.	General principles	
b.	Spectacle lenses	
с.	Contact lenses	
d.	Intraocular lenses	
e.	Principles of refractive surgery	
<b>Clinical Refra</b>	ction	
<b>Objective Ref</b>	raction: Retinoscopy	C2, C3
1. List th	e principles and indications for retinoscopy.	
Subjective Re	fraction Techniques	
1. Descri	be the major types of refractive errors.	
2. Descri	be the indications for and use of trial lenses for simple	
refrac	tive error.	
Cycloplegic R	efraction	
1. Descri	be medication concentrations according to age (e.g.	C2, C3
cyclop	entolate, atropine).	

	Psychomotor Domain	Level	of
		Competence	
Geometric Optics			
Reflection (Mirrors)			
<ol> <li>Illustrate reflection at a plane surface (i.e., image and field of a plane mirror).</li> </ol>		3	
<ol> <li>Illustrate reflection at curved surfaces (i.e., focal point and foca length of a spherical mirror).</li> </ol>			
3. Demonstrate a multiple lens system			
Refraction			



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1. Illustrate refraction at a plane	P2 3	3
surface.		
2. Illustrate refraction at curved		
surfaces.		
3. Demonstrate image jump and		
displacement.		
Prisms		
1. Demonstrate the types of prisms		
(e.g., plane, parallel, plate).		
2. Illustrate refraction of light through		
a prism.		
Spherical Lenses		
1. Demonstrate binocular balancing.		
Astigmatic Lenses	3	3
1. Demonstrate how the Maddox rod		
works.	P2	
2. Locate the conoid of Sturm.		
Notation of Lenses		
1. Design myopic, hyperopic, and		
astigmatic lenses.	3	3
2. Perform simple transposition.		
3. Perform toric transposition.	P2	
4. Calculate a lens prescription.		
Aberration of Lenses		
1. Correct aberrations relevant to the		
eye, including spherical, coma,		
astigmatism, and distortion.	3	3
2. Describe color aberrations and		
perform the duochrome test.	P2	
Clinical Optics		
1. Illustrate optics of the eye, including		
the dioptric power of different		
structures.		
2. Draw a schematic eye and reduced		
eye.		
3. Demonstrate contrast sensitivity		
measurements.		
4. Demonstrate the calculation of		3
intraocular lens power.		
	P2	
Clinical Refraction		



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Object	ive Refraction: Retinoscopy		
-	Perform the technique of		
	retinoscopy.		
2.	Perform an integrated refraction		
	based upon retinoscope results.		
3.	Identify media opacities with		
	retinoscopy.		3
4.	Perform cycloplegia.		
5.	Prescribe refractive correction based	P2	
	on the obtained objective and		
	subjective measurements.		
Subjec	tive Refraction Techniques		
1.	Perform elementary refraction		
	techniques for myopia, hyperopia,		
	and near-vision add.		
2.	Perform techniques for the		
	correction for presbyopia (i.e.,		
	measuring for near adds).		
	ments and Tests		
1.	Demonstrate the use of the direct		
	ophthalmoscope.		
2.	Demonstrate the use of the indirect		
	ophthalmoscope.		
3.	Demonstrate the use of the		
	retinoscope.		
4.	Demonstrate glare and contrast		3
_	sensitivity testing.	P2	
5.	Demonstrate the use of the automated refractor.	P2	
6	Demonstrate measurement of		
0.	higher-order aberrations.		
7	Demonstrate the use of stereoacuity		
/.	testing.		
8	Demonstrate the use of corneal		
	topography (e.g., placido disc,		
	keratometer, automated corneal		
	topography).		
9.	Demonstrate the use of the Hess		
_	screen or describe its use if not		
	available.		



10. Demonstrate the use of the	3
synoptophore.	-C-1
11. Demonstrate the use of color vision	P2
tests (e.g., Ishihara color plates;	
Farnsworth-Munsell test).	

# 2<sup>nd</sup> Year Resident Outcomes

By the completion of their 2<sup>nd</sup> year residency, the residents should be able to:

		Cognitive
		Domain
Optics		
Specta	licles	
1.	Describe materials index.	
2.	Describe the principles underlying progressive spectacle lens design.	C2, C3
3.	Describe progressive lenses measurements.	
4.	Describe spectacles specificities in children.	
Lasers		
1.	Describe the technology behind the excimer laser and the femtosecond laser.	
2.	List different wavelengths used in ophthalmic lasers.	
3.	Describe indications for refractive surgery.	
Aberro	ometer Technology	
1.	Explain the principles underlying Hartmann-Shack aberrometers.	
2.	Describe the concept of Zernike polynomials.	
Diagno	ostic Equipment	
1.	List indications for and the use of intraocular lens (IOL) calculation algorithms.	C2, C3
2.	List indications for the use of corneal pachymetry.	
3.	List indications for the use of specular microscopy.	
4.	List indications for the use of corneal tomography with anterior segment optical coherence tomography (OCT).	
5.	List indications for the use of topographic/elevation corneal evaluation (i.e., Penta Cam, Orbscan II, Galilei).	
6.	List indications for the use of laser interferometry for macular testing.	
Refrac	tion	
1.	Describe and prescribe more complex types of refractive errors,	
	including postoperative refractive errors.	



2.	Describe the more advanced ophthalmic optics and optical principles of	
	refraction and retinoscopy (e.g., post keratoplasty, post-cataract	
	extraction).	
3.	Describe how to test muscle balance.	

		Psychomotor	Level of
		Domain	Competence
Optics			
Aberro	ometry Technology		
1.	Estimate the clinical incidence of higher-order aberrations.	P3	4
Diagno	ostic Equipment		
1.	Demonstrate the use of IOL calculation algorithms.		
2.	Demonstrate the use of corneal pachymetry.		
3.	Demonstrate the use of specular microscopy.		
4.	Demonstrate the use of corneal tomography with		
	anterior segment optical coherence tomography (OCT).		
5.	Demonstrate the use of topographic/elevation corneal		
	evaluation (i.e., Pentacam, Orbscan II, Galilei).		
6.	Demonstrate the use of accommodometer.	Р3	4
7.	Demonstrate the use of laser interferometry for		
	macular testing.		
Refrac	tion		
1.	Perform more advanced refraction techniques (e.g.,		
	astigmatism, complex refractions, asymmetric		
	accommodative add).		
2.	Perform objective and subjective refraction		
	techniques for more complex refractive errors,		
	including astigmatism, irregular astigmatism (e.g.,	P3	
	keratoconus, keratectasia, post corneal graft), and		
	postoperative refractive error.		
3.	•		
4.	Demonstrate the measurement of interpupillary		
	distance (IPD).		
5.	Demonstrate the prescribing of multifocal lenses.		
6.	Demonstrate the prescribing of lenses for children.		



7. Describe binocular balance.		
8. Describe how to use more advanced techniques using	Р3	4
trial lenses or the phoropter for more complex		
refractive errors, including modification and		
refinement of subjective manifest refractive error and		
more complex refractive errors (e.g., advanced and		
irregular astigmatism, vertex distance).		

# Specific Learning Outcomes Clinical Ophthalmology

## 1<sup>st</sup> Year Resident Outcomes

By the completion of their 1<sup>st</sup> year residency, the residents should be able to:

### **Cataract and Lens**

		Cognitive
		domain
1.	Describe the lens anatomy, physiology, and accommodation.	C2, C3
2.	Identify the most common causes and types of cataracts (e.g., anterior	
	polar, cortical, nuclear sclerotic, posterior sub capsular, posterior polar,	
	mature lenses such as the morgagnian cataract).	
3.	Describe the relationship between the lens and systemic disease (e.g.,	
	diabetes, myotonic dystrophy).	
4.	List ocular conditions that are associated with cataract (e.g., uveitis,	
	Wilson disease, ocular ischemia, and ocular tumors, including	
	treatment for tumors such as radiotherapy).	
5.	List systemic and topical medicine that can cause pathologic changes in	
	the lens (e.g., oral and topical corticosteroid use).	
6.	List the basic history and examination steps for preoperative cataract	
	and posterior capsular opacification evaluation.	
7.	Describe the principles and mechanisms of the following instruments in	C2, C3
	the evaluation of cataract:	
	a. Lensometer	
	b. Autorefractor	
	c. Retinoscope	
	d. Phoropter or loose lenses	
	e. Keratometer	
	f. Slit-lamp biomicroscope	
	g. Glare and contrast testing devices	



	h. Potential acuity meter	
8.	Describe the basics of IOL power estimation, including:	
	<ul> <li>a. Linear regression formulas (e.g., Sanders-Retzlaff-Kraff [SRK] and SRKII)</li> </ul>	
	<ul> <li>Theoretical eye model prediction formulas (e.g., SRKT, Holladay, and Haigis)</li> </ul>	
9.	Describe the methods to estimate axial eye length, including:	
	a. Contact ultrasound	
	b. Immersion ultrasound	
	<ul> <li>c. IOLMaster, LENSTAR, or equivalent, even if equipment is unavailable</li> </ul>	
10.	List the steps of routine intracapsular cataract extraction (ICCE), ECCE,	
	and phacoemulsification.	
11.	Define the elementary refraction techniques to obtain best-corrected	
	vision prior to considering cataract extraction.	
12.	Describe the major etiologies of dislocated or subluxated lens (e.g.,	C2, C3
	pseudo exfoliation syndrome, trauma, Marfan syndrome,	
	homocystinuria, Weill-Marchesani syndrome, syphilis).	
13.	Describe the following:	
	a. Basic ophthalmic optics as related to cataract	
	b. Types of refractive error in cataract	
	c. Retinoscopy techniques for cataract	
	d. Subjective refraction techniques for cataract patients	
14.	Describe methods to prevent postoperative infection, including	
	presurgical preparation, intraoperative antibiotics, and postoperative	
	antibiotic techniques.	
15.	Describe postoperative medications used for cataract surgery,	
	including antibiotics, non-steroidal anti-inflammatory drugs, and	
	corticosteroid therapy.	
16.	Describe the risk factors for intraoperative floppy iris syndrome (IFIS)	
	and intraoperative techniques to limit the risk of this syndrome (e.g.,	
4-	alpha blockers, use of rings, hooks)	
17.	Describe the special considerations when dealing with a unilateral cataract (trauma,	
	history of uveitis, history of topical steroid use, past surgeries)	



		Psychomotor	Level of
		domain	competency
1.	Perform basic slit-lamp biomicroscopy, retinoscopy, and ophthalmoscopy.	P2	3
2.	Classify common types of lens opacities.		
3.	Perform subjective refraction techniques and retinoscopy in patients with cataract.		
4.	Perform laser capsulotomy on routine cases of posterior capsule opacification.		
5.	Perform direct and indirect ophthalmoscopy prior to and following cataract surgery.		
6.	Perform the basic steps of cataract surgery (e.g., incision, wound closure) in the practice lab, if available.		
7.	Assist with cataract surgery.		
8.	Prepare patient for surgery including sterile draping and anesthesia.	P2	2
9.	Implement the basic preparatory procedures for cataract surgery (e.g., obtaining informed consent, identification of instruments, sterile technique, gloving and gowning, prep and drape, and other preoperative preparation).		
10.	Use the operating microscope for basic cataract surgery.		
11.	Perform some of the steps of cataract surgery under direct supervision.		

## **Contact Lenses**

	Cognitive domain
1. List advantages and disadvantages of contact lens (CL) wear.	C2, C3
2. List indications and contraindications for CL wear.	
<ol> <li>Describe a systematic and comprehensive ophthalmic examination oriented for CL fitting, including complex and</li> </ol>	
challenging cases.	





4.	Describe the various CL indications and options for each contact	
	lens type (e.g., soft CL [SCL], rigid gas permeable [RGP] CL, toric	
	CL, multifocal CL, scleral CL).	
5.	Decide which CL categories (e.g., SCL, RGP CL, hybrid CL, and	
	subgroups within each category (e.g., sphere, toric, bifocal,	
	frequent planned replacement) are best suited for a particular	
	patient.	
6.	Describe how to convey the basic CL parameters for SCL and	
	RGP CL:	
	a. Base curve	
	b. Diameter refractive power	
	c. Lens materials	
	i. Center thickness	
	ii. Peripheral curvature	
7.	Explain the concept and clinical relevance of oxygen	
	permeability (Dk) and oxygen transmissibility (Dk/center	C2, C3
	thickness).	
8.	Describe various materials used in the manufacture of CL.	
9.	Explain the optics of SCL and RGP CL:	
	a. Base curve changes	
	b. Lacrimal lens	
	c. Vertex distance	
	d. Optic zone.	
10.	Recognize the importance of obtaining central keratometry in	
	CL fitting of patients without complex needs, and explain the	
	conversion between radians and diopters.	
11.	Identify different methods of obtaining central keratometry	
	readings (e.g., manual keratometry, computerized corneal	
	topography).	
12.	Explain the importance of using diagnostic staining agents (e.g.,	
	fluorescein, lissamine green, rose Bengal) to assess corneal and	
	conjunctival staining patterns.	
13.	Describe basic tests to assess the tear film properties (e.g.,	
	Schirmer test, tear break-up time, phenol red thread tear test,	
	and meibomian gland assessment).	
14.	Describe conversion of a spectacle prescription (Rx) to a CL Rx,	
	including method of converting from plus to minus cylinder and	
- <del>-</del>	vertex distance calculations.	
15.	Describe basic steps for SCL fitting.	





16. Identify the main characteristics to be present in a CL	
prescription (eye designation, brand identification, base curve,	
diameter, and refractive power).	
17. Describe CL care guidelines to be given to the patient related to	
insertion, removal, and disinfection of CL.	
18. Describe risk factors for CL-related complications (e.g.,	
overnight wear, non-preserved saline solution usage).	
19. Describe treatment of CL-related complications (e.g., tight lens	
syndrome, overwear syndrome, giant papillary conjunctivitis,	
infectious keratitis).	

		Psychomotor/Affective	Level of
		domain	competency
1.	Perform a basic CL history.	P2	3
2.	Perform all the steps of a basic clinical		
	examination oriented for CL fitting (i.e.,		
	refraction, keratometry, visual acuity		
	assessment).		
3.	Perform a routine comprehensive slit-lamp		
	examination of the anterior segment as		
	applied to CL fitting.		
4.	Perform tear film assessment required for CL		
	patients.		
5.	Perform the techniques of retinoscopy,		
	refraction, and over-refraction in the routine		
	CL patient.		
	Perform central keratometry.		
7.	Discuss with the patient the most appropriate		
	choice for their particular clinical case.		
8.	Perform initial SCL fitting, evaluation of fit		
	(loose CL versus tight CL), and over refraction.		
	Insert and remove a trial SCL.		
10.	Instruct patients regarding safe CL insertion		
	and removal, CL wearing schedule, lens care	A2	
	regimens, CL disinfection care, indications,		
	contraindications, and possible		
	complications.		
11.	Work effectively within a medical care team.		



## **Cornea and External Diseases**

	Cognitive domain
<ol> <li>Describe the basic anatomy, embryology, physiology, pathology, microbiology, immunology, genetics, epidemiology, and pharmacology of the cornea, conjunctiva, sclera, eyelids, lacrimal apparatus, and ocular adnexa.</li> </ol>	C2, C3
<ol> <li>Describe the fundamentals of corneal optics and refraction (e.g., astigmatism, keratoconus).</li> </ol>	
<ol> <li>Describe congenital abnormalities of the cornea, sclera, and globe (e.g., Peter anomaly, microphthalmos, birth trauma, and buphthalmos).</li> </ol>	
<ol> <li>Describe characteristic corneal and conjunctival degenerations (e.g., pterygium, pinguecula, Salzmann nodular degeneration, senile plaques of the sclera).</li> </ol>	
<ol> <li>Recognize the classic corneal dystrophies (e.g., map-dot-fingerprint dystrophy, lattice, dystrophy, granular dystrophy, macular dystrophy, Fuchs dystrophy).</li> </ol>	
5. Describe the fundamentals of ocular microbiology and recognize corneal and conjunctival inflammations and infections (e.g., staphylococcal hypersensitivity, simple microbial keratitis, fungal corneal ulcers, trachoma, ophthalmia neonatorum, herpes zoster ophthalmicus, herpes simplex keratitis, adenovirus keratoconjunctivitis and conjunctivitis).	
7. Describe the basic principles of ocular pharmacology of anti-infective, anti-inflammatory and immune modulating agents (e.g., indications and contraindications for topical corticosteroids, non-steroidal anti- inflammatory agents, and antibiotics).	
<ol> <li>Manage lid margin disease (e.g., staphylococcal blepharitis, meibomian gland dysfunction).</li> </ol>	
<ol> <li>Describe the basic differential diagnosis of acute and chronic conjunctivitis or red eye (e.g., scleritis, episcleritis, conjunctivitis, orbital cellulitis, gonococcal and chlamydial conjunctivitis).</li> </ol>	
10. Manage pyogenic granuloma.	





11. Recognize the basic presentations of ocular allergy (e.g., phlyctenules, seasonal hay fever, vernal conjunctivitis, allergic and	
atopic conjunctivitis, giant papillary conjunctivitis).	
12. Comprehend the mechanisms of ocular immunology and the external	C2, C3
manifestations of anterior segment inflammation (e.g., red eye	
associated with acute and chronic iritis).	
13. Describe the symptoms, signs, testing, and evaluation for dry eye	
(e.g., Schirmer test, tarsorrhaphy); and treatment for dry eye.	
14. Describe the etiologies and treatment of superficial punctate	
keratopathy (e.g., dry eye, Thygeson superficial punctate	
keratopathy, neurotrophic keratitis, blepharitis, toxicity, ultraviolet	
photo keratopathy, contact lens-related keratitis).	
15. Enlist the etiologies of hyphema and microhyphema.	
16. Describe the basic mechanisms of traumatic and toxic injury to the	
anterior segment and treatment (e.g., chemical and thermal burns,	
lid laceration, and orbital fracture).	
17. Recognize corneal lacerations (perforating and non-perforating),	
anterior segment trauma, corneal and conjunctival foreign bodies.	
18. Describe the epidemiology, differential diagnosis, evaluation, and	
management of common benign and malignant lid lesions, including	
pigmented lesions of the conjunctiva and lid (e.g., nevi, melanoma,	
primary acquired melanosis, ocular surface squamous neoplasia).	

		Psychomotor	Level of
		domain	competence
1.	Perform external examination (illuminated and magnified) and slit-lamp biomicroscopy, including drawing of anterior segment findings.	P2	3
2.	Administer topical anesthesia, as well as special topical stains of the cornea (e.g., fluorescein dye and rose Bengal).		
3.	Perform tests for dry eye (e.g., Schirmer test, tear film breakup, and dye disappearance).		
4.	Perform punctal occlusion (temporary or permanent) or insert plugs.		
5.	Perform simple corneal sensation testing (e.g., cotton-tipped swab).		



6.	Perform tonometry (e.g., applanation, Tono-	
	Pen, Schiøtz, pneumotonometry).	
7.	Perform techniques of sampling for viral,	
	bacterial, fungal, and protozoal ocular	
	infections (e.g., corneal scraping and	
	appropriate culture techniques).	
8.	Interpret simple stains of the cornea and	
	conjunctiva (e.g., Gram stain, Giemsa stain).	
9.	Manage corneal epithelial defects (e.g.,	
	pressure patching and bandage contact	
	lenses).	
10.	D. Perform removal of a conjunctival or corneal	
	foreign body (e.g., rust ring).	
11.	1. Perform simple (non-recurrent) pterygium	
	excision (e.g., with autologous conjunctival	
	transplantation).	
12.	2. Perform an isolated lid laceration repair.	
13.	3. Perform an isolated corneal laceration repair	
	(e.g., linear laceration not extending to limbus,	
	not involving uveal or intraocular structures).	
14.	4. Perform epilation.	
15.	5. Perform a lateral tarsorrhaphy.	
16.	5. Perform incision, drainage, and/or remove a	
	primary chalazion/stye.	
17.	7. Perform a simple incisional or excisional biopsy	
	of a lid lesion.	
18.	3. Perform irrigation of chemical burn to the eye.	
19.	9. Perform Seidel test.	

# **Refractive Surgery**

	Cognitive
	domain
1. Describe simple types of refractive errors:	C2, C3
a. Myopia	
b. Hyperopia	
c. Astigmatism	
d. Presbyopia	





2.	Describe basic optic principles, such as line of sight and Purkinje	
	image.	
3.	Explain theories of accommodation.	C2, C3
4.	Describe the basics of ophthalmic optics, including how the following	
	affect the optics of the eye:	
	a. Low and high order aberrations	
	b. Corneal layers	
	c. Shape of cornea	
	d. Shape of lens	
5.	Describe basic refraction techniques using trial lenses or phoropter	
	for basic refractive errors, including:	
	a. Retinoscopy	
	b. Modification and refinement of subjective refraction	
	c. Cycloplegic retinoscopy and refraction	
	d. Postcycloplegic refraction	
6.	Describe the optical principles of common refractive surgery	
	diagnostic tools, including:	
	a. Ultrasonic pachymetry	
	b. Keratometer	
	c. Lensometer	
	d. Pupillometry	
	e. Corneal topography	
	f. Scheimpflug imaging and elevation maps	
	g. Optical coherence tomography (OCT)	
7.	Describe the following topographic maps using different scales (i.e.,	
	absolute, normalized, adjustable)	
8.	Describe normal corneal topographic patterns, as well as topographic	
	signs of keratoconus and ectasia.	
9.	Describe elevation topography maps and their importance in	
	screening refractive surgery candidates.	
10	. Describe indications and limitations of corneal topography in	
	refractive surgery.	
	. Enlist the mandatory diagnostic tests necessary for refractive surgery.	
	. Describe the basics of laser biophysics and laser tissue interaction.	
13.	. Describe the complications of high myopia, high hyperopia, and pathologies related to high astigmatism.	
14	. Define the clinical stages of keratoconus and forme fruste	
	keratoconus using clinical and topographic tests.	
15	. Describe the milestones in refractive surgery development, including	
	radial keratotomy, keratomileusis, and phakic intraocular lenses	
	(IOLs).	



16. Enlist current refractive procedures, their mechanisms of action,	
indications, and limitations, including:	
a. Types of excimer laser procedures	C2, C3
b. Phakic IOLs	
c. Implantation of intracorneal ring segments	
d. Corneal inlays	
e. Accommodative lenses	
17. Describe the main IOL calculation formulas.	
18. Describe the principles and different types (i.e., linear, rotational,	
pendular) of mechanical microkeratomes, including their	
characteristics, indications, risks, and possible complications.	
19. Describe the role of femtosecond technology in refractive surgery,	
including advantages and limitations of flap creation with a	
femtosecond laser.	
20. Describe different techniques of keratoplasty and their relation with	
refractive surgery.	

		Psychomotor/affective domain	Level of competence
1.	Perform objective and subjective refraction,	P2	3
2.	including cross cylinder and Worth 4-dot test. Diagnose refractive defects.		
	Apply different prescription formulas.		
	Prescribe spectacles for at least 20 patients with simple refractive errors (e.g., myopia, hyperopia, regular astigmatism).		
5.	Perform refraction on patients with extreme errors of refraction (e.g., 5 patients with hyperopia over 8.0 D, and 5 patients with myopia above 20.0 D).		
6.	Use the lensometer to measure spectacle power.		
7.	Use the keratometer to make corneal measurements.		
8.	Use the ultrasonic pachymeter to measure corneal thickness.		
9.	Validate corneal topography maps, including elevation topography. Recognize signs of ectatic disorders and/or candidates at risk for		



an unsatisfactory refractive surgery outcome,		
and rule out poor-quality tests (e.g., artifacts,		
alignment, and corneal exposure issues).		
10. Interpret an aberration map and evaluate its	P2	3
significance in the refractive defect of a patient,		
as well as the need to treat or not.		
11. Validate a manual refraction as a real refractive		
defect of a patient, comparing results with		
keratometers, aberrometers, and topography.		
12. Analyze tear film and tear deficiency.		
13. Unmask astigmatism from higher order		
aberrations, such as coma.		
14. Demonstrate how informed consent should be	A3	3
explained.		

## Glaucoma

		Cognitive domain
Basic S	cience	
1.	Describe the anatomy of the anterior chamber, angle, and ciliary	C2, C3
	body.	
2.	Describe the anatomy of the retinal nerve fiber layer, optic nerve	
	head, and visual pathway from the retina to the visual cortex.	
3.	Describe the mechanisms and dynamics of aqueous humor	
	inflow and outflow.	
4.	Describe the microscopic anatomy of the retina from inner to	
	outer portions, with attention to the retinal ganglion cell layer	
	and nerve fiber layer.	
5.	Describe the blood supply of the optic nerve and ciliary body.	
6.	Describe the apoptotic mechanism of retinal ganglion cell death.	
7.	Describe the physiology underlying visual-field examination and	
	its interpretation.	
8.	Describe the fundamentals of Goldmann static, kinetic	
	perimetry, and standard automated perimetry.	
9.	Describe basic principles of tonometry and aqueous outflow, and	
	applications of tonometric data (e.g., diurnal curve, peak and	
	trough values).	
Clinica	l Science	





- Describe the major features of primary open-angle glaucoma (high and low tension), angle-closure glaucoma, glaucoma suspects, and ocular hypertension.
- 2. Describe the major risk factors for primary open-angle glaucoma and angle-closure glaucoma.
- 3. Describe the steps in evaluating primary open-angle glaucoma and angle-closure glaucoma.
- Define glaucoma as a progressive neural degeneration of retinal ganglion cells, their axons and their connections to central visual centers.
- 2. Describe the features of glaucomatous optic neuropathy.
- Describe the basic features of the major glaucoma's: primary open-angle glaucoma, angle closure glaucoma, exfoliative glaucoma, and pigmentary glaucoma.
- 4. Describe the role of intraocular pressure (IOP) in the development and progression of glaucoma.
- 5. Enlist the factors that influence IOP.
- 6. Describe and understand basic principles of Goldmann applanation tonometry.
- 7. Describe tonometers (e.g., Schiøtz, Tono-Pen) and recognize artifacts of testing.
- 8. Describe principles and basic techniques of gonioscopy (3 or 4 mirror lenses) to evaluate angle structures.
- 9. Describe normal and abnormal angle findings.
- 10. Enlist risk factors other than IOP for primary open-angle glaucoma.
- 11. Classify angle-closure glaucoma (e.g., pupillary block, plateau iris, lens-related angle-closure, and malignant glaucoma).
- 12. Describe corneal pachymetry and how biomechanics and measurements of corneal thickness affect IOP interpretations.
- 13. Highlight the principles of indirect ophthalmoscopy to evaluate the optic nerve and retinal nerve fiber layer.
- 14. Describe the most common types of visual field defects in glaucoma.
- 15. Describe principles and mechanisms of medical management of glaucoma.
- 16. Describe major classes of glaucoma medications, their mechanisms of action, indications, contraindications, and side effects (topical and systemic).
- 17. Enlist drug interactions between systemic drugs and glaucoma drugs.



18. Interpret major glaucoma studies.	
19. Describe the major results of large prospective clinical trials in	
addition to those appropriate to the practice region.	
a. The Glaucoma Laser Trial (GLT)	
b. The Ocular Hypertension Treatment Study (OHTS)	
c. The Collaborative Initial Glaucoma Treatment Study	
(CIGTS)	
d. The Fluorouracil Filtering Surgery Study (FFSS)	
e. The Normal Tension Glaucoma Study (NTGS)	
f. The Advanced Glaucoma Intervention Study (AGIS)	
g. The European Glaucoma Prevention Study (EGPS)	
h. The Early Manifest Glaucoma Trial (EMGT)	

		Psychomotor/affective	Level of
		domain	competence
1.	Take a relevant patient history.	P2	3
2.	Identify the signs and symptoms of glaucoma.		
3.	Perform basic slit-lamp biomicroscopy (including peripheral anterior chamber depth evaluation, Van Herick test).		
4.	Perform basic tonometry (e.g., applanation, Schiøtz, Tono-Pen, airpuff).		
5.	Correct artifacts while performing tonometry.		
6.	Disinfect tonometer		
7.	Check calibration of Goldmann's applanation tonometer.		
8.	Perform basic gonioscopy with Goldmann- type and indentation lenses.		
9.	Evaluate angle structures, abnormalities, and appositional and synechial angle closure.		
10	. Perform central corneal pachymetry and relate to IOP findings.		
11.	Recognize the common features of the glaucomatous optic nerve including the significance of optic nerve head size, and perform stereo examination, using direct		



ophthalmoscope, fundus lens, and indirect		
lenses (i.e., 60, 66, 78, or 90 diopter lens).		
12. Recognize typical features of glaucomatous		
optic neuropathy (e.g., neuroretinal rim	P2	3
changes, disc hemorrhage, and peripapillary		
atrophy).		
13. Recognize optic nerve features of disorders		
that cause visual field loss (e.g., optic nerve		
head drusen, optic neuritis).		
14. Describe slit-lamp findings of secondary		
glaucomas (e.g., iridocorneal endothelial		
syndrome, pigment dispersion syndrome,		
exfoliation syndrome, and angle recession).		
15. Interpret visual field results for Goldmann		
kinetic perimetry and Humphrey or Octopus		
standard automated perimetry.		
16. Test for leaking filtering bleb using the Seidel		
method.		
17. Test for relative afferent pupillary defect.		
18. Recognize ocular emergencies of acute angle		
closure, and blebitis/endophthalmitis.		
19. Perform paracentesis to lower acute IOP.		

# Neuro-Ophthalmology

		Cognitive domain
1.	Describe the neuroanatomy of the visual pathways.	C2, C3
2.	Describe the anatomy and functions of cranial nerves 2-8.	
3.	Describe the anatomy of the bony orbit.	
4.	Describe the pupillary and accommodative neuroanatomy.	
5.	Describe ocular motility and related neuronal pathways.	
6.	Describe the typical features, evaluation, and management of the most common optic neuropathies (e.g., infectious, demyelinating, ischemic, inflammatory, hereditary, toxic, nutritional, compressive, and infiltrative).	
7.	Describe the typical features, evaluation, and management of the most common ocular motor neuropathies (e.g., third, fourth, sixth nerve palsy).	





<ol> <li>Describe the typical features of cavernous sinus syndrome and superior orbital fissure syndrome.</li> <li>Describe and distinguish congenital nystagmus versus acquired nystagmus.</li> <li>Describe the typical features, evaluation, and management of the most common efferent pupillary abnormalities (e.g., Horner syndrome, third nerve palsy, tonic pupil, light-near dissociation).</li> <li>Describe the typical features and evaluation of the most common visual field defects (e.g., optic nerve, optic chiasm, optic radiation, occipital cortex).</li> <li>Describe the clinical features and evaluation of ocular myasthenia gravis.</li> <li>Describe the clinical features and evaluation of carotid-cavernous fistula.</li> <li>Describe the differential diagnosis, evaluation, and management of congenital optic nerve abnormalities (e.g., optic pit, disc coloboma, papillorenal syndrome, morning glory syndrome, tilted disc, optic nerve hypoplasia, myelinated nerve fiber layer, melanocytoma, disc drusen, Bergmeister papilla).</li> <li>Describe the indications for obtaining neuroimaging studies, including computerized tomography (CT) scanning, magnetic resonance imaging (MRI), orbital ultrasonography and catheter angiography.</li> <li>Describe the signs and symptoms of giant cell arteritis and the indications for performing a temporal artery biopsy.</li> <li>Describe the clinical features, evaluation and neuro-ophthalmic aspects of thyroid ophthalmopathy.</li> <li>Describe ta systematic, sign-and-symptom-oriented neuro-ophthalmic patient interrogation (i.e., history taking) and recording techniques.</li> <li>Describe features of common headache and facial pain syndromes (e.g., migraine, trigeminal neuralgia).</li> </ol>			
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Psychomotor	Level	of
domain	competence	



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1.	Perform basic visual function tests (e.g., color	P3	3
	vision testing, Amsler grid, photostress test,		
	contrast sensitivity testing).		
2.	Perform tests of binocularity and fusion (e.g.,		
	polarized Titmus stereo test, worth 4-dot test).		
3.	Perform a basic pupillary examination.	P3	3
4.	Perform basic pharmacologic pupillary testing		
	for Horner syndrome, pharmacologic dilation,		
	and tonic pupil.		
5.	Detect a relative afferent pupillary defect.		
6.	Detect light-near dissociation.		
7.	Perform a basic assessment of ocular		
	alignment.		
8.	Use simple observational techniques (e.g.,		
	Hirschberg test, Krimsky method).		
9.	Perform basic cover/uncover testing for tropia.		
10	. Perform alternate cover testing for phoria.		
	. Perform simultaneous prism and cover testing.		
12	. Perform measurement of deviations with		
	prisms.		
13	. Describe the indications for and apply Fresnel		
	and grind-in prisms.		
14	. Describe the indications for and in a clinical		
	setting perform forced duction and forced		
	generation testing.		
15	. Perform a complete evaluation of the major		
	ocular motor systems (e.g., fixation, pursuit,		
	saccades, convergence, and vestibuloocular		
	reflex).		
16	Perform an evaluation of eyelids (e.g., assess		
	lid position, measure palpebral fissure,		
	quantify levator function).		
17	. Perform confrontational field testing (e.g.,		
	static and kinetic, central and peripheral, red		
4.0	and white targets).		
18	. Perform basic kinetic perimetry and interpret		
4.0	results.		
19	. Perform basic automated perimetry and		
	interpret results.		



<ul> <li>20. Describe the format of standard clinical tests (e.g., light stimulus, background illumination, and test points).</li> <li>21. Perform basic direct, indirect, and magnified ophthalmoscopy examination of the optic disc, macula, vessels, and periphery of the retina (e.g., recognize optic disc swelling, optic atrophy, neuroretinitis, nerve head vascular abnormalities, and macular abnormalities, such as edema, pigmentary changes, subretinal fluid, vessel abnormalities, pigmentary</li> </ul>
<ul> <li>and test points).</li> <li>21. Perform basic direct, indirect, and magnified ophthalmoscopy examination of the optic disc, macula, vessels, and periphery of the retina (e.g., recognize optic disc swelling, optic atrophy, neuroretinitis, nerve head vascular abnormalities, and macular abnormalities, such as edema, pigmentary changes, subretinal fluid, vessel abnormalities, pigmentary</li> </ul>
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abnormalities, and macular abnormalities, such as edema, pigmentary changes, subretinal fluid, vessel abnormalities, pigmentary
such as edema, pigmentary changes, subretinal fluid, vessel abnormalities, pigmentary
fluid, vessel abnormalities, pigmentary
changes).
22. Use the findings to generate a differential
diagnosis.
23. Interpret basic echography (ultrasound) of the
orbits.
24. Perform exophthalmometery.
25. Check pulse, blood pressure in both arms,
carotid bruit, and heart sounds.

# **Ophthalmic Pathology**

	Cognitive domain
<ol> <li>Describe the professional duties and specific and unique aspects of professionalism of ophthalmic pathology, and the significance of ophthalmic pathology to the practice of ophthalmology.</li> </ol>	C2, C3
<ol><li>Describe basic ocular anatomy and histology of the major structures of the eye and its adnexa</li></ol>	
<ul> <li>3. Describe basic pathophysiology of the common disease processes of the eye and its adnexa, and identify the major histologic findings: <ul> <li>a. Degeneration (e.g., pterygium, keratoconus)</li> <li>b. Dystrophy (e.g., Fuchs dystrophy, TGFBI-associated dystrophies)</li> <li>c. Infection (e.g., fungal keratitis, bacterial endophthalmitis)</li> <li>d. Inflammation (e.g., chalazion, idiopathic orbital inflammation)</li> </ul> </li> </ul>	





	e. Neoplasm and proliferation (e.g., basal and squamous cell	
	carcinoma, uveal melanoma, retinoblastoma)	
4.	Describe common methods of specimen acquisition and handling for	
	ophthalmic pathology, especially handling methods that avoid	
	artifacts and ensure representative sampling:	
	a. Surgical biopsy, with special emphasis on the eyelids and	
	conjunctiva, cornea, and vitreous	
	b. Resection margin marking	C2, C3
	c. Enucleation	
	d. Exenteration	
	e. Impression cytology	
	f. Fine needle aspiration biopsy	
5.	Describe basic information necessary to communicate to the	
	ophthalmic pathologist regarding study of these specimens.	
6.	Describe common indications for frozen sections in ophthalmic	
	pathology (e.g., complete resection margins in basal and squamous	
	cell carcinoma, demonstration of lipid in sebaceous gland	
	carcinoma).	
7.	Describe basic steps in handling and processing of gross specimens	
	in the ophthalmic pathology laboratory through a site visit, with	
	relevance to ophthalmic surgery.	

#### C. Psychomotor Domain

	Psychomotor	Level	of
	domain	competence	
1. Process specimens for submitting to an	P2	3	
ophthalmic pathology laboratory.			
2. Interpret reports from these specimens			
written by the ophthalmic pathologist.			
Participate as an observer through a site visit in the			
macroscopic and microscopic examination of			
ophthalmic pathology specimens from active cases			

# Oculoplastic Surgery and Orbit

	Cognitive
	domain
General	C2, C3



1.	Perform preoperative and postoperative assessment of patients with	
	common oculoplastic disorders.	
	Eyelid	
1.	Describe basic anatomy and physiology (e.g., orbicularis, meibomian	
	glands, Zeis glands, orbital septum, levator muscle, Müller muscle,	
	Whitnall ligament, Lockwood ligament, preaponeurotic fat, scalp,	
	face).	
2.	Describe basic mechanisms and indications for treatment of eyelid	
	trauma (lid margin sparing, lid margin involving, and canaliculus	C2, C3
	involving).	
3.	Describe mechanisms and indications for treatment of ptosis.	
4.	Describe mechanisms and indications for treatment of upper and	
	lower eyelid retraction.	
5.	Describe mechanisms and indications for treatment of entropion.	
6.	Describe mechanisms and indications for treatment of ectropion.	
7.	Identify floppy eyelid syndrome and its systemic associations.	
8.	Identify blepharospasm and hemifacial spasm.	
	Describe history and examination findings for benign and malignant	
	lid lesions.	
	Lacrimal	
1.	Describe basic anatomy and physiology (e.g., puncta, canaliculi,	
	lacrimal sac, nasolacrimal duct, endonasal anatomy, lacrimal glands).	
2.	Identify dacryocystitis.	
3.	Describe mechanisms of tearing.	
4.	Describe mechanisms and indications for treatment of congenital and	
	acquired nasolacrimal duct obstruction.	
5.	Recite the differential diagnosis of lacrimal gland mass (e.g.,	
	inflammatory, neoplastic, congenital, infectious).	
	Orbital	
1.	Describe basic anatomy (e.g., orbital bones, orbital foramina,	
	paranasal sinuses, annulus of Zinn, arterial and venous vascular	
	supply, nerves, and extraocular muscles).	
2.	Identify normal orbital and relevant nasal and paranasal sinus	
	anatomy on imaging studies (e.g., computed tomography, magnetic	
	resonance imaging).	
3.	Describe basic mechanisms and indications for treatment of orbital	
	trauma (e.g., medial wall and floor fractures, retrobulbar	
	hemorrhage).	
4.	Describe the pathophysiology of thyroid eye disease.	
5.	Recite the differential diagnosis of common orbital tumors in children	
	and adults.	



6. Recite the differential diagnosis of proptosis in children and adults.

7. Describe typical features of orbital cellulitis.

		Psychomotor / affective	Level of competency
	e . P.1	domain	
4	Eyelid	P2	3
1.	Describe indications for and perform the basic		
	office examination techniques for the most		
	common eyelid abnormalities (e.g., margin reflex		
	distance, palpebral fissure height, levator		
	function, lagophthalmos, lid crease, lid laxity		
	assessment, brow height, dermatochalasis,		
2	eversion, double eversion). Perform minor lid and conjunctival procedures		
۷.	(e.g., repair of small eyelid laceration including		
	marginal, removal of benign eyelid lesions,		
	chalazion curettage or excision, conjunctival		
	biopsy).		
3.	Treat complications of minor operating room		
	procedures (e.g., incision and drainage of		
	chalazia, excision of small eyelid lesions).		
4.	Manage trichiasis (e.g., epilation, cryotherapy,		
	surgical therapy).		
5.	Perform a temporary tarsorrhaphy.		
6.	Perform everting sutures (Quickert sutures).		
7.	Perform a lateral canthotomy/cantholysis.		
	Lacrimal		
1.	Perform the basic office examination techniques		
	for the most common lacrimal abnormalities		
	(e.g., Schirmer test, dye disappearance test,		
	punctal position, punctal dilation, canalicular		
	probing, lacrimal probing and irrigation).		
2.	Describe indications for and perform an incision		
	and drainage of the lacrimal sac.		
3.	Perform punctal plug insertion or removal.		
	Orbital		



- Describe indications for and perform the basic office examination techniques for the most common orbital abnormalities (e.g., Hertel measurement, inspection, palpation, auscultation).
  - Identify indications for and perform the basic anophthalmic socket assessment (e.g., types of implants, implant movement, socket health, socket surface, socket volume, fornices, prosthesis type and fit).

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## Pediatric Ophthalmology and Strabismus

		Cognitive domain
1.	Describe basic examination techniques for strabismus (e.g.,	C2, C3
	ductions and versions, cover and uncover testing, alternate	
	cover testing, prism cover testing).	
2.	Describe basic visual development and visual assessment of the	
	pediatric ophthalmology patient (e.g., central, steady,	
	maintained fixation), including any one matching card,	
	resolution and recognition acuity, and crowding using standard	
	vision testing (e.g., tumbling E eye chart, Allen cards, Landolt "C"	
	Broken Ring vision chart).	
3.	Describe the basic anatomy and physiology of strabismus:	
	a. Innervation of extraocular muscles	
	b. Primary, secondary, and tertiary actions	
	c. Laws governing the muscle actions	
	d. Comitant and incomitant deviations	
	e. Overaction and underaction	
	f. Restrictive and paretic saccades	
	g. Vergence	
	h. Pursuit movements	
4.	Describe basic sensory adaptations for binocular vision,	
	including:	
	a. Normal and anomalous retinal correspondence	
	b. Suppression	
	c. Horopter	
	d. Panum area	



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	e.	Fusion	
	f.	Stereopsis	
5		be pseudostrabismus.	
5. 6.		be the different etiologies of amblyopia, including:	
0.		Deprivation	
		Ametropic	
	с.	Strabismic	
	-		C2, C3
		Organic	02,00
7.		be various forms of esotropia, such as:	
		Comitant and incomitant	
	c.	Accommodative and nonaccommodative	
	d.	Decompensated	
		Sensory	
	f.	Neurogenic	
	g.	Myogenic	
	h.	Neuromuscular junction	
	i.	Restrictive	
	j.	Nystagmus and esotropia	
	k.	Spasm of the near	
	١.	Monofixation syndrome	
	m.	Consecutive	
8.	Descri	be various forms of exotropia, such as:	
	a.	Congenital	
	b.	Comitant and incomitant	
	с.	Decompensated	
	d.	Sensory	
	e.	Neurogenic	
	f.	Myogenic	
	g.	Neuromuscular junction	
	h.	Restrictive	
	i.	Basic divergence excess	
	j.	Exophoria	
-		Convergence insufficiency	
9.		be the nonsurgical treatment of strabismus and	
	-	opia, such as:	
		Patching	
	b.		
	C.	Fresnel and grind-in prism therapy	
	d.	Convergence exercises	





_	10. Describe different forms of childhood nystagmus.	
	11. Describe features, classification, and treatment indications for	
	retinopathy of prematurity.	
	12. Describe etiologies and types of pediatric cataract with	
	consideration of:	
	a. Age of onset	
	b. When do you treat and types of treatment	
	c. Postoperative rehabilitation	C2, C3
	13. Describe ocular findings in child abuse (e.g., retinal	
	hemorrhages)	
	14. Describe basic evaluation of decreased vision in infants and	
	children, such as:	
	a. Delayed maturation of vision	
	b. Leber congenital amaurosis	
	c. Other hereditary retinal disorders	
	d. Congenital glaucoma	
	e. Congenital rubella syndrome	
	f. Retinopathy of prematurity (ROP)	
	g. Various globe anomalies	
	15. Describe the symptoms, associations, findings, and treatment of	
	childhood glaucoma.	
	16. Summarize ocular embryology development (i.e., lens	
	development, fetal vasculature, anterior segment	
	development, closure of embryonic fissure).	
	17. Describe common causes of conjunctivitis in infants and	
	children in terms of symptoms, diagnosis, and treatment.	
	18. Assess subluxated and dislocated lenses and their systemic	
	associations (e.g., Marfan syndrome, homocystinuria, Weill-	
	Marchesani syndrome).	
	19. Describe management of epiphora in children, including	
	congenital nasolacrimal duct obstruction.	
	20. Describe refractive errors and spectacle correction in childhood	
	(recognizing that it is arguably the most common cause of	
	preventable visual impairment in children worldwide).	
	21. Describe accommodation and drugs used for cycloplegia.	
	22. Describe indications and uses of contact lenses in childhood.	
	23. Describe normal visual development milestones.	
	24. Describe the basic principles of genetics.	





				-	
			Psychomotor	Level	of
			domain	competence	
1.	Perform	an extraocular muscle examination based	Р3	3	
	on knov	vledge of the anatomy and physiology of			
	ocular n	notility.			
2.	Assess of	ocular motility using duction and version			
	testing.		Р3	3	
3.	Apply H	ering law and Sherrington law.			
4.	Perform	basic measurement of strabismus (e.g.,			
	Hirschbe	erg test, Krimsky method, cover testing,			
	prism c	over testing, simultaneous prism cover			
	testing,	alternate cover testing).			
5.	Perform	assessment of vision in the neonate,			
	infant, a	and child, including:			
	a. F	ixation preference test			
	b. S	itandard subjective visual acuity tests			
	c. I	nduced tropia test			
6.	Perform	cycloplegic retinoscopy in children using			
	loose le	nses, lens stick, or phoropter, depending			
	on the	age of the child and availability of the			
	devices	in the clinic.			
7.	Measur	e the refractive condition of a patient's			
	eyes usi	ng a retinoscope.			
8.	Apply in	a clinical setting the following skills in the			
	ocular n	notility examination:			
	a.	Stereoacuity testing			
	b.	Accommodative			
		convergence/accommodation ratio (e.g.,			
		heterophoria method, gradient method)			
	с.	Tests of binocularity and retinal			
		correspondence			
	d.	Cycloplegic refraction (i.e., retinoscopy)			
	e.	Anterior and posterior segment			
		examination			
	f.	Basic and advanced measurement of			
		strabismus			
	g.	Teller acuity cards			
9.	Assist a	primary surgeon in performing extraocular			
	muscles	surgery, including:			
	a.	Recession			
	ч.				



C.	Muscle weakening (e.g., tenotomy) and	
	strengthening (e.g., tuck) procedures	
d.	Transposition	
e.	Use of adjustable sutures	
f.	Intraoperative forced duction test (FDT)	
10. Probe	tear ducts to diagnose and treat an	
obstruc	ction.	
11. Medica	ally and, if indicated, surgically manage	
chalazi	ons.	
Treat molluscu	m contagiosum with curettage, if indicated	

## **Vitreoretinal Diseases**

		Cognitive
		domain
1.	Describe basic principles of retinal anatomy and physiology (i.e., basic retinal and choroidal anatomy, retinal and choroidal physiology), with emphasis on macular anatomy and physiology.	C2, C3
2.	Describe fundamentals of ancillary testing and demonstrate basic understanding of fluorescein angiography (angiographic phases), optical coherence tomography (OCT) (e.g., macular anatomy, determine pathophysiology behind structural alterations).	
3.	<ul> <li>Describe pathological anatomy, physiopathology, and clinical pictures</li> <li>of the most common vascular diseases: <ul> <li>a. Diabetic retinopathy</li> <li>b. Central vein occlusion</li> <li>c. Branch vein occlusion</li> <li>d. Arterial occlusion</li> <li>e. Hypertensive retinopathy</li> </ul> </li> </ul>	
4.	Describe features of different types of retinal detachment (i.e., rhegmatogenous, tractional, exudative).	
5.	Describe typical features of common macular diseases (e.g., age- related macular degeneration [AMD], macular whole, macular pucker, central serous chorioretinopathy, chloroquine maculopathy, pseudophakic cystoid macular edema).	
6.	<ul> <li>Describe features of traumatic pathologies, including:</li> <li>a. Commotio retinae</li> <li>b. Traumatic choroidal rupture</li> <li>c. Purtscher retinopathy</li> </ul>	



<ol> <li>Describe typical features of retinitis pigmentosa, main macular dystrophies (e.g., Stargardt, Best, cone dystrophy), and other hereditary pathologies.</li> <li>Describe basic principles of laser photocoagulation (e.g., laser response to change in power, duration, and spot size) and photodynamic therapy for retinal treatment.</li> <li>Describe basic principles, techniques, and safety of intravitreal injections.</li> <li>Manage postoperative/posttraumatic endophthalmitis.</li> </ol>	
<ul> <li>hereditary pathologies.</li> <li>8. Describe basic principles of laser photocoagulation (e.g., laser response to change in power, duration, and spot size) and photodynamic therapy for retinal treatment.</li> <li>9. Describe basic principles, techniques, and safety of intravitreal injections.</li> </ul>	7. Describe typical features of retinitis pigmentosa, main macular
<ol> <li>8. Describe basic principles of laser photocoagulation (e.g., laser response to change in power, duration, and spot size) and photodynamic therapy for retinal treatment.</li> <li>9. Describe basic principles, techniques, and safety of intravitreal injections.</li> </ol>	dystrophies (e.g., Stargardt, Best, cone dystrophy), and other
<ul><li>response to change in power, duration, and spot size) and photodynamic therapy for retinal treatment.</li><li>9. Describe basic principles, techniques, and safety of intravitreal injections.</li></ul>	hereditary pathologies.
<ul><li>photodynamic therapy for retinal treatment.</li><li>9. Describe basic principles, techniques, and safety of intravitreal injections.</li></ul>	8. Describe basic principles of laser photocoagulation (e.g., laser
<ol> <li>Describe basic principles, techniques, and safety of intravitreal injections.</li> </ol>	response to change in power, duration, and spot size) and
injections.	photodynamic therapy for retinal treatment.
-	9. Describe basic principles, techniques, and safety of intravitreal
10. Manage postoperative/posttraumatic endophthalmitis.	injections.
	10. Manage postoperative/posttraumatic endophthalmitis.

	Psychomotor domain	Level of competence
<ol> <li>Perform direct ophthalmoscopy.</li> <li>Perform indirect ophthalmoscopy.</li> <li>Perform slit-lamp biomicroscopy with precorneal lenses, 3-mirror contact lenses, or other wide-field contact lenses.</li> <li>Diagnose the presence of common retinal</li> </ol>	P2	3
4. Diagnose the presence of common retinal disorders such as exudative AMD, diabetic retinopathy, cystoid macular edema, central serous retinopathy, based on results of fundus examination, fundus photographs, OCT, and fluorescein angiography.		

## **Uveitis and Ocular Inflammation**

	Cognitive domain
1. Describe the definition and classification of intraocular	C2, C3
inflammation.	
2. Describe the basic principles of history taking:	
a. Ocular history	
i. Correlate with possible anatomical	
diagnosis (e.g., photophobia and anterior	
ii. uveitis; floaters and posterior uveitis)	
iii. Describe the onset (sudden or insidious)	



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iv.	Describe the duration (limited or persistent)			
v.	Describe the course (acute, recurrent,			
	chronic)			
vi.	Investigation and treatment history			
b. Systen				
-	Known diseases, including			
	immunosuppressed states, such as HIV,			
	malignancy,			
ii.	diabetes mellitus			
iii.	Symptoms of recent onset for (e.g., fever,			
	chills, and rigors may suggest sepsis)			
iv.	Systems review, including all medications,			
	past and current			
v.	List the clinical features of:			
vi.	Anterior uveitis			
vii.	Intermediate uveitis			
viii.	Posterior or panuveitis			
ix.	Episcleritis and scleritis (e.g., red eye,			
	blurred vision)			
х.	Anterior segment cell and flare			
xi.	Keratic precipitates (non-granulomatous or			
	granulomatous)			
xii.	Posterior synechiae			
xiii.	Vitreous cell and flare			
xiv.	Vitreous opacities			
xv.	Snowbank			
xvi.	Retinal and/or choroidal lesions			
xvii.	Retinal vasculitic			
xviii.	Retinal detachment (exudative, tractional,			
	and rhegmatogenous)			
xix.	Optic disc changes (e.g., optic disc edema,			
	optic neuritis).			
Describe the	e typical demographic features, clinical			
features, and	differential diagnosis of common, rapidly			
blinding causes for items 3a–3n (based on local				
epidemiological data). For example:				
	terior uveitis			
i.	Infectious (e.g., bacterial, viral, protozoal,			
	parasitic)			

3.





	ii.	Inflammatory (e.g., sarcoidosis, HLA B27-	
		associated, juvenile idiopathic	
	iii.	arthritis, Behçet disease, collagen vascular	
		disease)	
	iv.	Postsurgical uveitis	
	v.	Posttraumatic	
	vi.	Fuchs uveitis syndrome	
		Posner-Schlossman syndrome	
		ermediate uveitis	
		Pars planitis	
		Toxocariasis	
	iii.	Sarcoidosis	
		iv. Multiple sclerosis	
	c. Po	sterior or panuveitis	
	i.	Infectious (e.g., toxoplasmosis,	
		toxocariasis, tuberculosis, acquired and	
		congenital ocular syphilis, acute retinal	
		necrosis)	
	ii.	Inflammatory (e.g., sarcoidosis, Behçet	
		disease, Vogt-Koyanagi-Harada disease,	
		sympathetic ophthalmia)	
		Postoperative uveitis	
	iv.	Endophthalmitis (e.g., postoperative,	
		traumatic, endogenous, fungal,	
		phacoanaphylactic)	
		biscleritis and scleritis	
	i.	Collagen vascular diseases (e.g.,	
		rheumatoid arthritis, Wegener	
	ii.	granulomatosis) Infection (e.g., syphilis, tuberculosis,	
		Infection (e.g., syphilis, tuberculosis, fungal, parasitic, bacterial)	
л	Doscribo india	cations for ancillary testing in the evaluation	
4.		., fluorescein angiography [FA], indocyanine	
		ngiography, optical coherence tomography	
		ultrasonography).	
5.		cations for a tailored approach (based on	
5.		res) to laboratory investigations, including	
	obtaining tiss		
	_		
	-	ging studies (e.g., x-ray of chest, sacroiliac omputerized axial tomography [CT or CAT]	
	scan).		



6.	Describe the indications and contraindications of topical
	steroids, nonsteroidal antiinflammator drugs (NSAIDs),
	and cycloplegics.

			Psychomotor	Level	of
			domain	competence	
1.	Evalua	te clinical features of anterior uveitis,	P2	3	
	includ	ing:			
	a.	Corneal pathology (active keratitis or			
		scars, endotheliitis, band keratopathy)			
	b.	Pattern of keratic precipitates			
		(nongranulomatous, granulomatous)			
	C.	Iris changes (rubeosis iridis, gross iris atrophy)			
	d.	Anterior chamber evaluation of cells			
		and flare, including grading according			
		to standardization of uveitis			
		nomenclature (SUN) working group			
		grading system			
	e.	Differentiate episcleritis from scleritis			
2.	Descri	be the activity (active or quiescent)			
3.	Perfor	m:			
	a.	Dilated examination of the posterior			
		segment with slit-lamp biomicroscopy			
		using noncontact and contact lenses,			
		indirect ophthalmoscopy.			
	b.	Vitreous evaluation for cells and flare,			
		including grading of vitreous haze			
		according to SUN working group			
		grading system			
	С.	Retina/choroid (retinal detachment,			
		choroidal or retinal inflammation)			
	d.	Retinal vasculature (vascular			
		inflammation)			
		Optic disc (swelling, pallor)			
4.		be the regional epidemiology of uveitis			
_		late this information to the diagnosis.			
5.	Enist t	he following:			



	a. Uveitis in immunosuppresse	1
	individuals with active and recovered	1
	acquired immune	
	b. deficiency syndrome or pharmacologi	
	immunosuppression (e.g.	,
	cytomegalovirus retinitis	,
	pneumocystis (carinii) jiroveci)	
	c. Unusual infectious etiologies for uveiti	5
	(e.g., Lyme disease, West-Nile fever)	
	d. Masquerade syndromes such a	5
	vitreoretinal lymphoma	
	e. Differentiate infective from	1
	noninfective causes of uveitis.	
6.	Perform pars plana evaluation and sclera	1
	depression.	
7.	Interpret fluorescein angiography, B-sca	1
, i	ultrasonography, and correlate clinically.	
8.	Provide patient with all relevant information	1
i	about proposed ancillary testing procedure	5
t t	for uveitis, including risks and complications.	

# Ocular Oncology

		Cognitive domain
1.	Describe the basic categorization of common conjunctival and intraocular tumors.	C2, C3
2.	Describe the clinical features of the major types of ocular tumor.	
3.	Describe the symptoms and clinical manifestations indicating the presence of an ocular tumor (e.g., leukocoria, sentinel vessels).	
4.	Describe the differential diagnosis of the major tumors.	
5.	Describe the examinations and tests by which ocular tumors are diagnosed.	
6.	Describe the systemic features of ocular tumors and how these features are detected.	
7.	Describe the basic management principles of ocular tumors.	
8.	Describe the epidemiology of the more common tumors (e.g., melanoma).	



9. Describe the methods, risks, and benefits of tumor biopsy.

#### D. Technical skills

	Psychomotor domain	Level of competence
1. Perform slit-lamp and ophthalmoscopic examination of patients with an ocular tumor.	P2	3
<ol> <li>Recognize an ocular tumor and refer to an ocular oncology subspecialist.</li> </ol>		
<ol><li>Contribute to the care of patients after treatment.</li></ol>		

### Low Vision Rehabilitation

	Cognitive
	domain
1. Describe the definition, categories (types), and degrees of low vision.	C2, C3
2. Describe the most common causes of low vision (global and regional	
epidemiology and its impact on different age groups).	
3. Describe the role of the ophthalmologist in recognizing the need for	
referring patients to a low vision rehabilitation service.	
4. Describe the special aspects of vision-assessment techniques for	
children and adults with low vision (e.g., Early Treatment of Diabetic	
Retinopathy Study charts, Log MAR visual acuity chart, SOSH low	
vision chart set, LEA test eye charts).	
5. Describe significant co-morbidities that impact low vision	
rehabilitation.	
6. Describe various low vision aids.	
7. Describe the basic optics of low-vision devices.	
8. Demonstrate sensitivity to psychological and emotional aspects of	
visual impairment.	
9. Describe challenges commonly encountered by individuals with visual	
impairments.	
10. Describe how low vision impacts safety, including risk of falls, errors	
in medication, and driving accidents.	
11. Describe the importance of different visual functions, including:	
a. Visual acuity (far and near distance)	
b. Contrast sensitivity	



- c. Central and peripheral visual field
- d. Light and dark adaptation
- e. Depth perception
- f. Color vision

		Psychomotor domain	Level of competence
1.	Perform an evaluation of visual function in patients with low vision.	P2	3
2.	Describe how to use high-add reading glasses with and without a base-in (BI) prism.		
3.	Prescribe simple but appropriate rehabilitative therapies and optical devices to help the patient meet their goals (e.g., magnification, illumination).		
4.	Encourage patients with low vision to actively participate in visual rehabilitation.		
5.	Describe the functional losses of vision that may occur with various ocular diseases.		
6.	Describe the functional losses that might result from certain treatments.		
Ethics	s and Professionalism in Ophthalmology		
1.	Provide the definition and basic concepts behind the following terms used in medical ethics:		
	<ul> <li>a. Morality versus ethics (intent-based standards versus conduct-based standards)</li> </ul>		
	b. Autonomy and surrogacy		
	c. Beneficence		
	d. Nonmaleficence		
	e. Truth telling		
	f. Distributive justice		
	<ul> <li>g. Fiduciary responsibility to patients</li> <li>b. Compassion</li> </ul>		
	h. Compassion		
2.	Describe the ethical principles listed in the following key medical documents:		
	a. Hippocratic Oath		
	b. Declaration of Geneva		



c. Code of Ethics, American Academy of	
Ophthalmology	
3. Describe the basics of ophthalmic practice	
management:	
a. Partnership arrangements	
b. Contractual negotiations	
c. Hiring and supervising of employees	
d. Basic accounting	
i. Profit/loss statements	
ii. Billing	
iii. Collections	
e. Financial management	
4. Describe the basics of the health care system and	
reimbursement for services as appropriate to the	
local, regional, and national market of the trainee.	

## Community Eye Health

		Cognitive
		domain
Princip	les for the prevention of blindness	C2, C3
1.	Explain the World Health Organization (WHO) definition of blindness and low vision.	
2.	Describe the magnitude of blindness in different economic settings.	
3.	List the major causes of blindness in different economic settings.	
4.	Describe the magnitude of blindness in the resident's own country.	
5.	List the major causes of blindness in the resident's own country.	
6.	Define the concept of blind-person years.	
7.	Outline the structure of the health service, and how eye care services	
	are integrated into this structure.	
8.	Outline the social and economic implications of visual impairment and	
	the impact on quality of life.	
9.	Outline the barriers to the uptake of eye care services.	
10	. Describe the principles of primary health care and their application for	
	primary eye care.	
Inclusi	ve practice	
1.	Explain the WHO definition and conceptualization of disability.	
2.	Appraise the epidemiology of disability (including due to visual	
	impairment) and its impact in different economic settings.	





3.	Describe the intersection of blindness and visual impairment with
	other issues that may cause marginalization, including the patient's
	age, gender, other impairments, poverty, ethnic group, and faith
	community.
4.	Critically appraise the impact of disability in peoples lives (eg, poverty,
	education, quality of life [social and economic], and occupation).
5.	Describe the barriers to the uptake of eye care services within health
	systems by marginalized groups.
6.	Describe the principles of rehabilitation and community-based
	rehabilitation with relevance to people with visual impairment and the
	integration of rehabilitation within a health system.
7.	Describe strategies and partnerships with disability support services
	that can improve quality of life (eg, health, education, livelihoods,
	economic security, social inclusion) of people with long term visual
	impairment.
Catara	
	Describe the prevalence and incidence of blindness due to cataract.
2.	
3.	Describe the desired CSR required to eliminate blindness due to
	cataract.
	List the barriers to the uptake of cataract surgery.
	Outline the rationale for the monitoring of cataract services.
6.	Describe the components of a system for the monitoring of cataract
_	services.
7.	
	following cataract surgery.
	ctive error
	Define significant refractive error.
2.	Describe the prevalence of significant refractive error in children and
Э	in adults.
3.	Outline the strategy for including refractive error in a blindness prevention program, including a system for screening of school
	children to detect refractive error.
Л	List the barriers to the uptake of refractive error services.
4. Low vi	
	Define low vision.
	Describe the prevalence of low vision.
	Outline the strategy for including low vision in a blindness prevention
ى.	program.
4	List the barriers to the uptake of low vision services.
••	List the same of the uptake of four vision services.



5.	Describe the impact of low vision on the affected person and how it	
	impacts their access to wider health, education, economic, and social	
	inclusion.	
6.	List the resources available for people with low vision (eg, low-vision	
	devices, low vision training, and access to wider opportunities in	
	education, livelihoods, and social inclusion).	
Childh	ood blindness	
1.	Define childhood blindness.	
2.	Describe the prevalence of childhood blindness in different economic	
	settings.	
3.	Describe the incidence of childhood blindness.	
4.	Describe the classification of the causes of childhood blindness.	
5.	Outline the blind school survey method and the key informant method	
	for identifying the causes of childhood blindness.	
6.	Summarize the results of blind school surveys that have been	
	conducted.	
7.	List the barriers to the uptake of services for childhood eye problems.	
8.	Outline the role of primary eye care in the prevention and treatment	
	of childhood blindness.	
9.	Outline how to partner with services that can improve quality of life	
	(eg, health, education, livelihoods, and social inclusion) of children	
	with long term visual impairment.	
Tracho	oma	
1.	Describe the risk factors for trachoma.	
2.	Outline the WHO clinical grading of trachoma.	
3.	Outline the surgery, antibiotics, facial cleanliness, and environmental	
	changes (SAFE) strategy for the control of trachoma.	
4.	Describe the magnitude of trachoma, and describe the affected	
	regions.	
5.	Outline the role of primary health care in the prevention and	
	treatment of trachoma.	
Glauco	oma	
1.	Describe the prevalence of glaucoma and blindness due to glaucoma.	
Diabet	tic retinopathy	
2.	Describe the prevalence of diabetes and diabetic retinopathy.	
3.	Human resources for blindness prevention program	
4.	Describe the role and distribution of different cadres working in eye	
	care.	
_		1

5. Planning of blindness prevention programs



Describe the steps in developing a one-year operational plan for a blindness
prevention program for a health district with a population of one million
people

		Psychomotor	Level of
		domain	competence
Princip	bles of prevention of blindness		
1.	Calculate prevalence rates from given data sets.	P2	3
2.	Calculate numbers blind from given prevalence rates.		
3.	Calculate blind-person years from given data sets.		
4.	Calculate estimates of numbers of persons who are blind.		
5.	Calculate estimates of blind-person years.		
6.	Calculate an estimate of the number of		
	persons who are irreversibly blind and require		
	rehabilitation services.		
Catara	ct		
1.	Calculate an estimate of the number blind		
	due to cataract.		
2.	Calculate cataract surgery rate.		
3.	Calculate cataract surgery coverage from	53	2
	given data sets.	P2	3
4.	Calculate and comment on visual acuity		
	outcomes following cataract surgery from		
	given data sets.		
Refrac	tive error		
1.	Calculate estimates of numbers of children		
	and adults with significant refractive error.		
2.	Calculate estimates of numbers of children		
	and adults with low vision.		
Childh	ood blindness		
1.	Calculate estimates of the numbers of		
	children blind due to different causes.		



## 2<sup>nd</sup> Year Resident Goals

By the completion of their 2<sup>nd</sup> year residency, the residents should be able to:

## **Cataract and Lens**

		Cognitive domain
1.	Describe the less common causes of lens abnormalities (e.g.,	C2, C3
	spherophakia, lenticonus, ectopia lentis, coloboma).	
2.	Describe the preoperative evaluation of the cataract patient,	
	including:	
	a. Systemic diseases of interest or relevance to cataract surgery.	
	b. Systemic medication of relevance to cataract surgery (e.g.,	
	alpha 1 adrenergic blocking agent, blood thinning agents, corticosteroids)	
	c. Relationship of external and corneal diseases of relevance	
	to cataract and cataract surgery (e.g., lid abnormalities, dry eye)	
	d. Management of uveitis prior to and following cataract	
	surgery	
	e. Management of glaucoma prior to and following cataract	
	surgery, including options for postoperative intraocular	
	pressure (IOP) control	
3.	Describe glare analysis testing for cataract surgery.	
4.	Describe the use of A-scan and B-scan contact and immersion	
	ultrasonography and optical coherence techniques in cataract	
	surgery to measure axial eye length.	
5.	Describe the instruments and techniques of cataract extraction,	
	including extracapsular surgery and phacoemulsification.	
6.	Describe the important parameters of the phacoemulsification	
	machine and how to alter them for particular conditions of	
	surgery.	
7.	Describe the types, indications, and techniques of anesthesia for	
	cataract surgery (e.g., topical, local, general).	
8.	Describe indications, techniques, and complications of surgical	
	procedures, including: ECCE, ICCE, and phacoemulsification,	
_	paracentesis, and IOL placement.	
9.	Describe the pathogenesis and strategies for prevention of	
	posterior capsular opacification.	
10.	. Describe history and techniques of basic IOL implantation.	C2, C3





- 11. Correlate the level of visual acuity with the lens or capsular opacities.
- 12. Describe the pathogenesis, clinical presentation, differential diagnosis, evaluation, clinical course, treatment, and outcome of the common complications of cataract and anterior segment surgery (e.g., intraoperative floppy iris syndrome, corneal edema, IOP elevation, hyphema, endophthalmitis, toxic anterior segment syndrome (TASS), cystoid macular edema (CME), retinal detachment, IOL dislocation, lens-induced glaucoma, and uveitis).
- 13. Describe the indications for, principles of, and techniques of yttrium aluminium garnet (YAG) laser capsulotomy, and understand the proper timing of YAG laser capsulotomy.
- Describe advanced IOL power calculation (e.g., after radial keratotomy [RK], myopic laser-assisted in situ keratomileusis [LASIK]/photorefractive keratectomy [PRK], hyperopic LASIK/PRK).
- 15. Describe the properties of different ophthalmic viscoelastic devices (OVDs) (e.g., dispersive, cohesive, adaptive) and the advantages and disadvantages for certain phases of surgery.
- 16. Describe the fluid dynamics in phacoemulsification, including the difference between peristaltic and venture pump types.
- Manage common postoperative complications of cataract surgery (e.g., endophthalmitis, toxic anterior segment syndrome, elevated IOP, CME, wound leak, uveitis, and capsular block syndrome).
- 18. Define the more complex indications for cataract surgery (e.g., better view of posterior segment, lens-induced glaucoma).
- 19. Describe the techniques to manage a small pupil, including mechanical manipulation, management of iris membrane, iris hooks, viscoelastic, and phaco techniques.
- 20. Describe techniques to diagnose and operate on patients with posterior polar cataract.
- 21. Describe the preoperative preparations for surgery and special intraoperative considerations for patients with uveitis.
- 22. Describe techniques for prevention of capsular opacification and phimosis (before, during, after surgery), including the use of capsular tension rings and IOL factors.



		Psychomotor	Level of
		domain	competence
1.	Perform local injections of corticosteroids, antibiotics, and anesthetics, including	P3	4
	retrobulbar and subtenon.		
2.	Perform extracapsular surgery in a practice		
	setting (e.g., animal or practice lab).		
3.	Practice surgery in the operating room under		3
	supervision, including mastery of the cataract	P2	
	surgical skills:		
2.	Perform paracentesis of the anterior chamber.		
3.	Implement advanced applications of		
	viscoelastics in surgery (e.g., control of iris		3
	prolapses, elevation of dropped nucleus,		
	viscodissection, aspiration of residual/retained		
	viscoelastic, soft-shell technique).		

## **Contact Lenses**

		Cognitive domain
1.	Explain applied anatomy and physiology (e.g., corneal	C2, C3
	metabolism and temperature, oxygen consumption, stromal	
	acidosis, tear osmolality, tissue fragility, cell apoptosis, corneal	
	sensitivity, closed eyelid-related ocular surface repercussions).	
2.	Recognize signs and symptoms of CL intolerance and over wear.	
3.	Explain the importance of assessing tear film and ocular surface	
	condition with more complex auxiliary tests in certain CL fitting	
	situations (e.g., tear film osmolality and biochemical	
	composition, impression cytology).	
1.	Identify CL fitting situations requiring corneal topography (e.g.,	
	computerized/Placido rings).	
2.	Explain the rationale underlying different topography profiles	
	and how these relate to the manifest refraction.	
3.	Analyze topography maps.	
4.	Explain physical properties of CL materials:	





a. International Organization for Standardization (ISO) classification	C2, C3
5. Explain advantages and disadvantages of SCL materials.	
<ol> <li>Explain advantages and disadvantages of SEE materials.</li> <li>Explain advantages and disadvantages of RGP CL materials.</li> </ol>	
7. Explain RGP/SCL geometry relation with corneal geometry (i.e.,	
lacrimal meniscus, refraction, and ocular surface implications).	
8. Explain main principles to fit RGP CL (e.g., first trial CL choice,	
fluorescein patterns, alignment, movement, wearing and	
replacement schedule, fitting motivation, and follow up).	
9. Explain main principles to fit toric SCL:	
a. Stabilization	
i. LARS rule (i.e., Left Add, Right Subtract)	
ii. Movement	
iii. Rotation	
iv. Possible refitting needs	
10. Appraise clinical situations best suited for RGP CL fitting versus	
toric SCL fitting.	
11. Explain when CL refitting is indicated and perform refitting	
when needed.	
<ol> <li>Recognize signs and symptoms of a tight, optimal, and loose CL fitting.</li> </ol>	
13. Explain advantages and disadvantages of different wearing	
schedules (e.g., conventional, frequent planned replacement,	
flexible, daily).	
14. Describe ocular impact and physiological needs regarding	
different CL wearing schedules.	
15. Describe CL requirements for materials needed for	
extended/flexible CL wearing.	
2. Explain patient and CL selection and fitting techniques as	
applied to fit presbyopia.	
3. Explain how to keep a CL fitting trial set (i.e., CL, equipment, and	
disinfection care).	
4. Describe and evaluate different CL care systems.	
5. Explain the clinical importance of CL environment (i.e., CL	
patient surrounding, ocular surface, and storage case).	

Psychomotor	Level	of
domain	competence	





1.	Perform a CL history in patients requiring	P3	3
	more complex CL fitting (e.g., subclinical		
	ectatic corneal disorders such as		
	keratoconus and pellucid marginal		
	degeneration, regular moderate		
	astigmatism, presbyopia, ocular surface		
	disease, and post-refractive surgery).		
2.			
Ζ.	Perform a clinical examination, including		
	retinoscopy and refraction techniques to		
	verify and inspect CL in patients requiring		
	more complex CL fitting (e.g., subclinical		
	ectatic corneal disorders such as		
	keratoconus and pellucid marginal		
	degeneration, regular moderate		
	astigmatism, presbyopia, ocular surface		
	disease, and post-refractive surgery).		
3.	Indicate more complex additional		
	auxiliary tests (e.g., computer-based		
	corneal topography, tear film osmolarity,		
	impression cytology) in patients requiring		
	more complex CL fitting (e.g., subclinical		
	ectatic corneal disorders such as		
	keratoconus, pellucid marginal		
	degeneration, regular moderate		
	astigmatism, presbyopia, and ocular		
	surface disease, and post refractive		
	surgery).		
4.	Perform RGP CL fitting (spherical).		
5.	Perform SCL toric fitting.		
6.	Perform presbyopia CL fitting.		
7.	Perform appropriate CL selection and		
	material or parameters modification in CL		
	refit.		
8.	Perform CL verification for visual acuity,		
0.	fitting, and comfort in patients requiring		
	more complex CL fitting.		
9.	Educate patients regarding CL-related		
9.	complications.		
10	•		
10.	Manage CL-related complications.		
			85   P a g e



11. Perform the skills needed for long-term	3
management and follow up of CL	
patients.	

## **Cornea and External Diseases**

		Cognitive domain
1.	Describe the more complex anatomy, embryology, physiology,	C2,C3
	pathology, microbiology, immunology, genetics, epidemiology, and	
	pharmacology of the cornea, conjunctiva, sclera, eyelids, lacrimal	
	apparatus, and ocular adnexa.	
2.	Describe the more complex congenital abnormalities of the cornea,	
	sclera, anterior segment and globe and their associated systemic	
	manifestations (e.g., Axenfeld, Rieger, and Peter anomalies,	
	aniridia, hamartomas and choristomas).	
3.	Correlate the concordance of the visual acuity with the density of	
	media opacity (e.g., cataract, corneal scars, edema), and evaluate	
	the etiology of discordance between acuity and findings from	
	examination of the media.	
4.	Recognize and treat less common corneal or conjunctival	
	presentations of degenerations and common conjunctival	
	neoplasms (e.g., inflamed, atypical, or recurrent pterygium, band	
	keratopathy, benign and malignant tumors).	
5.	Describe the epidemiology, clinical features, pathology, evaluation,	
	and treatment of peripheral corneal thinning disorders or	
	ulceration (e.g., Terrien marginal degeneration, Mooren ulcer,	
	rheumatoid arthritis-related corneal melt, dellen).	
6.	Describe the epidemiology, differential diagnosis, evaluation, and	
	management of vitamin A deficiency (e.g., Bitot spot, dry eye,	
	slowed dark adaptation) and neurotrophic corneal diseases.	
7.	Manage recurrent corneal erosions.	
8.	Manage chronic conjunctivitis (e.g., chlamydia, trachoma,	
	molluscum contagiosum, Parinaud oculoglandular syndrome,	
	ocular rosacea).	
9.	Describe the more complex principles of ocular pharmacology of	
	anti-infective, anti-inflammatory, and immune modulating agents	
	(e.g., use of topical non-steroidal and steroidal agents,	
	cyclosporine, and anti-tumor necrosis factor agents).	





<ul> <li>10. Describe more complex differential diagnosis of red eye (e.g., autoimmune and inflammatory disorders causing scleritis, episcleritis, conjunctivitis, orbital cellulitis).</li> <li>11. Describe key features of trachoma, including epidemiology, clinical features, staging, and its complications (e.g., cicatrization), prevention (e.g., facial hygiene), and topical and systemic antibiotic treatment (especially in hyperendemic regions), and surgery (e.g., tarsal rotation).</li> <li>12. Manage interstitial keratitis (e.g., syphilis, viral diseases, and noninfectious, immunologic, inflammation).</li> <li>13. Describe the differential diagnosis and the external manifestations of more complex anterior segment inflammation (e.g., acute and chronic iritis with and without systemic disease).</li> <li>14. Manage the ocular complications of severe diseases, such as chronic exposure keratopathy, contact dermatitis, and rosacea.</li> <li>15. Describe the classification, pathology, indications for surgery, and prognosis of common eyelid abnormalities (e.g., blepharoptosis, trichiasis, distichiasis, essential blepharospasm, entropion, ectropion) and understand their relationship to secondary diseases of the cornea and conjunctiva (e.g., exposure keratopathy).</li> <li>16. Manage foreign body, animal, and plant substance injuries and understand the risk of injury with organic material.</li> <li>17. Describe more complex mechanisms of traumatic and toxic injury to the anterior segment (e.g., long-term sequelae of acid and alkali burn, complex lid laceration involving the lacrimal system, full-thickness laceration).</li> <li>18. Manage corneal lacerations (perforating and nonperforating).</li> <li>19. Manage hyphema (e.g., surgical indications, evacuation).</li> <li>20. Recognize the anterior segment manifestations of systemic diseases (e.g., Wilson disease) and pharmacologic side effects (e.g., amiodarone vortex keratopathy).</li> <li>21. Manage common and uncommon benign and malignant lid lesions.</li> </ul>		
<ul> <li>episcleritis, conjunctivitis, orbital cellulitis).</li> <li>11. Describe key features of trachoma, including epidemiology, clinical features, staging, and its complications (e.g., cicatrization), prevention (e.g., facial hygiene), and topical and systemic antibiotic treatment (especially in hyperendemic regions), and surgery (e.g., tarsal rotation).</li> <li>12. Manage interstitial keratitis (e.g., syphilis, viral diseases, and noninfectious, immunologic, inflammation).</li> <li>13. Describe the differential diagnosis and the external manifestations of more complex anterior segment inflammation (e.g., acute and chronic iritis with and without systemic disease).</li> <li>14. Manage the ocular complications of severe diseases, such as chronic exposure keratopathy, contact dermatitis, and rosacea.</li> <li>15. Describe the classification, pathology, indications for surgery, and prognosis of common eyelid abnormalities (e.g., blepharoptosis, trichiasis, distichiasis, essential blepharospasm, entropion, ectropion) and understand their relationship to secondary diseases of the cornea and conjunctiva (e.g., exposure keratopathy).</li> <li>16. Manage foreign body, animal, and plant substance injuries and understand the risk of injury with organic material.</li> <li>17. Describe more complex mechanisms of traumatic and toxic injury to the anterior segment (e.g., long-term sequelae of acid and alkali burn, complex lid laceration involving the lacrimal system, full-thickness laceration).</li> <li>18. Manage corneal lacerations (perforating and nonperforating).</li> <li>19. Manage toreal lacerations and nonperforating).</li> <li>10. Recognize the anterior segment manifestations of systemic diseases (e.g., Wilson disease) and pharmacologic side effects (e.g., amiodarone vortex keratopathy).</li> </ul>	10. Describe more complex differential diagnosis of red eye (e.g.,	C2,C3
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21. Manage common and uncommon benign and malignant lid lesions.	amiodarone vortex keratopathy).	
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	Psychomotor	Level	of
	domain	competence	
1. Perform more advanced techniques, including	P2	3	
keratometry, keratoscopy, endothelial cell			
count and/or evaluation, specular			
microscopy, and pachymetry.			



2.	Perform more complex and recurrent	
	pterygium excision, including conjunctival	
	grafting.	
3.	Perform more complex lid laceration repair.	
4.	Perform more complex corneal laceration	
	repair (e.g., stellate perforating laceration).	
5.	Perform stains of the cornea and conjunctiva	
	(e.g., calcofluor white, acid fast).	
6.	Repair simple lacerations of the lacrimal	
	drainage apparatus (e.g., perform intubations	
	and primary closure).	
7.	Treat hyphaema and microhyphaema with	
	associated increased intraocular pressure	
	and/or blood staining (e.g., surgical	
	evacuation).	

# **Refractive Surgery**

		Cognitive domain
1.	Describe various types of refractive errors.	C2,C3
2.	Define the possible corrective solutions for all types of refractive errors.	
3.	Describe basic diagnostic tools used in refractive surgery, including topography, pachymetry, and biometry.	
4.	Describe more complex types of refractive errors, including postoperative refractive errors following cataract surgery, keratoplasty, refractive surgeries, ectatic conditions, and irregular astigmatism.	
5.	Explain basics of wavefront analysis, including ray tracing and dynamic skiascopy, and graphical representation of wavefront errors, including corneal and entire eye high-order aberration maps, point-spread function, and modulation-transfer function.	
6.	Describe the basics of Zernike polynomials and Fourier analysis.	
7.	Use different topographic maps and scales for different purposes (e.g., screening, postoperative evaluation, detection of complications).	
8.	Describe the basics of measuring contrast sensitivity.	



9. Describe laser-tissue interaction Munnerlyn formula.
10. Describe corneal biomechanics, including biomechanical responses to keratorefractive surgery, corneal healing after excimer laser procedures, corneal hysteresis, and corneal resistance factor.
11. Identify post laser in-situ keratomileusis (LASIK) ectasia.
12. Describe the mechanism of action, indications, advantages, and potential complications of mitomycin C application in surface ablation.
13. Describe the effect of corneal crosslinking on the biomechanical properties of the cornea, including its indications and how it can be combined with other refractive surgery procedures.

	Psychomotor	Competence
	Domain	Level
<ol> <li>Perform refraction techniques using trial lenses or phoropter for basic and more complex cases, including:         <ul> <li>a. Modification and refinement of subjective manifest refractive error</li> <li>b. Cycloplegic retinoscopy and refraction</li> <li>c. Post cycloplegic refraction</li> <li>d. Contact lens use</li> <li>e. Irregular astigmatism</li> </ul> </li> </ol>	Р1	3
<ul> <li>f. Post-keratoplasty</li> <li>g. Refractive surgery cases</li> <li>2. Apply the basics of optics and optical principles of refraction and retinoscopy in the clinical setting, including higher order aberrations.</li> </ul>	P2	3
<ul> <li>3. Gather accurate information essential for preoperative evaluation of patients seeking refractive surgery, including: <ul> <li>a. Medical interview</li> <li>i. Patient expectation</li> <li>ii. Social history</li> <li>iii. Medical history</li> <li>iv. Pertinent ocular history</li> <li>b. Physical examination</li> </ul> </li> </ul>	P1	4



	i. Uncorrected visual acuity		
	ii. manifest and cycloplegic visual		
	acuity		
	iii. Intraocular pressure		
	iv. Slit-lamp examination		
	v. Fundus examination		
4. Diagnose	and manage dry eye prior to surgery.	P2	3
5. Use th		P2	3
	nents in more complex patients (e.g.,		
·	neal surgery or corneal disease), and		
	results with corneal topography maps,		
	ity, and quality of vision.		
	refractive instruments and techniques	P2	3
. –	to refractor, pachymetry, automated		
	opography, aberrometer, pupillometry,		
	ens refraction, OCT, corneal hysteresis,		
	eal resistance factor) in the clinical		
-	r refractive surgery patients.	D4	2
	developing patient care management	P1	2
-	simple refractive errors (e.g., myopia, a, regular astigmatism), and define the		
	benefits for each procedure.		
	various types of refractive surgery,	P2	2
including		Γ <b>Δ</b>	2
	venty surface ablation procedures		
	venty LASIK procedures		
с. Т			
	plantation procedures		
	en phakic IOL surgeries		

## Glaucoma

Α.	Cognitive Skills	
1.	Describe epidemiology of congenital glaucoma, primary open-	C2
	angle glaucoma, exfoliation syndrome and exfoliative	
	glaucoma, and angle-closure glaucoma.	
2.	Describe the genetics of:	C2
	a. Primary congenital glaucoma (CYP1B1)	



b. Syndromes associated with congenital/developmenta	
glaucoma	
i. Lowe syndrome	
ii Nail-patella syndrome	
iii. Aniridia (PAX 6)	
iv. Axenfeld-Rieger syndrome (PITX2, FOXC1, FKHL7)	
c. Primary open-angle glaucoma	
i. GLC1A and the molecular biology of myocilin	
ii. Optineurin	
iii. Other genes as they become identified	C2
3. Describe the features of primary infantile and juvenile	
glaucomas.	C2
4. Describe etiologies and major risk factors for secondary open-	
angle glaucomas.	C3
5. Recognize secondary glaucomas (e.g., angle recession,	
inflammatory, steroid induced, pigmentary, and exfoliative,	
phacolytic, neovascular, postoperative, malignant, lens	
particle glaucomas, plateau iris, glaucomatocyclitic crisis, and	
iridocorneal endothelial syndrome) with attention to	
appropriate pathophysiology.	С3
6. Manage complex secondary glaucomas (e.g., exfoliation,	
angle recession, inflammatory, steroid induced, pigmentary,	
phacolytic, neovascular, postoperative, malignant, lens-	
particle glaucomas; plateau iris; glaucomatocyclitic crisis;	
iridocorneal endothelial syndromes; aqueous	
misdirection/ciliary block).	C2
7. Describe diurnal fluctuations in IOP and ocular perfusion	
pressure and their application in the approach to therapy.	C2
8. Describe more advanced optic nerve and nerve fiber layer	
anatomy in glaucoma and typical and atypical features	
associated with glaucomatous cupping (e.g., rim pallor, disc	
hemorrhage, parapapillary atrophy, rim thinning, notching,	
circumlinear vessels, central acuity loss, hemianopic or other	
nonglaucomatous types of visual field loss).	C2
9. Describe tools and techniques for quantitative anterior	
segment imaging such as ultrasound biomicroscopy and	
anterior segment optical coherence tomography (OCT).	
10. Describe basic principles of tools to analyze optic nerve and	
retinal nerve fiber layer such as OCT, Heidelberg Retina	C2
Tomograph (HRT), and GDx.	



11. Interpret HRT, OCT, and GDx scans.	
12. Interpret more advanced forms of perimetry (kinetic and	C2
automated static), including various perimetry strategies such	C2
as threshold testing, suprathreshold testing, and special	
algorithms.	
13. Describe the principals involved in determining glaucomatous	
progression both clinically and perimetrically.	C2
14. Describe the principles, and more advanced anatomic	
gonioscopic features of primary and secondary glaucomas	C2
(e.g., plateau iris, appositional closure).	
15. Describe target IOP and its use in glaucoma management.	
16. Describe the principles of medical management of more	C2
advanced glaucomas (e.g., advanced primary open-angle	C2
glaucoma, secondary open and closed angle glaucomas, and	-
normal tension glaucoma).	
17. Describe pitfalls of medical treatment, in particular poor	
compliance and adherence.	C2
18. Describe the features of angle-closure glaucomas and	
aqueous misdirection.	C2
19. Describe the most common clinical features and etiologies of	
ocular hypotonic.	C2
20. Describe differential diagnosis and management of hypotony.	
21. Apply the results of major clinical trials in glaucoma to clinical	C2
practice (e.g., GLT, OHTS, CIGTS, FFSS, NTGS, AGIS, EGPS, and	C2
EMGT).	
22. Apply specific medical treatments in more advanced	
glaucoma.	C3
23. Describe the principles, indications, and techniques of various	
types of laser energy, spot size, and laser wavelengths.	C2
24. Describe the principles, indications, and techniques of	C2
trabeculectomy (with or without cataract surgery, with or	
without antimetabolites), glaucoma drainage devices, and	
cyclodestructive procedures.	
25. Describe the major etiologies of dislocated or subluxated lens	
associated with glaucoma (e.g., trauma, Marfan syndrome,	C2
homocystinuria, Weill-Marchesani syndrome, syphilis).	
26. Describe the less common causes of lens abnormalities	
associated with glaucoma (e.g., spherophakia, lenticonus, and	
ectopia lentis).	
27. Define the relationships of glaucoma and uveitis.	C1





28. Describe diagnostic accuracy, false positive and false negative	
diagnoses and their significance at individual and societal	
levels, differences between case-based and community-based	C2
screening, including an understanding of sensitivity and	
specificity, number needed to treat, <i>t</i> tests, life-table analysis,	
prospective versus retrospective studies, case control and	
cohort studies.	

1.	Perform argon and selective laser	P1	2
	trabeculoplasty for open-angle glaucoma.		
2.	Perform argon or YAG laser for angle-closure	P1	2
	glaucoma.		
3.	Perform surgical peripheral iridoectomy for	P1	2
	angle-closure glaucoma.		
4.	Perform peripheral iridoplasty for non-pupillary	P1	2
	block angle-closure glaucoma.		
5.	Perform laser suture lysis.	P1	2
6.	Perform cyclodestructive surgery	P1	2
	(photocoagulation or cryotherapy).		
7.	Assist with trabeculectomy and glaucoma	P1	2
	drainage device surgery in the operating room.		
8.	Describe and manage a flat anterior chamber.	P1	2
9.	Perform routine trabeculectomy.	P1	1

## Neuro-Ophthalmology

Α.		Cognitive	e Skills						
	1.	Describe	typical	and	atypical	features,	evaluation,	and	C2
	management of the most common optic neuropathies (e.g.,								





	papilledema, optic neuritis, ischemic, inflammatory, infectious, infiltrative, compressive, hereditary optic neuropathies).	
2.	Describe features, evaluation, and management of the more	C2
	complex supranuclear and internuclear palsies (e.g., progressive	
	supranuclear palsy and subtle internuclear ophthalmoplegia, one-and-half syndrome).	C2
3.	List the common causes of an acute versus chronic isolated	
	ocular motor neuropathy and define general management of	
	each.	C2
4.	List the common causes of cavernous sinus syndrome and	
5.	superior orbital fissure syndrome. Differentiate among different forms of acquired nystagmus	С3
0.	(e.g., downbeat, upbeat, pendular, gaze evoked, rebound,	
	convergence, retraction).	C2
6.	List the different mechanism causing nonphysiologic anisocoria	
	and describe characteristics features and evaluation of the less common disorders (e.g., mixed sympathetic and	
	parasympathetic denervation of iris, aberrant regeneration in	<b>2</b> 2
	third nerve palsy, pharmacologic miosis).	C2
7.	List mechanism and causes of central versus peripheral light	
	near dissociation (e.g., Argyll-Robertson pupil, diabetic neuropathy, tonic pupil,	
	Parinaud syndrome).	C1
8.	Describe features and evaluation of the less commonly	-
	encountered visual field defects (e.g., sectoranopia,	C1
9.	checkerboard, monocular temporal crescent). Describe more advanced aspects of visual field testing	
-	indications, selection, and interpretation (e.g., artifacts of	
	automated perimetry, testing, and thresholding strategies).	C2
10	Describe neuro-ophthalmic aspects of common systemic	
	diseases (e.g., hypertension, diabetes, thyroid disease, myasthenia gravis, temporal arteritis, sarcoidosis, systemic	
	infections, and inflammation).	C2
11.	Describe neuro-ophthalmic findings that are common following	
	head trauma (e.g., traumatic optic neuropathy, bilateral fourth	<b>C</b> 2
12	nerve palsy, traumatic brain injury). Describe evaluation and management of inherited neuro-	C2
	ophthalmic diseases (e.g., Leber hereditary optic neuropathy,	
	autosomal dominant optic atrophy, spinocerebellar	
	degenerations).	C2





13. Describe evaluation and management of ocular myasthenia	
gravis.	C3
14. Recognize common pathologic findings of brain and orbits on CT	
and MRI related to neuro-ophthalmology.	<b>2</b> 2
15. Describe the typical features, evaluation, and management of	C2
urgent neuro-ophthalmic pathologies (e.g., giant cell arteritis,	
cavernous sinus thrombosis, orbital apex syndrome, pituitary	
apoplexy).	

1.	Perform a detailed cranial nerve evaluation	P1	3
	other than the oculomotor nerve evaluation		
	(e.g., trigeminal, and facial and acoustic nerve		
	function).		
2.	Interpretneuro-radiologic images (e.g.,	P2	3
	indications and interpretation of orbital		
	tumors, thyroid eye disease, pituitary		
	adenoma, optic nerve glioma, optic nerve		
	sheath meningioma).		
3.	Perform the evaluation, management, and	P3	3
	specific testing (e.g., stereopsis, mirror test,		
	red-green testing, monocular prism test) of		
	patients with "functional" (i.e., nonorganic		
	visual loss (e.g., recognize nonorganic spiral or		
	tunnel visual fields).		
4.	Interpret basic ocular coherence tomography	P1	3
	(OCT) imaging of the eye (e.g., optic disc,		
	retinal nerve fiber layer, and macula).		
5.	interpret basic ocular electrophysiology (e.g.,	P1	3
	visually-evoked potential [VEP],		
	electroretinogram [ERG], electrooculogram		
	[EOG]).		
6.	Perform basic neurologic screening	P2	3
	examination (e.g., tandem walk, sensory		
	examination, cerebellar function testing, basic		
	cognitive evaluation).		
		P1	3



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7. 8.	Identify patients with "functional" visual loss (i.e., nonorganic visual loss) and provide appropriate approach and follow up. Quantify relative afferent pupillary defect (RAPD) with neutral density filter and be able	P2	3
9.	to detect RAPD in patients with only one working pupil. Interpret fluorescein angiography images.	P1	3

# **Ophthalmic Pathology**

Α.	Cognitive Skills	
1.	Describe more advanced ocular anatomy (e.g., common variants), and identify the histology of the major structures of the eye and its adnexa relevant to specific clinical rotation(s) (e.g., oculoplastics, cornea, glaucoma, retina, ophthalmic oncology).	C2
2.	Identify the major histologic findings of common diseases of the eye (e.g., keratitis, exfoliation syndrome, corneal and retinal dystrophies and degenerations, frequent neoplasms) relevant to specific clinical rotation(s) (e.g., oculoplastics, cornea, glaucoma, retina, ophthalmic oncology).	C2
3.	Describe the pathophysiology and histology of potentially vision or life-threatening diseases (e.g., temporal arteritis, endophthalmitis, retinoblastoma, ocular melanoma, extraocular or orbital spread of an intraocular or periorbital tumor, metastasis to the eye and orbit) relevant to specific clinical rotation(s) (e.g., oculoplastics, cornea, glaucoma, retina, and ophthalmic oncology).	C2

1. Process appropriately more advanced specimens	P1	3
for submitting to an ophthalmic pathology		
laboratory, including writing of the accompanying		
letter to the ophthalmic pathologist (e.g.,		
impression cytology, fine needle aspiration		



	biopsy,	vitreous	biopsy,	evisceration,		
	exenterati	ion specimer	ı).			
2.	Perform a	biopsy for fro	ozen section	study in ocular	P1	1
	pathology					
3.	Participate	e under supe	ervision thro	ough a site visit	P1	2
	in a macro	oscopic and r	nicroscopic	examination of		
	ophthalmi	ic specimens	from active	cases, working		
	from low t	to high powe	r.			

# **Oculoplastic Surgery and Orbit**

Eyelid	
1. Describe more advanced eyelid anatomy and physiology (e.	g., <b>C2</b>
lymphatics).	
2. Describe the mechanisms of and indications for eyel	id <b>C2</b>
reconstruction.	
3. Described the genetics (where known), clinical feature	es, <b>C2</b>
evaluation, and treatment of congenital eyelid deformition	es
(e.g., coloboma, distichiasis, epicanthus, telecanthu	
blepharophimosis, ankyloblepharon, epiblepharo	
euryblepharon, cryptophthalmia, Goldenhar syndrom	е,
Treacher-Collins syndrome, Waardenburg syndrome).	
4. Describe clinical features, evaluation, syndromic associatio	
and management of congenital ptosis (e.g., simpl	
blepharophimosis-ptosis-epicanthus inversus syndron	le
[BPES], jaw wink, congenital fibrosis).	
5. Describe the genetics (when applicable), clinical feature	
evaluation, and treatment of acquired myogenic ptosis (e.g oculopharyngeal muscular dystrophy, mitochondri	
myopathies, myotonic dystrophy, myasthenia gravis).	ai
6. Describe the clinical features, evaluation, and treatment	of <b>C2</b>
acquired neurogenic ptosis (e.g., third nerve palsy, Horn	
syndrome).	
7. Describe the mechanisms and indications for treatment	of
more advanced eyelid trauma (e.g., chemical burns, therm	
burns, canthal avulsions, eyelid avulsions).	C2
8. Describe features, evaluation, and treatment of presept	
cellulitis versus orbital cellulitis.	
	C2



	Γ
Lacrimal	
1. Describe more advanced lacrimal anatomy and physiology	C2
(e.g., lacrimal pump theories).	
2. Describe the mechanisms and indications for treatment of	C2
more advanced lacrimal trauma (e.g., nasolacrimal duct	
obstructions resulting from facial fractures).	
3. Describe features, evaluation, and treatment of more	C2
complicated cases of nasolacrimal duct obstruction,	
canaliculitis, dacryocystitis, and acute and chronic	
dacryoadenitis.	
4. Describe the genetics, clinical features, evaluation, and	C2
management of lacrimal dysgenesis.	

Orbita	1	
1.	Describe more advanced orbital anatomy and physiology (e.g.,	C2
	vascular anatomy, neural anatomy, orbital septa).	
2.	Describe the clinical features, evaluation, and management of	C2
	congenital orbital deformities (e.g., anophthalmia,	
	microphthalmia, hypotelorism, hypertelorism versus	
	telecanthus).	
3.	Describe the genetics, clinical features, evaluation, and	C2
	management of common craniosynostoses and other	
	congenital malformations (e.g., Crouzon syndrome, Apert	
	syndrome).	
4.	Describe the mechanisms and indications for treatment of more	C2
	advanced orbital trauma (e.g., zygomaticomaxillary complex	
	fractures, naso-orbital ethmoid fractures, Le Fort fractures).	
5.	Describe management plan for thyroid ophthalmopathy (e.g.,	C2
	epidemiology, symptoms and signs, associated systemic	
	diseases, orbital imaging, differential diagnosis, surgical,	
	medical, and radiation indications, side effects of treatment).	
6.	Describe Management nonspecific orbital inflammation (e.g.,	
	symptoms and signs, orbital imaging, differential diagnosis,	C2
	biopsy indications, choice of treatments).	



Eyelids

1.	Perform more advanced examination techniques	P2	3
	for less common eyelid abnormalities (e.g.,		
	decreased blink, orbicularis weakness, contour		
	abnormalities, marginal entropion).		
2.	Perform more complicated minor lid procedures	P2	3
	(e.g., larger benign skin lesions, recurrent chalazia).		
3.	Perform more complicated eyelid surgery (e.g.,	P2	2
	upper blepharoplasty, lower lid tightening).		
4.	Perform more advanced eyelid reconstruction (e.g.,	P2	2
	wedge/pentagonal block resection).		
5.	Identify indications for and complications of, and	P2	3
	treat blepharospasm and hemifacial spasm.		
6.	Identify histopathological features of common	P2	3
	eyelid conditions.		

### Lacrimal

1.	Perform more advanced lacrimal assessment (e.g.,	P2	2
	interpretation of dye testing, canalicular probing in		
	trauma).		
2.	Perform basic lacrimal procedures (e.g., lacrimal	P2	2
	drainage testing [irrigation, Jones Dye Tests 1 and		
	2], lacrimal probing, lacrimal intubation, incision		
	and drainage of lacrimal sac abscess).		
3.	Identify indications for and interpret lacrimal	P2	3
	imaging (e.g., scintigraphy, cystography).		
4.	Identify histopathological features of common	P2	3
	lacrimal conditions.		

Orbit



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1.	Perform more advanced assessment of the orbit	P2	2
	(e.g., hypoglobus, facial asymmetry, enophthalmos,		
	and proptosis).		
2.	Perform enucleation and evisceration.	P2	3
3.	Perform more advanced socket assessment (e.g.,	P2	2
	extrusion of implants, anophthalmic socket		
	complications).		
4.	Identify common orbital pathology (e.g., orbital	P1	3
	fractures, orbital tumors) on imaging studies (e.g.,		
	magnetic resonance imaging, computed		
	tomography, ultrasound).		
5.	Treat common presentations of orbital cellulitis.	P2	3
6.	Identify histopathological features of common	P1	3
	orbital conditions.		

# Pediatric Ophthalmology and Strabismus

1. Describe basic and more advanced strabismus examination	C2
techniques (e.g., combined vertical and horizontal prism cover	
testing, double Maddox rod testing).	
2. Describe basic and more advanced visual development and	C2
visual assessment of the pediatric ophthalmology patient (e.g.,	
blink to light or threat, measures of fixation and following	
behavior, objective measures of visual acuity) using the	
optokinetic nystagmus (OKN) drum to assess fixation and	
electrophysiological techniques such as sweep visual evoked	
potential (VEP) evaluation.	
3. Describe basics of binocular sensory testing (e.g., Titmus stereo	C2
testing, Randot stereo testing, Worth 4-dot test, Bagolini lenses).	
4. Describe etiologies, evaluation, and management of vertical	C2
strabismus, including:	
a. Neurogenic	
b. Myogenic	
c. Neuromuscular junction	
d. Oblique overaction or underaction	
e. Dissociated vertical deviation	
f. Restrictive	
	C2





5.	Describe various strabismus patterns (e.g., A or V pattern) and	
	associations with various types of comitant strabismus; the	
	anatomic role of muscle pulleys; and the potential role of	
	radiology in assessing complex strabismus.	C2
6.	Describe common hereditary or congenital ocular motility or lid	
	syndromes (e.g., Duane syndrome, Marcus Gunn jaw-winking	
	syndrome, Brown syndrome).	C2
7.	Describe typical features of retinoblastoma (e.g., differential	
	diagnosis, evaluation, treatment indications, and types).	C2
8.	Describe basic evaluation and differential diagnosis of decreased	
	vision in infants and children (e.g., retinal and optic nerve	
	etiologies, amblyopia).	C2
9.	Describe recognizable causes of blindness in infants (e.g.,	
	albinism, optic nerve hypoplasia, achromatopsia, Leber	
	congenital amaurosis, retinal dystrophy, congenital optic	
	atrophy) and appropriate work up and associated diseases.	C2
10.	Describe cortical visual impairment and periventricular	
-	leukomalacia.	C2
11.	Interpret diplopia charts (e.g., Hess charts, Lees chart, and Harms	-
	screen).	C2
12	Evaluate a child with congenital blindness, including VEP and	
	interpretation of an electroretinogram (ERG).	

<ol> <li>Perform more advanced strabismus testing, such a Parks-Bielschowsky 3-step test, Lancaster red-green test, Maddox rod testing, double Maddox roo testing, and measurement of dissociated vertica deviation (DVD).</li> </ol>	n d	3
<ol><li>Perform forced duction test (FDT) and force generation test (FGT) in the clinic.</li></ol>	e <b>P2</b>	2
<ol> <li>Perform basic extraocular muscle surgery, and exercise surgical judgment for the indications and contraindications for strabismus surgery.</li> </ol>		2
<ol> <li>Perform preoperative extraocular muscle surger assessment, intraoperative techniques, and describe intraoperative and postoperative complications of strabismus surgery.</li> </ol>	d	3
<ol> <li>Perform the following strabismus surgeries:</li> <li>a. Recession</li> </ol>	P2	2

b. Resection



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### Vitreoretinal Diseases

1.	Descri	be more advanced retinal anatomy and physiology.	C2
2.	Descri	be more advanced ancillary testing concepts of fluorescein	C2
	and in	docyanine green (ICG) angiography as applied to retinal	
	vascul	ar and other diseases (e.g., indications, basic differential	
	diagno	sis based on angiographic patterns).	
3.	Descri	be the fundamentals of retinal electrophysiology and basic	C2
	ophtha	almic echography.	
4.	Descri	be the findings of major studies in vascular retinal diseases,	C2
	includi	ng the following:	
	a.	Diabetic retinopathy	
	b.	Early Treatment Diabetic Retinopathy Study (ETDRS)	
	с.	Diabetes Control and Complications Trial (DCCT)	
	d.	United Kingdom Prospective Diabetes Study (UKPDS)	
	e.	Diabetic Retinopathy Clinical Research Network (DRCRnet)	
		Trials	
	f.	Central vein occlusion	
	g.	Central Vein Occlusion Study (CVOS)	
	h.	Standard Care vs. Corticosteroid for Retinal Vein Occlusion	
		(SCORE)	
	i.	Global Evaluation of implaNtable dExamethasone in retinal	
	j.	Vein occlusion with macular edemA (GENEVA) Study Group	
	k.	Central Retinal Vein Occlusion (CRUISE) Study	
	I.	Branch vein occlusion	
	m.	Branch Vein Occlusion Study (BVOS)	
	n.		
		(SCORE)	
	0.	GENEVA Study Group	
	р.	BRAnch Retinal Vein Occlusion (BRAVO) Trial	
	q.	Retinopathy of prematurity	
	r.	Cryotherapy for Retinopathy of Prematurity (CRYO-ROP)	
	s.	Early Treatment for Retinopathy of Prematurity (ETROP)	
5.		be management plan for peripheral retinal diseases and	
	vitreou	us pathologies (e.g., vitreous hemorrhage, posterior vitreous	C2





	detachment, retinal tears, giant retinal tears, lattice degeneration with atrophic holes).	
6.	Describe the techniques for retinal detachment repair, including	C2
	indications, mechanics, instruments, basic techniques, and surgical	
	adjuvants, including heavy liquids, expandable gases, and silicone	
	oil for the following:	
	a. Pneumatic retinopexy	
	b. Scleral buckling	
	c. Vitrectomy	
7.	Describe typical features of less common macular diseases:	C2
	a. Myopic maculopathy	
	b. Serous retinal detachment secondary to optic disc pit	
	c. Ocular histoplasmosis syndrome	
	d. Phenothiazine/tamoxifen toxicity	
	e. Diagnose, evaluate, treat, and classify open and closed	
	globe trauma (e.g., Birmingham Eye Trauma Terminology	
	System).	
8.	Describe management plan for postoperative/posttraumatic	C2
	choroidal detachments and sympathetic ophthalmia.	
9.		C2
9.	Describe the indications/complications of laser treatment for	
	diabetic retinopathy (e.g., panretinal photocoagulation, macular	
	grid).	

1.	Perform indirect ophthalmoscopy with scleral	P1	2
	indentation.		
2.	Perform ophthalmoscopic examination with contact	P2	2
	lenses, including panfunduscopic lenses.		
3.	Interpret fluorescein and indocyanine green (ICG)	P1	3
	angiography and correlate findings with differential		
	diagnosis.		
4.	Diagnose the presence of pigment granules in the	P2	2
	anterior vitreous (i.e., Shafer sign) during a retinal		
	detachment or retinal break.		
5.	Interpret retinal imaging technology (e.g., OCT,	P2	3
	retinal thickness analysis).		
6.	Perform posterior segment photocoagulation.	P2	2
7.	Perform diabetic focal/grid macular laser treatment.	P2	2



8. Perform peripheral scatter photocoagulation	P2	2
(panretinal).		
9. Perform laser retinopexy (demarcation) for isolated	P2	2
retinal breaks.		
10. Interpret basic echographic patterns (e.g.,	P2	3
rhegmatogenous retinal detachment, tractional		
retinal detachment, posterior vitreous detachment,		
choroidal detachment, intraocular foreign body).		
11. Perform fundus drawings of the retina, showing	P2	3
vitreoretinal relationships and findings.		
12. Perform (or assist during) cryotherapy of retinal	P2	2
holes and other pathology.		
13. Perform (or assist during) vitreous tap and	P2	3
intravitreal antibiotic injections for the treatment of		
endophthalmitis.		
14. Perform subtenon injections of triamcinolone	P2	3
acetonide for the treatment of macular edema.	12	5
	P2	3
15. Perform intravitreal injection of anti-vascular	PZ	5
endothelial growth factor (VEGF) drugs for the		
treatment of AMD.		

## **Uveitis and Ocular Inflammation**

1.	Describe the pathophysiology of intraocular inflammation.	C2
2.	Describe the principles of history taking of patients with uveitis according to SUN.	C2
3.	Describe the importance of being guided by clinical findings from	C2
	the ocular examination and taking a more specific history in	
	order to generate a list of differential diagnoses.	
4.	Describe more advanced principles of examination of patients	C2
	with uveitis and differential diagnoses of the clinical signs:	
	a. Anterior segment (e.g., iris nodules, pupillary membrane,	
	peripheral anterior synechiae, iris bombe)	
	b. Posterior segment (e.g., pars plana signs of inflammation	
	[snowballs], retinal detachment, retinal vasculitis, optic	
	swelling [differentiate optic neuritis from	
	hyperemia], macula [macular edema])	



5.	Describe the regional epidemiology of uveitis	
6.	Describe the typical demographic feature, clinical features, and	
	differential diagnosis of:	C2
	a. Common uveitis in immunosuppressed individuals (e.g.,	C2
	cytomegalovirus retinitis, endogenous endophthalmitis)	
	b. Masquerade syndromes such as vitreoretinal lymphoma	
7.	Differentiate serious infective from noninfective causes of	
	uveitis. (e.g., recognize an endogenous endophthalmitis and	
	differentiate this from an immune-mediated uveitis, such as	C2
	Behçet disease).	
8.	Describe angiographic features of retinitis, choroiditis, and	
	vasculitis.	
9.	Describe the B-scan features of certain retinal, choroidal, and	C2
	scleral diseases.	
10.	Describe the OCT features of macular edema.	C2
11.	Describe the common complications of common uveitis	
	syndromes (e.g., intraocular pressure elevation, cataract, band	C2
	keratopathy, macular edema).	C2
12.	Describe indications and contraindications for corticosteroid	
	treatment of uveitis (e.g., topical, local, systemic), including risks	
	and benefits of therapy.	C2
13.	Describe the management of common uveitic syndromes.	

1. Perform a more advanced examination of the	P2	3
anterior and posterior segment in addition to that		
described for Year 1.		
a. Anterior segment (e.g., iris nodules,		
pupillary membrane, peripheral anterior		
synechiae, iris bombe)		
b. Posterior segment (e.g., pars plana signs		
of inflammation [snowballs], retinal		
detachment, retinal vasculitis, optic		
swelling [differentiate optic neuritis from		
hyperemia], macula [macular edema])		
2. Recognize and evaluate the typical demographic		
features, clinical features, and differential	P2	3
diagnosis of common, rapidly blinding causes of		



<ul> <li>uveitis (based on local epidemiological data), as described in the curriculum of Year 1.</li> <li>a. Administer topical steroids, NSAIDs, and cycloplegics in the treatment of uveitis.</li> <li>b. Interpret the results of ancillary tests (e.g., fluorescein angiography, OCT, B-scan ultrasonography) for diagnosis.</li> <li>3. Perform a major investigational work up (e.g., P2 3</li> </ul>	
<ul> <li>a. Administer topical steroids, NSAIDs, and cycloplegics in the treatment of uveitis.</li> <li>b. Interpret the results of ancillary tests (e.g., fluorescein angiography, OCT, B-scan ultrasonography) for diagnosis.</li> </ul>	
cycloplegics in the treatment of uveitis. b. Interpret the results of ancillary tests (e.g., fluorescein angiography, OCT, B-scan ultrasonography) for diagnosis.	
<ul> <li>b. Interpret the results of ancillary tests (e.g., fluorescein angiography, OCT, B-scan ultrasonography) for diagnosis.</li> </ul>	
fluorescein angiography, OCT, B-scan ultrasonography) for diagnosis.	
ultrasonography) for diagnosis.	
3. Perform a major investigational work up (e.g., P2 3	
laboratory testing, radiologic testing) according to	
epidemiologic data, history, and clinical	
examination.	
4. Evaluate uveitis associated with P2 3	
immunosuppressed individuals (e.g., active and	
recovered acquired immune deficiency	
syndrome, pharmacologic immunosuppression).	
5. Perform posterior subtenon or transseptal P2 3	
injection of corticosteroids.	
6. Administer oral corticosteroids in the treatmentP23	
of uveitis.	
7. Manage side effects of immunosuppressiveP23	
therapy.	
8. Perform an anterior chamber and vitreous tap for <b>P2 3</b>	
diagnostic purposes	
9. Administer intravitreal injection antibiotics in P2 3	
cases of bacterial endophthalmitis.	

# Ocular Oncology

1. Describe the classification of ocular tumors (i.e., conjunctival and intraocular).	C2
2. Describe the clinical features of ocular tumors and their secondary effects.	C2
3. List the differential diagnosis for each of the ocular tumors.	C2
4. Describe diagnostic techniques for ocular tumors (e.g., examination under anesthesia for pediatric tumors, imaging, biopsy, laboratory tests, and oncology referral).	C2





5. Describe indications (e.g., biopsy for lymphoma) and contraindications (e.g., biopsy for retinoblastoma) for the	
various diagnostic techniques.	
6. Describe the management options for ocular tumors with	C2
indications and contraindications for each form of	
management.	C2
7. Describe the complications of ocular therapy and their	
management.	C2
8. Describe basic histopathology of tumors, including	C2
immunohistochemistry.	
9. Describe the prognosis of the different types of ocular tumor.	C2
10. Describe the epidemiology of the more common tumors (e.g.,	
melanoma).	
11. Describe the methods, risks, and benefits of tumor biopsy.	

	examination (e.	g., to	P2	3
recognize oculodermal	melanosis).			
2. Perform palpation of ce	2. Perform palpation of cervical lymph nodes.		P2	3
3. Perform slit-lamp exam	nination, gonioscop	oy, and	P2	3
indirect ophthalmoscop	y to diagnose and I	ocalize		
ocular tumors.				
4. Perform transillumination	on for intraocular tu	umors.	P2	3
5. Perform B-scan ultrasc	onography to dete	ct and	P2	3
measure intraocular tur	nors.			
6. Perform sequential ex	amination to asse	ess the	P2	3
tumor over time (e.g., a	typical nevus).			
7. Plan to evaluate for	systemic disease	(e.g.,	P2	3
metastases, primary tur	nor, syndromes).			
8. Provide short-term and	long-term postop	erative	P2	3
care to patients w	ith an ocular	tumor,		
collaborating with a	subspecialist and	other		
health care workers as a	appropriate.			
9. Manage ocular complica	ations as appropriat	e (e.g.,	P2	3
radiation retinopathy,	macular edema, ca	itaract,		
glaucoma).				
10. Interpret the results of	laboratory investig	gations	P2	3
and adjust managemen	t accordingly.			
			P2	3



11. Council about prognosis and various management	
options to patients and their families in a detailed,	
ethical, and compassionate manner.	

# **Refractive Surgery**

1.	Describe and diagnose various types of refractive problems,	C2
	including irregular astigmatism, and identify the best solution for	
	each.	
2.	Describe the most complex types of refractive errors, including	C2
	postoperative refractive errors, post keratoplasty, and refractive	
	surgery.	
3.	Describe the most advanced optics and optical principles of	C2
	refraction and retinoscopy, including higher-order aberrations.	
4.	List the indications for and interpret preoperative and	C2
	postoperative diagnostic testing, including:	
	a. Corneal topography	
	b. Wavefront analysis	
	c. Pachymetry	
	d. Calculation of stromal-bed thickness before and after	
	LASIK	
_	e. Aspheric profile of ablation	
5.	Formulate informed diagnostic and therapeutic decisions based	C2
	on patient information, current scientific evidence, clinical	
6	judgment, and patient expectations.	
6.	Describe accommodative and non-accommodative treatments of	C2
	presbyopia, including:	
	a. Monovision	
	b. Excimer laser correction	
	c. Conductive keratoplasty	
	d. Corneal inlays	
	<ul><li>e. Accommodating IOLs</li><li>f. Multifocal IOLs</li></ul>	
7.	Describe the advanced formulas for IOL calculation in extreme	C2
7.		
	myopia, hyperopia, and after corneal refractive surgery.	
		1





8.	Describe the basics of modulation transfer function (MTF), point	C2
	speed function (PSF) and Strehl ratio as objective ways to measure	
	quality of vision.	
9.	Describe the basics of topography-guided, wavefront-guided,	
	and optimized ablations as compared to standard ablations.	

1.	Perform basic refractive surgery procedures, such	P2	2
	as low myopia or low hyperopia with LASIK		
	(microkeratome) and surface ablation (LASIK or		
	photorefractive keratectomy [PRK]).		
2.	Perform the most advanced objective and	P2	3
	subjective refraction techniques using trial lenses or		
	the phoropter, including:		
	a. Contact lens refraction for more complex		
	refractive errors, including modification		
	b. and refinement of subjective manifest		
	refractive error		
	c. Cycloplegic retinoscopy and refraction		
	d. Postcycloplegic refraction		
	e. Irregular astigmatism		
	f. Postkeratoplasty		
	g. Refractive surgery cases		
3.	Utilize the most advanced optics and optical	P2	3
	principles for refraction and retinoscopy, including		
	higher order aberrations.		
4.	Utilize the keratometer for detection of subtle or	P2	3
	complex advanced corneal refractive errors.		
5.	•		
	instruments and techniques (e.g., corneal	P2	3
	topography, pupillometry, aberrometry,		
-	Scheimpflug imaging, OCT).		
6.	Fit contact lenses in patients with irregular corneas,		
	irregular astigmatism, and following refractive	P2	2
_	surgery.		
7.	Assist in advanced refractive surgeries, including		
	topography-guided ablation, wavefrontguided	P1	2
	ablation, and combined refractive surgeries.		





8.	Encourage patients to actively participate in their	P1	3
	own care by providing disease and treatment		
	information, and counsel patients on how to		
	prevent postoperative injury.		
9.	Correct refractive error after surgeries, such as	P2	3
	penetrating keratoplasty, deep anterior lamellar		
	keratoplasty, and radial keratotomy		

# Low Vision Rehabilitation

## A. Cognitive Skills

		Γ
1.	Describe clinical applications, indications, and limitations of the various low vision aids (e.g., electronic and optical magnification, large print, Braille, computers with artificial speech, text to speech).	C2
2.	Describe the more advanced optics of low vision devices.	C2
3.	Describe visual acuity and visual field evaluation methods for different levels of disability.	C2
4.	Describe the evaluation of and rationale for licensing automobile drivers who are visually impaired, and explain the local licensing regulations.	C2

### B. Technical skills

1.	Prescribe more complex rehabilitative therapies and optical devices to help the patient meet their goals.	P2	3
2.	Perform evaluation of vision assessment in licensing drivers who are visually impaired.	P2	3
3.	Demonstrate low vision devices and educate low vision patients on the uses and limitations of these devices.	Ρ2	3

# Ethics and Professionalism in Ophthalmology



1.	Describe basic medical ethics in the ophthalmic practice,	C2
	including:	
	a. Confidentiality of health information	
	b. Professional competence and maintenance of	
	competence c. Informed consent	
	d. Responsibility to report the unethical conduct of others	
	e. Adequate patient assessment and avoidance of	
	under/over treatment and under/over testing	
2.	Describe elements of effective physician-patient communication,	C2
2.	including:	02
	a. Relevant cultural and linguistic differences that potentially	
	influence ethical delivery of services	
3.	Describe advanced aspects of practice management (e.g.,	C2
	business models, documentation requirements and coding,	
	privacy requirements, accommodating patients or employees	
	with disabilities).	
4.	Describe advanced aspects of health care reimbursement (e.g.,	C2
	physicians' role in managed care organizations, administrative	
	role, third-party reimbursement, capitated programs).	
5.	Describe the framework of patient-care quality as it relates to	C2
	patient safety, patient advocacy, effectiveness, efficiency,	
6	timeliness, and equity.	
6.	Describe how ophthalmologists are responsible for ensuring that	<u></u>
	all those in the service area of the practice have access to affordable eye care.	C2
7.	Describe the various missions of ophthalmology organizations	
7.	with respect to service to members, patients, clinical education,	C2
	and quality of care.	
8.	Describe how participation of ophthalmologists in ophthalmology	
_	organizations serves the profession and society.	C2
9.	Describe the responsibilities of ophthalmologists and	
	ophthalmology societies to ensure that everyone has the right to	C2
	sight.	

# Community Eye Health





Enlist t	he Principles for the prevention of blindness	C2
1.	Outline the magnitude and distribution of global blindness.	
2.	List the major causes of global blindness.	
3.	Describe primary, secondary, and tertiary prevention strategies	
	that are applicable to the leading causes of low vision and	
	blindness.	
4.	Outline the different possible approaches (ie, disease	
	orientated, service orientated, strategy orientated, community	
	orientated) to blindness prevention.	
5.	Describe the integrated approach to blindness prevention that	
	is recommended for use in VISION 2020.	
6.	Describe the structure and function of a generic VISION 2020	
	program for a health service unit with a population of one	
	million.	
7.	In line with the WHO Universal Eye Health: A Global Action Plan	
	2014–2019, describe strategies to strengthen inclusive practices	
	related to gender, disability, and other groups within a generic	
	VISION 2020 program.	

### Cataract

1.	Describe the prevalence and incidence of blindness due to	C2
	cataract in different economic settings.	
2.	Describe the cataract surgery rates in different economic	C2
	settings.	
3.	Describe cataract surgery coverage, including its use and	C2
	limitations as an indicator to measure program output.	
4.	Outline the possible strategies to overcome the barriers to	C2
	cataract surgery.	
5.	Define cataract surgery efficiency and cataract surgery volume.	C2
6.	Outline the factors affecting cataract surgery capacity.	C2
7.	Outline the principles of an efficient cataract surgical service.	C2
8.	Describe a model for the staffing and running of a cataract	C2
	surgical unit.	
9.	Describe the components of a model for the costing of cataract	C2
	surgery.	





10. Describe the possible strategies for cataract surgery cost	C2
containment.	
11. Describe the possible strategies for cataract surgery cost	
recovery.	

### **Refractive error**

1.	Describe the prevalence of refractive error in different	C2
	countries/regions.	
2.	Outline the possible strategies for the provision of spectacles in	C2
	a blindness prevention program.	

### Low vision

1.	Describe	the	prevalence	of	low	vision	in	different	C2
	countries/	'regior	ıs.						
2.	Outline th	ie pos	sible strategie	es fo	r the	provisior	n of	low-vision	C2
	aids in a b	lindne	ss prevention	pro	gram.				

### Childhood blindness

1.	List the main causes of childhood blindness in different	C2
	socioeconomic settings.	
2.	Describe the primary, secondary, and tertiary prevention	
	strategies for the control of childhood blindness due to corneal	C2
	scar, cataract, glaucoma, and retinopathy of prematurity.	
3.	Describe the main barriers for children with visual disabilities to	
	access health, education, and social inclusion.	C2
4.	Outline the models/strategies for supporting education for	
	children with visual impairments through mainstream schools	C2
	(eg, inclusive education) or "special" schools.	

### Glaucoma

1.	Describe the prevalence of glaucoma in different regions and in	C2
	different race groups.	
2.	Outline the possible strategies for the opportunistic case	C2
	detection of glaucoma.	
3.	Describe the advantages and disadvantages of medical, laser,	C2
	and surgical interventions for the management of glaucoma in	
	middle and low-income countries.	



4. Define glaucoma treatment/surgery rate.	C2
5. Outline the possible strategies for increasing the glaucoma	C2
follow-up rate.	

### Diabetic retinopathy

1.	Outline the possible strategies for the prevention of diabetic	C2
	retinopathy, including the use of appropriate educational health	
	materials for counseling.	
2.	Outline the possible strategies for screening for diabetic	C2
	retinopathy.	
3.	Outline the possible strategies for the treatment of diabetic	C2
	retinopathy.	
4.	Outline the possible strategies for increasing the diabetic	C2
	retinopathy follow-up rate.	

### Human resources for blindness prevention programs

1.	Describe the recommended cadres and numbers of human	C2
	resources required at the community level, primary level,	
	secondary level, and tertiary level for a generic blindness	
	prevention program for a health service unit of one million in the	
	resident's own country or health district.	
2.	Describe the roles of each of the cadres that are recommended	C2
	for a generic blindness prevention program.	
3.	Describe the available training facilities for a generic blindness	C2
	prevention program.	

## Infrastructure for blindness prevention programs

1.	From the International Agency for the Prevention of Blindness	C2
	(IAPB) standard list for VISION 2020, describe the recommended	
	instruments and equipment required at the primary, secondary,	
	and tertiary level for a generic blindness prevention program for	
	a health service unit of one million population.	
2.	Outline the strategies for the maintenance of the recommended	C2
	instruments and equipment.	

### Planning of blindness prevention programs



1. Describe the potential role of a VISION 2020 coordinator and a	C2
VISION 2020 committee.	

### Principles of blindness prevention

1. For planning purposes, integrate primary, secondary, and	P1	3
tertiary preventions for leading causes of low vision and		
blindness into a district blindness prevention program plan		
adhering to inclusive practices.		

### Cataract

1.	For planning purposes, calculate estimates of numbers of people blind due to cataract in different countries and regions.	P1	3
2.	For planning purposes, calculate cataract surgery rate in different countries and regions.	P1	3
3.	For planning purposes, identify and suitable strategies for overcoming the barriers to cataract surgery in a blindness prevention program.	Ρ	3
4.	For planning purposes, identify and include suitable strategies for improving the efficiency of a cataract surgical unit in a blindness prevention program.	P1	3

#### **Refractive error**

Calculate estimates of numbers of children and adults with significant refractive error in different countries and regions.	P2	3
For planning purposes, identify and include suitable strategies for including refractive error as a priority in a blindness prevention program.	P2	3

Low vision





1. Calculate estimates of numbers of children and adults with	P2	3
low vision in different countries and regions.		
2. For planning purposes, identify and include suitable	P2	3
strategies for including low vision as a priority in a blindness		
prevention program.		

### Childhood blindness

1. For planning purposes, use available program reports to	P2	3
identify key gaps in and barriers to service delivery.		

#### Trachoma

1. For planning purposes, use available program reports to	P2	3	
identify key gaps in and barriers to service delivery.			

### Glaucoma

1. Calculate estimates of numbers of people with glaucoma in	P2	3
different countries and regions.		
2. For planning purposes, identify suitable strategies for including glaucoma as a priority disease in a blindness	P2	3
prevention program.		

### Diabetic retinopathy

1. Calculate estimates of numbers of people with diabetic	P2	3
retinopathy in different countries and regions.		
2. For planning purposes, identify suitable strategies for	P2	3
including diabetic retinopathy as a priority disease in a		
blindness prevention program.		

#### Human resources

1. For planning purposes, identify suitable strategies	for P2	3
improving the human resource capacity in a blindr	ness	
prevention program.		

#### Infrastructure



1. For planning purposes, identify suitable strategies for	P2	3
improving the infrastructure capacity in a blindness		
prevention program.		

### Planning of blindness prevention programs

1. Develop activities plan for a one-year operational plan for a	P2	3
blindness prevention program for a health district with a		
population of one million.		

# 3<sup>rd</sup> Year Resident Goals

By the completion of their 3<sup>rd</sup> year residency, the residents should be able to:

### **Cataract and Lens**

1.	Describe the principles, indications for, mechanics of of contact	C2
	and immersion A-scan ultrasonography and calculation of IOL	
	power.	
2.	Describe the complications of more advanced anterior segment	
	surgery (e.g., pseudoexfoliation, small pupils, intraoperative	C2
	floppy iris syndrome, mature cataract, hard nucleus,	
	posttraumatic, zonular dehiscence, cataract surgery after pars	
	plana vitrectomy, short eye, corneal endothelial diseases).	
3.	Describe the use of special devices for cataract surgery in	
	complex situations such as specialized IOLs, capsular tension	C2
	rings and segments, iris hooks, use of indocyanine green/trypan	
	blue staining of the anterior capsule.	
4.	Describe IOL fixation options in the lack of capsular support for	
	in the bag fixation (anterior chamber [AC] IOL, sulcus fixation +/-	C2
	optic capture, iris fixation, and scleral fixation).	
5.	Describe the indications for, techniques of, and complications of	
	cataract extraction in the context of the subspecialty disciplines	C2
	of the following:	
	a. Glaucoma (e.g., combined cataract and glaucoma	
	procedures, glaucoma in cataractous eyes, cataract	
	surgery in patients with prior glaucoma surgery)	
	a. Retina (e.g., cataract surgery in patients with scleral	
	buckles or prior vitrectomy)	



<ul> <li>b. Cornea (e.g., cataract extraction in patients with corneal opacities) and the use of fiber optic for better visualization</li> </ul>	
<ul> <li>Ophthalmic plastic surgery (e.g., ptosis following cataract surgery)</li> </ul>	
<ul> <li>d. Refractive surgery (e.g., cataract surgery in eyes that have undergone refractive surgery)</li> </ul>	
6. List indications for and techniques of intracapsular surgery (e.g.,	
rare cases may requiremthis procedure, or patients may have	C2
had the procedure performed previously).	
7. Describe instrumentation and techniques used to implant	
foldable and non-foldable IOLs.	C2
8. Describe the evaluation and management of common and	
uncommon causes of postoperative endophthalmitis and TASS.	C2
9. Describe the causes and indication for performing, repositioning,	
removal, or exchange of IOLs.	C2
10. Describe the government and hospital regulations that apply to	
cataract surgery.	C2
11. Describe the indication and option for astigmatism management	
during cataract surgery (e.g., on axis incision, limbal relaxing	C2
incisions [LRI], opposite clear corneal incision [OCCI], toric IOL).	
12. Describe the use of corneal topography and wave front analysis	
to help select the best type of IOL for a patient especially	C2
following keratorefractive surgery.	
13. Describe the option for presbyopic correction solutions during	
cataract surgery (e.g., monovision. multifocal IOLs,	
accommodative IOLs, dual optic IOLs). 14. Describe the mechanisms of actions, indications,	<b>C</b> 2
14. Describe the mechanisms of actions, indications, contraindications, advantages, and disadvantages of premium	C2
IOLs (e.g., multifocal, accommodative, toric, aspheric, blue	
blocker, intraocular miniature telescope).	C2
15. Describe evaluation and management of IOL complications (e.g.,	
intraoperative damage to IOL, postoperative IOL opacification,	
dislocation, sublocation).	
16. Describe the advantages and disadvantages of the materials	C2
used for IOL fabrication (e.g., poly-methylmethacrylate [PMMA],	
silicone, hydrophobic acrylic, hydrophilic acrylic).	
17. Describe lens/IOL surgery solutions for myopia and hyperopia	C2
(e.g., refractive lens exchange, phakic IOLs).	



1. Assist in the teaching and supervision of basic and	P2	2
standard level learners.		
2. Perform phacoemulsification in a practice setting		2
(e.g., animal or practice lab) and then in the		
operating room, ideally 50-100 cases of a		
combination of phacoemulsification and ECCE,		
including mastery of the following skills:		
a. Wound construction		
b. Anterior capsulotomy/capsulorhexis		
c. Viscoelastics		
d. Intracapsular, extracapsular, and		
phacoemulsification techniques (e.g.,		
sculpting, divide and conquer, stop and chop,		
phaco chop)		
e. Instrumentation and techniques of irrigation		
and aspiration		
f. IOL implantation (e.g., anterior and		
posterior, foldable and non-foldable)		
g. IOL repositioning, removal, or exchange		
3. Perform intraoperative and postoperative		
management of any event that may occur during or		3
as a result of cataract surgery.		

# **Contact Lenses**

1.	Describe the various options for SCL, RPG CL, and hybrid CL	C2
	fitting in advanced ectatic corneal disorders such as	
	keratoconus and pellucid marginal degeneration, including	
	post-intracorneal ring segment implantation cases.	
2.	Describe the various options for SCL and RPG CL fitting in post	C2
	keratoplasty cases.	
3.	Describe the various options for SCL and RPG CL fitting in	C2
	complex post-refractive surgery, including corneal ectasia.	





1. Describe CL fitting in special clinical situations such as severe	C2
dry eye, glaucoma, diabetes, allergy, pregnancy, strabismus,	
and sports practice, adverse environmental and occupational	
conditions.	
2. Describe indications, fitting techniques, and long-term	C2
management of CL wear for children and adolescents.	
3. Describe CL options and most complex fitting techniques for	C2
medical CL indications such as aphakia, albinism, recurrent	
corneal erosions, neurotropic keratitis, corneal scarring,	
aniridia, and prosthetic cosmesis.	
4. Identify indications for scleral CL fitting.	C2
5. Explain reverse geometry RGP CL for post-graft or post-	С3
refractive surgery cases	
6. List the indications for therapeutic CL.	
7. Describe material selection, physiological implications,	C2
mechanisms of action, and adjuvant topical treatment	
associated with therapeutic CL.	
8. Describe the various possibilities of fitting with soft and hard	C2
therapeutic CL.	
9. Explain the importance of appreciating visual acuity, fit, and	C3
comfort in therapeutic CL.	
10. Describe the differences among CL material choices especially	C2
suited for more complex cases and its clinical correlation.	
11. Explain the influence of both systemic and topical medication	C3
on CL fitting and tolerance.	
12. Describe the methods of modifying a CL to improve comfort,	
vision, or physiological response.	C2
13. Evaluate CL-induced complications, and describe treatment	
strategies for their management, in particular acanthamoeba	С3
and fungi infections.	
14. Describe indications and methods for fitting front surface toric,	C2
back surface toric, and bitoric RGP CL.	





1.	Perform an advanced CL history and examination.	P2	3
2.	Obtain a full ocular history and conduct necessary		3
	tests to perform a complex CL fitting examination		
	(e.g., post keratoplasty, multiple surgeries, post-		
	refractive surgery, and corneal ectasia, advanced		
	corneal ectatic disorders such as keratoconus and		
	pellucid marginal degeneration, and active corneal		
	and ocular surface disease).		
3.	Perform refraction, retinoscopy, and over-refraction		3
	in complex cases.		
4.	Use advanced CL designs including reverse		2
	geometry.		
5.	Indicate the auxiliary CL instruments in patients with		2
	complex needs (e.g., computerized topography,		
	fluorescein patterns, and diagnostic lenses).		
6.	Interpret topography in complex CL fittings.		3
7.	Analyze aberrometry and endothelial/confocal		3
	biomicroscopy.		
8.	Indicate CL modification and refitting in complex		2
	cases, when needed.		
9.	Select the appropriate CL in complex clinical cases		2
	(e.g., post keratoplasty, multiple surgeries, post-		
	refractive surgery, and corneal ectasia, advanced		
	ectatic corneal disorders such as keratoconus,		
	pellucid marginal degeneration, and active corneal		
	and ocular surface disease).		
	Perform therapeutic CL fitting and follow up.		3
11.	Manage CL-induced complications, both infectious		
	and noninfectious (e.g., sterile infiltrates, corneal		3
	neovascularization, corneal permanent staining, and		
	giant papillary conjunctivitis).		
	op an educational skill set to effectively educate		
rotatin	g students and residents about CL topics.		

# **Cornea and External Diseases**





1.	Describe the most complex anatomy, embryology, physiology, histopathology, microbiology, immunology, genetics, epidemiology, and pharmacology of the cornea, conjunctiva, sclera, eyelids, lacrimal apparatus, and ocular adnexa.	C2
2.	Discuss the most complex corneal optics and refraction (e.g., post keratoplasty) and their methods of treatment (e.g., contact lenses, refractive surgery).	C2
3.	Describe the most complex and less common congenital abnormalities of the cornea, sclera, and globe (e.g., cornea plana, keratoglobus).	C2
4.	Recognize the less common corneal dystrophies and degenerations (e.g., Meesman dystrophy, Reis-Buckler dystrophy, François syndrome, Schnyder crystalline dystrophy, congenital hereditary stromal dystrophy, congenital hereditary endothelial dystrophy, posterior polymorphous dystrophy) in addition to the more common dystrophies (e.g., anterior membrane dystrophy, granular, lattice, and macular).	C3
5.	Recognize common and uncommon corneal and conjunctival neoplasms and degenerations (e.g., spheroidal degeneration, carcinoma in situ).	C2
6.	<ul> <li>Describe less common and rare ocular infections, and describe the differential diagnosis of the most complicated corneal and conjunctival infections (e.g., amoebas, leishmaniasis, and nematodes).</li> <li>a. In non-endemic areas, describe the basic features of onchocerciasis.</li> <li>b. In endemic areas, define the etiology, vector (e.g., black fly), and incidence, diagnostic features (e.g., microfilariae, keratitis, iritis), diagnosis (e.g., skin snip test), course and prognosis, treatment (e.g., ivermectin, nodulectomy), and</li> </ul>	C2
_	prevention (e.g., vector control, environmental and behavioral changes) of onchocerciasis.	
7.	Describe the most complex principles of ocular pharmacology of anti-infective, anti-inflammatory, and immune modulating agents (e.g., combination therapies of antiviral and anti-inflammatory agents).	C2
8.	Describe the most complex differential diagnosis of red eye (e.g., pemphigoid, pemphigus, Stevens-Johnson syndrome).	C2
9.	Describe the differential diagnosis and the external	~~ ~
	manifestations of the most complex or uncommon anterior	
	segment inflammations (e.g., syphilitic keratouveitis).	C2





10. Describe the indications for ocular surface transplantation,	
including conjunctival autograft/flap, amniotic membrane	C2
transplantation, and limbal stem cell transplantation.	
11. Describe the surgical indications (e.g., Fuchs dystrophy,	
aphakic/pseudophakic bullous keratopathy, and keratoconus),	C2
surgical techniques, and recognition and management of	
postoperative complications (especially immunologically-	
mediated rejection) of corneal transplantation (e.g., penetrating,	
lamellar).	

1.	Interpret the most advanced corneal techniques (e.g., endothelial microscopy, computerized	P2	3
	corneal topography and tomography, anterior		
	segment ocular coherence tomography).		
2.	Perform a thin conjunctival flap (e.g., Gunderson		2
	flap).		
3.	Perform specialized and complicated fitting of		3
	contact lenses (e.g., post keratoplasty, advanced		
	keratoconus).		
4.	Perform more complex corneal surgery (e.g.,		3
	penetrating or lamellar keratoplasty,		
	keratorefractive procedures, and		
	phototherapeutic keratectomy), and understand		
	the postoperative management including		
	postkeratoplasty astigmatism management and		
_	graft rejection.		
5.	Perform other complex conjunctival surgery		3
c	(e.g., autograft, stem cell transplant).		
6.	Manage more complex neoplasms of the		3
	conjunctiva (e.g., carcinoma, melanoma).		

# Glaucoma

1.	Describe	the	etiology,	pathophysiology,	and	clinical	C2
	characteris	tics of	f the most	complex glaucoma	as (e.g	., angle	





	recession, multi mechanism glaucoma, traumatic glaucoma,	
	neovascular, uveitic glaucoma, iridocorneal endothelial	
	syndrome).	
2.	Identify the key examination techniques and management of	C2
	complex medical and surgical problems in glaucoma (e.g.,	
	complicated or postoperative primary and secondary open-angle	
	and closed-angle glaucoma, uncommon visual field defects).	
3.	Describe visual field damage, progression, and rate of	C2
	progression, caveats, and their use in glaucoma management.	
4.	Describe medical management of the most advanced and	C2
	complex glaucoma (e.g., advanced primary open-angle glaucoma	
	previously treated with medicine, laser, or surgery; secondary	
	glaucomas).	
5.	Describe the clinical features of ocular hypotony, recognize and	C2
	know how to treat common and uncommon etiologies (e.g.,	
_	choroidal detachment, leaking trabeculectomy bleb).	
6.	Describe the results, apply the conclusions, and critically analyze	C2
	the major clinical trials in glaucoma (e.g., GLT, OHTS, CIGTS, FFSS,	
	NTGS, AGIS, EGPS, EMGT), as well as describe and use other	
-	publications in the management of glaucoma patients.	
7.	Describe the principles, indications, and complications of laser	
	treatment of more advanced or complex glaucoma (e.g., repeat	c2
0	procedures).	C2
٥.	Describe the more advanced surgical treatment of glaucoma:	
	(e.g., trabeculectomy, combined cataract and trabeculectomy, glaucoma drainage devices, and cyclodestructive procedures),	C2
	including indications, techniques, and complications.	C2
q	Describe use of antimetabolites and anti angiogenic agents and	
5.	potential complications from their use.	
10	Recognize glaucoma surgical complications, their etiologies, and	C3
10.	options for treatment.	
11	Describe new non-penetrating glaucoma surgery techniques:	C2
	principles, techniques, advantages, limitations, and	-
	complications.	C2
12.	Describe new microsurgical devices (e.g., EX-PRESS, iStent, gold	
	shunt, Trabectome) used in glaucoma surgery.	



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1.	Perform YAG or argon laser procedures in glaucoma	P2	2
	patients (e.g., monocular patient, repeat laser,		
	vitreolysis, and suture lysis).		
2.	Perform laser peripheral iridotomy for more		2
	advanced glaucoma (e.g., monocular patient, acute		
	angle closure, hazy cornea).		
3.	Perform laser treatments (e.g., argon laser		2
	trabeculoplasty, iridoplasty) for more advanced		
	glaucoma cases (e.g., repeat treatments,		
	monocular patient).		
4.	Perform cyclophotocoagulation for more advanced		2
	cases (e.g., prior surgery, monocular patient).		
5.	Perform routine and repeat trabeculectomy with or		2
	without antimetabolites.		
6.	Manage medically and/or surgically a flat anterior		3
	chamber as appropriate.		
7.	Perform small incision phaco/intraocular lens		
	surgery combined with trabeculectomy, at the		3
	same or different sites.		

# Neuro-Ophthalmology

1.	Describe the typical and atypical features, evaluation, and management of papilledema and raised intracranial pressure due to a variety of causes (e.g., sinus thrombosis, idiopathic, meningitis).	C2
2.	Describe the typical features, evaluation, and management of urgent neuro-ophthalmic pathologies (e.g., giant cell arteritis, cavernous sinus thrombosis, orbital apex syndrome, pituitary apoplexy).	C2
3.	Describe typical features of the most advanced and least common optic neuropathies (e.g., chronic recurrent inflammatory optic neuritis, posterior ischemic optic neuropathy, neuromyelitis optica, autoimmune optic neuropathy, toxic/nutritional).	C2





4.	Describe typical and atypical features, evaluation, and	C2
	management of the most complex and least common ocular	
	motor neuropathies and their mimics (e.g., patterns of aberrant	
	regeneration).	
5.	Describe typical and atypical features, evaluation, and	C2
	management of the most complex and least common forms of	
	nystagmus (e.g., spasmus nutans, see-saw nystagmus, periodic	
	alternating nystagmus).	
6.	Describe typical and atypical features, evaluation, and	C2
	management of the most advanced and least common pupillary	
	abnormalities (e.g., pupil findings in coma, transient pupillary	
	phenomenon).	
7.	Describe features, evaluation, and management of the most	
	complex and least common visual field defects and recognize	
	pattern mimics (e.g., combination of disc-related scotoma plus	C2
	hemianopia, binasal hemianopia, sectoranopia, bilateral inferior	
	altitudinal loss due to superior occipital lobe lesions and not	
	bilateral anterior ischemic optic neuropathy).	
8.	Describe syndromes of cortical visual dysfunction.	
9.	Detect early neuro-ophthalmic signs and symptoms of drug	
	toxicity for commonly used medications.	
10	. Describe the neuro-ophthalmic complications related to	
	pregnancy.	

<ol> <li>Interpret the complete cranial nerve evaluation in the context of neuroophthalmic localization and diseases.</li> </ol>	P2	3
2. Interpret neuro-radiologic images in neuro- ophthalmology (e.g., interpretation of orbital imaging for orbital pseudotumor and tumors, thyroid eye disease, intracranial imaging modalities and strategies for tumors, aneurysms, infection, inflammation, ischemia), and appropriately discuss, in advance of testing, the localizing clinicoradiological features with the neuroradiologist in order to obtain the best study and interpretation of the results.		3
		3



3.	Identify patients with "functional" visual loss (i.e.,	
	nonorganic visual loss) and provide appropriate	
	counseling and follow-up.	3
4.	Quantify RAPD with neutral density filter and detect	
	small RAPD in patients with only one working pupil.	1c1
5.	Perform optic nerve sheath decompression, if	
	trained, for papilledema.	3
6.	Perform neuro-ophthalmic evaluations for people	
	with special needs (e.g., comatose patients,	
	children, children with developmental and visual	
	maturation evaluations).	

# **Ophthalmic Pathology**

## A. Cognitive Skills

1.	Describe more advanced ocular anatomy	C2
2.	Describe the major histologic findings of common diseases of the	C2
	eye (e.g., keratitis, exfoliation syndrome, corneal and retinal	
	dystrophies and degenerations, frequent neoplasms) relevant to	
	specific clinical rotation(s) (e.g., oculoplastics, cornea, glaucoma,	
	retina, ophthalmic oncology).	
3.	Describe the pathophysiology and histology of potentially vision	C2
	or life-threatening diseases (e.g., temporal arteritis,	
	endophthalmitis, retinoblastoma, ocular melanoma, extraocular	
	or orbital spread of an intraocular or periorbital tumor, metastasis	
	to the eye and orbit) relevant to specific clinical rotation(s) (e.g.,	
	oculoplastics, cornea, glaucoma, retina, and ophthalmic	
	oncology).	
4.	Interpret reports of more advanced techniques in ophthalmic	С3
	histopathology (e.g., cytology, special stains, transmission	
	electron microscopy, immunohistochemistry, tumor free	
	margins) relevant to specific clinical rotation(s) (e.g.,	
	oculoplastics, cornea, glaucoma, retina, ophthalmic oncology),	
	including how the clinician communicates the need for these	
	studies.	





1.	Process appropriately more advanced specimens for submitting to an ophthalmic pathology laboratory, including writing of the accompanying letter to the ophthalmic pathologist (e.g., impression cytology, fine needle aspiration biopsy, vitreous biopsy, evisceration, exenteration specimen).	P2	3
2.	Perform a biopsy for frozen section study in ocular pathology.		3
3.	Participate under supervision through a site visit in a macroscopic and microscopic examination of ophthalmic specimens from active cases, working from low to high power.		3

# Oculoplastic Surgery and Orbit

# A. Cognitive skills

# Eyelid

1.	Descri	be the most advanced eyelid anatomy and physiology.	C2
2.	Descri	be the etiology, evaluation, and medical and surgical	C2
	treatm	nent of the following eyelid diseases:	
	a.	Complex ectropion (e.g., congenital, paralytic,	
		involutional, cicatricial, mechanical, allergic)	
	b.	Complex entropion (e.g., involutional, spastic,	
		cicatricial, congenital)	
	с.	Complex myogenic ptosis (e.g., myasthenia gravis,	
		chronic progressive external ophthalmoplegia [CPEO],	
		oculopharyngeal muscular dystrophy [OPMD],	
		myotonic dystrophy)	
	d.	Upper eyelid retraction	
	e.	Lower eyelid retraction	
	f.	Benign, pre-malignant, or malignant eyelid tumors	
		(e.g., papilloma, seborrheic keratosis, epidermal	
		inclusion cyst, molluscum contagiosum, verruca	
		vulgaris, keratoacanthoma, actinic keratosis, basal cell	
		carcinoma, squamous cell carcinoma, sebaceous cell	
		carcinoma, melanoma)	
	g.	Single or recurrent inflammatory lesions (e.g.,	
		recurrent chalazion or its mimics)	



h.	Facial nerve pals	y with	exposure kera	atopathy	(e.g.,
	tarsorrhaphy,	gold	weight,	lower	lid
	tightening/elevati	on)			

## Lacrimal

1. Describe the most advanced lacrimal anatomy and	C2
physiology.	
2. Describe the etiology, evaluation, and medical and surgical	C2
treatment of the following lacrimal diseases:	
a. Punctal stenosis	
b. Canalicular stenosis	
c. Common canalicular stenosis	

## Orbital

1. Describe the most advanced orbital anatomy and physiology.	C2
2. Describe the etiology, evaluation, and medical and surgical	C2
treatment of the following orbital diseases:	
a. Orbital trauma	
i. All orbital fractures	
ii. Retrobulbar hemorrhage	
iii. Orbital foreign bodies	
b. Orbital neoplasms	
i. All benign	
ii. All malignant	
c. Orbital inflammation	
i. Infectious	
1. Bacterial	
2. Fungal	
3. Mycoplasma	
ii. Noninfectious	
1. Thyroid eye disease	
2. Sarcoidosis	
3. Wegener granulomatosis	
4. Nonspecific orbital inflammation	
5. Describe epidemiology, clinical	
features, evaluation, and management	
of fetal alcohol syndrome.	



### Eyelid

1. Perform more complicated and advanced "in	P2	3
office" examination techniques for less common		
but important eyelid abnormalities.		
2. Perform more complicated lid procedures,		3
including:		
a. Frontalis sling		
b. Lateral tarsal strip		
c. Eyelid reconstruction		

### Lacrimal

1.	Perform more complicated and advanced "in	3	
	office" examination techniques for less common		
	but important lacrimal abnormalities.		
2.	Perform more advanced lacrimal assessment (e.g.,		
	intraoperative and postoperative testing, more complex trauma to lacrimal system).	3	
3.	Manage lacrimal system abnormalities, including		
	surgeries (e.g., lacrimal probing, dacryocystectomy, dacryocystorhinostomy).	3	

## Orbital

1.	Perform more complicated and advanced "in	P2	
	office" examination techniques for less common		
	but important orbital abnormalities (e.g., forced		
	duction testing).		
2.	Describe typical and atypical features and describe		
	the differential diagnosis, clinical features, and		C2
	treatment of more complicated orbital diseases,		
	including:		
	a. Complex orbital infections (e.g., orbital		
	cellulitis, mucormycosis, aspergillosis		
	b. Congenital tumors (e.g., dermoid)		



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			1
	c. Fibro-osseous disorders and tumors (e.g	,	
	fibrous dysplasia, osteoma	,	
	chondrosarcoma, osteosarcoma, Page	:	
	disease)		
	-	,	
	hemangioma, cavernous hemangioma		
	hemangiopericytoma, lymphangioma	,	
	Kaposi sarcoma)		
	e. Xanthomatous tumors (e.g., xanthelasma	,	
	juvenile xanthogranuloma)		
	f. Lacrimal gland tumors (e.g., benign mixed		
	tumor, adenoid cystic carcinoma		
	, , ,	,	
	malignant mixed tumor, lymphoma)		
	g. Neural tumors (e.g., optic nerv		
	glioma/meningioma, neurofibromatosis	,	
	neuroblastoma, schwannoma)		
	h. Sarcomas (e.g., rhabdomyosarcoma	,	
	leiomyosarcoma, liposarcoma	,	
	osteosarcoma)		
	i. Lymphoid lesions (e.g., lymphoid		
	hyperplasia, lymphoma, leukemia)		
	j. Metastatic lesions (e.g., from breast	,	
	prostate, lung, colon)		
	k. Thyroid eye disease		
	I. Nonspecific orbital inflammation		
	m. Trauma (e.g., fractures, foreign body	,	
	retrobulbar hemorrhage, traumatic opti		
	neuropathy)		
3.	Describe indications and complications of basi		
	orbital skills and procedures, including:		
	a. Anterior orbitotomy for tumo		
	,		C2
	biopsy/excision		C2
~	b. Orbital floor fracture repair		
4.	Describe indications and complications of differen		
	orbital approaches and incisions (e.g., Kronleir	,	
	Caldwell-Luc, transconjunctival, transnasal).		
5.	Describe indications for orbital ultrasound	,	
	computerized axial tomography (CT or CAT) scar	,	C2
	and magnetic resonance imaging (MRI) scan (e.g	,	
	orbital trauma, orbital lesions, and tumors).		



# Pediatric Ophthalmology and Strabismus

1. Describe more advanced anatomy (including pulleys) and physiology of strabismus (e.g., torsion, tertiary actions,	C2
consecutive deviations).	
2. Describe more advanced sensory adaptations (e.g., anomalous	C2
head position).	
3. Describe and recognize the different forms of childhood	C2
nystagmus (e.g., infantile nystagmus syndrome [INS], fixation	
maldevelopment nystagmus syndrome [FMNS], spasmus nutans syndrome [SNS]), and appropriate work up for different	
time of onset and age groups.	
4. Describe and recognize ROP (e.g., stages, treatment	C2
indications).	
5. List treatment options and indications of low-birth-weight	C2
children, and describe long-term ocular and systemic problems.	
6. Describe and recognize less common hereditary or	
malformative ocular anomalies and syndromes (e.g., Mobius	
syndrome, Goldenhar syndrome, Peter anomaly, including pedigree chart analysis).	
7. Describe etiology, evaluation, and management of congenital	C2
infections (e.g., TORCHES sequence: TOxoplasmosis, Rubella,	
Cytomegalovirus, HErpes simplex, Syphilis).	
8. Describe and recognize the common causes of pediatric uveitis	C2
with natural history, indicated work up, and treatment.	
9. Describe congenital optic nerve anomalies in children (e.g.,	C2
optic nerve coloboma, morning glory syndrome, optic nerve	
hypoplasia), and indicate necessary work up and associated diseases.	
10. Describe American Association for Pediatric Ophthalmology and	C2
Strabismus (AAPOS) etiology position statements on learning	
difficulties and dyslexia, and know how to locate educational	
support resources for parents.	
11. Identify referral centers for children with retinoblastoma, the	C2
work up for leukocoria, the evaluation of family members, and	
the principals of genetic counseling. 12. Describe typical features of childhood tumors (e.g.,	C2
12. Describe typical features of childhood tumors (e.g., hemangiomas, rhabdomyosarcoma) and their management.	





13. Describe identifiable congenital ocular anomalies (e.g.,	
microphthalmia, persistent fetal vasculature), and describe	C2
appropriate work up for etiology, criteria for intervention, and	
genetic counseling for parents.	
14. Describe indications for botulinum toxin use in strabismus.	

1.	Perform a more advanced extraocular muscle examination based on knowledge of the anatomy and physiology of ocular motility.	P2	3
2.	Assess more advanced ocular motility problems (e.g., bilateral or multiple cranial neuropathy, myasthenia gravis, thyroid eye disease).		
3.	Apply Hering law and Sherrington law in more advanced cases (e.g., pseudoparesis of the contralateral antagonist, enhancement of ptosis in myasthenia gravis).		3
4.	Perform more advanced measurements of strabismus (e.g., use of synoptophore or amblyoscope, when available).		3
5.	Perform assessment of vision in more difficult strabismus patients (e.g., uncooperative child, mentally impaired, nonverbal, or preverbal).		3
6.	<ul> <li>Perform the following surgical techniques:</li> <li>a. Muscle weakening (e.g., tenotomy) and strengthening (e.g., tuck) procedures of rectus muscles</li> <li>b. Inferior oblique weakening procedures</li> </ul>		3
7.	c. Use of adjustable sutures Manage the complications of strabismus surgery (e.g., slipped muscle, anterior segment ischemia, overcorrection, and under correction).		3

# **Vitreoretinal Diseases**



1.	Describe and apply retinal electrophysiology.	C2
2.	Evaluate, treat, or refer the most complex forms of retinal	C3
	vascular diseases:	
	a. Combined arterial and venous obstructions	
	b. Advanced diabetic retinopathy	
	c. Advanced hypertensive retinopathy	
	d. Peripheral retinal vascular occlusive disease	
3.	Describe the findings of major studies in age-related macular	C2
	degeneration:	
	a. Treatment of Age-Related Macular Degeneration with	
	Photodynamic Therapy Study (TAP)	
	b. Verteporfin in Photodynamic Therapy Study (VIP)	
	c. Minimally Classic/Occult Trial of the Anti-Vascular	
	Endothelial Growth Factor (VEGF) Antibody Ranibizumab	
	in the Treatment of Neovascular AMD (MARINA)	
	d. Anti-VEGF Antibody for the Treatment of Predominantly	
	Classic Choroidal Neovascularisation in AMD (ANCHOR)	
	e. The Comparisons of Age-Related Macular Degeneration	
	Treatments Trials (CATT)	
4.	Evaluate and diagnose complex cases of retinal detachment (e.g.,	C3
	acute retinal necrosis, proliferative vitreoretinopathy).	
5.	Diagnose and classify retinopathy of prematurity.	C2
6.	Diagnose and manage (or refer) complex trauma cases (e.g.,	C2
	chorioretinitis sclopetaria, intraocular foreign body, shaken baby	
	syndrome).	
7.	Diagnose hereditary vitreoretinal degenerations (e.g., Stickler	C2
	syndrome, Wagner syndrome, Goldmann-Favre degeneration).	
8.	Describe the treatment algorithm for each specific retinal	C2
	condition, with special emphasis on pros and cons.	

<ol> <li>Perform indirect ophthalmoscopy with scleral indentation in complex retinal cases (e.g., multiple holes, documented with detailed retinal drawing).</li> </ol>	Ρ2	3
2. Perform ophthalmoscopic examination with panfunduscopic or other lenses in complex retinal		3



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	conditions (e.g., giant retinal tears, proliferative		
	vitreoretinopathy).		3
3.	Interpret and apply in clinical practice the results		
	of fluorescein and ICG angiography and OCT in		
	complex retinal or choroidal pathology.		3
4.	Perform posterior segment photocoagulation in		
	more complicated retinal cases:		3
5.	Diabetic focal/grid macular treatment (e.g.,		
	monocular patient, repeat treatment)		3
6.	Repeat peripheral scatter photocoagulation		
	(panretinal)		3
7.	Laser retinopexy (demarcation) of large or		
	multiple breaks; cryotherapy		
8.	Interpret and apply in clinical practice		3
	electrophysiology (e.g., ERG, EOG, VEP, dark		
	adaptation) in more complicated retinal		
	pathology.		3
9.	Interpret and apply in clinical practice ocular		
	imaging techniques (e.g., B-scan echography) in		
	more complex cases (e.g., choroidal osteoma).		3
10.	Perform detailed fundus drawings of the retina		
	with vitreoretinal relationships in the most		
	complex retinal cases (e.g., recurrent retinal		
	detachment, retinoschisis with and without		
	retinal detachment).		
11.	Perform laser therapy or cryotherapy of retinal		
	holes and other more complex retinal		2
	pathologies.		
12.	Participate during scleral buckling and pars plana		2
	vitrectomy surgeries.		
		L	0

# **Uveitis and Ocular Inflammation**

1. Describe the more complex complications of common uveitis	C2
syndromes in addition to that mentioned in Year 2 (e.g., retinal	
vascular occlusion, retinal neovascularization and vitreous	





	hemorrhage, inflammatory choroidal neovascularization,	
	hypotony).	C2
2.	Describe indications and contraindications for corticosteroid	
	treatment of uveitis (e.g., topical, local, systemic), including	
	risks and benefits of therapy.	C2
3.	Describe the management of common uveitic syndromes.	C2
4.	Describe the techniques of anterior chamber and vitreous tap	
	and of intravitreal injection of antibiotics in cases of bacterial	
	endophthalmitis.	C2
5.	Describe more advanced examination principles for patients	
	with more subtle signs of uveitis, such as:	
	a. Anterior segment (e.g., conjunctival ulcer, iris	
	transillumination defects, granuloma)	
	b. Posterior segment (e.g., pars plana signs of	
	inflammation [snowbanks and snowballs], retinal	
	detachment [exudative, tractional, rhegmatogenous],	
	retinal vasculitis [periphlebitis or arteritis, occlusive or	
	nonocclusive], optic nerve [optic disc granuloma, optic	
	neuritis, disc neovascularization], macula [macular	
	edema, choroidal neovascularization])	C2
6.	Describe in greater detail the angiographic features of retinitis,	
	choroiditis, and vasculitis.	C2
7.	Describe indications and contraindications for commonly used	
	immunotherapy for uveitis in addition to corticosteroid therapy	
	(e.g., azathioprine, cyclosporine A), including risks and benefits	
	of therapy.	C2
8.	Describe the clinical features and differential diagnoses for less	
	common forms of uveitis (e.g., Whipple disease, Crohn disease).	

1. Perform a more advanced examination of the	P2	3
anterior and posterior segment, for example:		
a. Anterior segment (e.g., conjunctival ulcer,		
iris transillumination defects, granuloma)		
b. Posterior segment (e.g., pars plana signs of		
inflammation [snowbanks and snowballs],		
retinal detachment [exudative, tractional,		
rhegmatogenous], retinal vasculitis		
[periphlebitis or arteritis, occlusive or		



	· ·
nonocclusive], optic nerve [optic disc	
granuloma, optic neuritis, disc	
neovascularization], macula [macular	
edema, choroidal neovascularization])	
2. Differentiate active from inactive disease and	3
arterial from venous side disease.	
3. Recognize serious infective causes from	
noninfective causes of uveitis.	3
4. Recognize and evaluate the typical demographic	
features, clinical features, and differential diagnosis	3
of uveitis common in the region via the process of	
history taking, clinical examination, and the use of	
investigative tools (such as FA, ICG, B-scan, OCT).	
5. Recognize and evaluate the typical demographic	3
features, clinical features, and endophthalmitis)	
Masquerade syndromes, such as vitreoretinal	
lymphoma differential diagnosis of uveitis in:	
a. Immunosuppressed individuals (e.g.,	
cytomegalovirus retinitis, endogenous	
6. Evaluate the common complications of common	3
uveitic syndromes (e.g., glaucoma, cataract, band	
keratopathy, macular edema).	
7. Administer periocular corticosteroid injections in	3
addition to topical corticosteroids in the treatment	
of uveitis.	
8. Perform an anterior chamber and vitreous tap for	
diagnostic purposes and to give intravitreal	3
injection of antibiotics in cases of bacterial	
endophthalmitis.	
9. Perform cataract removal.	3
10. Perform filtration surgery with antimetabolites.	3
11. Provide patient with relevant information about	3
possible side effects of medications and proper	
monitoring of medications.	
	3

# Ocular Oncology





1.	Describe the applied surgical anatomy, histology, and physiology	C2
	of the eye and ocular adnexa with relevance to ocular oncology.	
2.	Enlist the most common conjunctival and intraocular tumors.	C2
3.		C2
	a. Nonneoplastic tumors (e.g., hamartomas)	
	b. Neoplastic tumors	
	c. Benign (e.g., nevus, hemangioma)	
	d. Malignant (e.g., melanoma, carcinoma, metastasis)	
	e. Traumatic lesions (e.g., implantation cysts, hemorrhages)	
	<ul> <li>f. Degenerative lesions (e.g., disciform, sclero-choroidal calcification)</li> </ul>	
	g. Idiopathic disease (e.g., juvenile xanthogranuloma,	
	vasoproliferative tumor)	
	h. Paraneoplastic disease (e.g., Bilateral diffuse uveal	
	melanocytic proliferation)	
	i. latrogenic disease (e.g., radiation-induced disease)	
4.	Describe relevant pathological techniques (e.g., fixation,	C2
	histology, immunohistochemistry).	
5.	Describe relevant genetic abnormalities and techniques:	C2
	a. Germinal and somatic mutations relevant to oncology	
	(e.g., retinoblastoma)	
	b. Important genetic techniques (e.g., fluorescence in situ	
	hybridization)	
6.	Describe the relevance of staging tumors (e.g., TNM [Tumor,	C2
	lymph Nodes, Metastasis] Classification of Malignant Tumors).	
7.	Describe the etiology of ocular tumors, such as:	C2
	a. Environmental factors (e.g., conjunctival squamous cell	
	carcinoma)	
	b. Genetic factors (e.g., retinoblastoma)	
	c. Syndromes (e.g., von Hippel-Lindau disease)	
	d. Malformations (e.g., choroidal osteoma)	
8.	Describe the pathogenesis of ocular tumors (i.e., how tumors	C2
	cause harm):	
	a. Ocular effects (e.g., neovascular glaucoma)	
	b. Systemic effects (e.g., metastatic disease)	
9.	Describe the epidemiology of the more common ocular tumors	C2
	(e.g., melanoma).	
10.	Describe the principles of examination techniques:	C2
	a. Inspection	
	b. Transillumination	
	c. Color photography	



d. Optical coherence tomography	
e. Autofluorescence	
f. Angiography (indocyanine green and fluorescein)	
g. Ultrasonography	
h. Magnetic resonance imaging	
i. Computerized tomography	
j. Positron emission tomography	
k. Biopsy	
i. Incisional	
ii. Aspiration	
iii. Excisional	
iv. Impression cytology	
v. Systemic investigation according to ocular tumor	
diagnosis	
vi. History	
vii. Clinical examination	
viii. Hematology and biochemistry	
ix. Radiography	
x. Ultrasonography	
xi. Computerized tomography	
xii. Magnetic resonance imaging	
xiii. Genetic testing	
11. Describe the clinical features of each tumor type:	C2
a. Inspection/color photography	
<ul> <li>Investigational (i.e., angiography, echography)</li> </ul>	
c. List the differential diagnosis of each tumor, and describe	
the investigational approach for each condition.	
12. Describe how the following therapeutic modalities and their	C2
effects are relevant to ocular tumors:	
a. Radiotherapy (e.g., brachytherapy, external beam	
radiotherapy, proton beam)	
b. Chemotherapy (e.g., topical, intraocular, systemic)	
c. Phototherapy (e.g., photocoagulation, photodynamic	
therapy)	
d. Cryotherapy (e.g., liquid nitrogen, carbon dioxide)	
e. Surgical resection (e.g., local resection, enucleation)	C2
13. Describe how statistics can be applied to ocular oncology (e.g., survival analysis).	
14. Describe the methods, risks, and benefits of tumor biopsy and	C2
how these can be avoided (e.g., biopsy of retinoblastoma,	
incisional biopsy of conjunctival tumor).	
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1.	Perform or request appropriate examinations	P2	3
	and investigations according to differential		
	diagnosis.		
2.	Perform or refer for treatment for conjunctival		3
	or intraocular tumors, demonstrating awareness		
	of the indications, contraindications, and		
	complications of each treatment and having skill		
	to administer short-term and long-term		
	postoperative care:		
	a. Radiotherapy (e.g., brachytherapy,		
	external beam radiotherapy)		
	b. Phototherapy (e.g., photodynamic		
	therapy, transpupillary thermotherapy)		
	c. Surgical excision (e.g., local resection,		
	enucleation, exenteration)		
	d. Ocular pharmacological therapy by		
	various routes (i.e., topical, intravitreal,		
	ophthalmic artery infusion, sub tenon,		
	systemic)		
	i. Chemotherapy and biological		
	therapy		
	ii. Antiangiogenic agents		
_	iii. Steroids		
3.	Interpret results of relevant laboratory tests and		
	communicate results to patients, relatives, and		3
	health care workers; and adjust patient		
	management accordingly.		
4.	Communicate prognosis with patients, relatives,		
	and health care workers; and adjust patient		2
	management accordingly in collaboration, if		3
-	necessary, with a subspecialist.		
5.	Use information technology and other aids to		
C	cope with lack of expert knowledge.		4
0.	Assist patients with selecting the most appropriate management in collaboration, if		4
	necessary, with a subspecialist in ocular		-
	oncology.		



7. Provide or organize appropriate psychological	
support, demonstrating empathy and an	3
adequate awareness of the principles of this	
aspect of care (e.g., giving bad news).	
8. Collaborate with subspecialists and other health	
care professionals to provide patient focused	
care.	4
Develop protocols and infrastructure for practice-based	
learning and improvement (e.g., access to information,	
outcomes data).	

## Low Vision Rehabilitation

### A. Cognitive Skills

1.	Describe significant comorbidities that impact low vision	C2
	rehabilitation.	
2.	Describe the role of visual processing and perception deficits	C2
	(e.g., cerebral visual	
	Impairment, acquired brain injury, stroke).	
3.	Describe indications for the most complex low vision aids.	C2
4.	Apply more complex principles of optics of low vision devices.	C3
5.	Describe vision related quality of life measurements.	
6.	Describe social or public consequences and implications of low	C2
	vision.	
7.	Describe the role of the electrophysiological examinations as	C2
	diagnostic and prognostic tools for low vision patients.	
8.	Describe the implications of low vision in the education of	C2
	children.	

1. Evaluate visual acuity and visual field for	P2	3
determination of disability for legal and		
insurance purposes.		
2. Prescribe the most complex rehabilitative		3
therapies and optical devices to help the		
patient meet their goals.		



3.	Apply and prescribe visual field enhancing	3
	techniques, including scanning training for	
	hemianopic field loss.	
4.	Perform short cognitive assessment of elderly	
	patients with visual impairments for drivers'	3
	license approval.	

# Ethics and Professionalism in Ophthalmology

1.	Recognize and use advanced medical ethics in the ophthalmic		C2
	practice:		
	a. Applicable informed consent documents (e.g., clinical		
	research, off-label use disclosures)		
	b. Management (offering and rendering) of second		
	opinions		
	c. Individual and institutional responsibilities regarding		
	impaired physicians		
	d. Responsibility for postoperative care, including		
	appropriate transfer of care to other physicians		
	e. Management of conflicts of interest (clinical and		
	nonclinical)		
	i. Disclosures		
	ii. Gifts to physicians		
	f. Appropriate advertising (and applicable laws)		
	i. Appropriate conduct as a medical-expert witness		
	in litigation		
2.	Identify applicable insurance coverage responsibilities in a		C2
	practice situation.		
3.	Utilize more advanced aspects of health care reimbursement in		
	a clinical practice (e.g., denials of claims, hospital contracting,		
_	electronic billing).		
4.	Work within integrated eye care delivery systems (both within	P2	
_	eye care specialties and within general medicine and surgery).		
5.	Participate in all of the foregoing aspects of practice		
	management to the best ability with in a medical education		
~	setting.		
6.	Utilize all of the foregoing ethical principles and knowledge in		
	direct patient care.	P2	





7. Describe the responsibility of ophthalmologists to share their	C2
knowledge of clinical arts and sciences for the benefit of	
patients, the profession, and society.	

# **Community Eye Health**

### A. Cognitive Skills

### Principles of prevention of blindness

1.	Outline the different health service models in different (	C2
	countries and regions, and how eye care services might be	
	integrated into these.	
2.	Describe the components of a rapid assessment of avoidable	C2
	blindness (RAAB) survey.	
3.	Outline the government and nongovernment funding that are	C2
	available for eye care.	
4.	Describe the key practices and policies that will ensure the	C2
	principles of prevention of blindness are inclusive relating to	
	gender, disability, and other potential causes of	
	marginalization.	

#### Cataract

1. Outline the components of a system for monitoring the visual	C2
acuity outcomes following cataract surgery.	
2. Outline the components of the cataract surgery costs.	C2

### Trachoma

1. Describe the components of a rapid assessment of trachoma	C2
(RAT) survey.	

### B. Psychomotor Skills

#### Cataract

1.	Set up a system for the monitoring of the visual acuity	P2	
	outcomes following cataract surgery.		



2. Calculate cataract surgery costs with recommendations for	
strategies to decrease unit costs.	

#### **Refractive error**

1.	Evaluate the coverage and impact of school screening, and	С3
	make recommendations for improvement.	
2.	Evaluate the services for the provision of presbyopic	
	correction, and make recommendations for improvement.	

#### Low vision

1. Evaluate the coverage and impact of low-vision services.	1. Evaluate the coverage and impact of low-vision services.		С3
---	---	--	----

#### Childhood blindness

1. Where appropriate, set up a system for the screening and	P1	
treatment of retinopathy of prematurity.		

#### Trachoma

1. Where appropriate, network and advocate with agencies and	P2	
communities to implement the F (facial cleanliness) and E		
(environmental changes) components in the SAFE strategy.		

## Planning of blindness prevention programs

1. Develop a budget for a one-year operational plan for a	P2	
blindness prevention program for a health district with a		
population of one million.		

### 4<sup>th</sup> Year Resident Goals

By the completion of their 4<sup>th</sup> year residency, the residents should be able to:

### **Cataract and Lens**

1. Describe the issues of pediatric cataract surgery, including the	C2
indications for surgery (posterior capsulotomy +/- anterior	





vitrectomy), IOL implantation, unilateral and bilater	ral
congenital cataract, and IOL calculation in young children.	
2. Describe the management of cataract associated with anirid	
3. Describe the treatment options for "dropped IOL" a	nd C2
indications for referral to a vitreoretinal surgeon.	
4. Describe the advantages and strategies for advance	ed <b>C2</b>
phacoemulsification techniques such as torsional	or
transversal ultrasound, small incision and microincisi	on
cataract surgery (MICS), biaxial MICS cataract surgery.	
5. Describe the parameters, power, and fluidics in MICS.	C1
6. List the indications for triple procedures or combined surgeri	ies C1
(e.g., Phaco plus trabeculectomy, keratoplasty, silicone-	oil
removal).	
7. List the Indications for "premium" IOLs (e.g., multifoc	al, <b>C2</b>
accommodating, toric).	
8. Describe the surgical difficulties of hyper mature (Morgagnia	an) <b>C2</b>
cataract.	, -
9. Describe the treatment options for eyes with shallow anteri	ior C2
chamber and cataract including high-degree hyperopic ey	
and piggyback IOL implantation.	
10. Describe the treatment of cataract in patients with	an <b>C2</b>
· · · · · ·	
intraocular tumor (e.g., melanoma, retinoblastoma).	ad <b>C2</b>
11. Describe the methods to determine typical surgically induc	ed C2
astigmatism and surgeon specific a-constant.	
12. Describe the etiology and management of unexpect	
postoperative refractive errors, including hyperopic a	
myopic shifts (e.g., capsular phimosis, capsular block, a	nd
upside down IOL).	
13. Describe the management strategies to reposition	of <b>C2</b>
decentered, tilted, subluxated, and dislocated IOLs.	

1.	Perform surgery on congenital cataract,	P2	3
	including IOL power calculation.		
2.	Perform and teach small incision and MICS,		4
	torsional, or transversal ultrasound.		
3.	Perform and teach triple procedures or		3
	combined surgeries (e.g., phaco and		



trabeculectomy, keratoplasty, silicone-oil removal).	
<ol> <li>Implant "premium" IOLs (e.g., multifocal, accommodating, toric) and counsel patients preoperatively and postoperatively.</li> </ol>	3
<ul> <li>5. Perform surgery on patients with complex lens issues, including:</li> <li>a. Aniridia, iris coloboma, iris dialysis</li> <li>b. Hyper mature (Morgagnian) cataract</li> <li>c. Eyes with shallow anterior chamber</li> </ul>	3
<ul> <li>d. High-degree myopic eyes</li> <li>6. Perform reposition of malpositioned IOLs and late subluxation of IOL/capsule.</li> </ul>	3

### **Cornea and External Diseases**

1.	Recognize acute and chronic blepharitis, including both infectious and noninfectious etiologies, with emphasis on microbial blepharitis, meibomian gland dysfunction, and rosacea.	C2
2.	Recognize acute and chronic conjunctivitis, neonatal conjunctivitis, chlamydial disease, adenoviral conjunctivitis, allergic conjunctivitis, and bacterial conjunctivitis.	C2
3.	Recognize acute and chronic infectious keratitis including bacterial, viral, fungal, and parasitic, with emphasis on herpes simplex, herpes zoster, adenovirus, acanthamoeba, and contact lens-associated problems.	C2
4.	Recognize noninfectious keratitis including marginal keratitis, central ulcerative keratitis, epitheliopathy, endothelialitis, and interstitial keratitis.	C2
5.	Recognize anterior segment anomalies, including various anomalies associated with specific genetic abnormalities, corneal dystrophies, and corneal degenerations.	C2
6.	Recognize autoimmune and immunologic diseases of the anterior segment including allergy, corneal graft rejection, and cicatrizing conjunctivitis.	C2





7.	Recognize and be familiar with oral and topical	C2
	immunosuppression and anti-allergy medications.	
8.	Describe fundamentals of anterior segment anatomy,	C2
	chemistry, physiology, and wound healing including tear	
	formation and function, corneal topography/tomography,	
	endothelial cell function, and maintenance of corneal clarity.	
9.	Discuss principles of anterior segment pharmacology including	C2
	antimicrobial, anti-inflammatory, ocular hypotensive and	
	immunosuppressive agents, with emphasis on bioavailability,	
	mechanism of actions, relative efficacy, safety, and potential	
	complications.	

### Glaucoma

1.	List the main population-based studies in glaucoma prevalence,	C2
	incidence, and risk factors (e.g., Baltimore Eye Survey, Blue	
	Mountains Eye Study, Barbados Eye Study, Rotterdam Eye	
	Study, Thessaloniki Eye Study, Latinos Eye Study, Singapore	
	Malay Eye Study).	
2.	Describe and critically discuss results of the above-mentioned	C2
	studies on glaucoma prevalence, incidence, and risk factors.	
3.	Describe rate of progression and use of special algorithms (e.g.,	C2
	value function iteration, PROGRESSOR, Garway-Heath map).	
4.	Describe and critically discuss literature on structure-function	C2
	correlation.	
5.	Describe use of other tonometers (e.g., ocular response	C2
	analyzer, dynamic contour tonometry, pneumotonometer).	
6.	Describe mechanisms of ganglion cell damage and potential	C2
	pathways for neuroprotection.	
7.	Describe and know specific medical and surgical treatments in	C2
	the most complex and most advanced glaucoma cases (e.g.,	
	refractory glaucoma, monocular patients, and noncompliant	
	patients).	
8.	Describe and know the specific management of complications	C2
	related to the surgical intervention of the most complex and	
	most advanced glaucoma's.	



1.	Perform goniotomy, trabeculotomy, and	P2	3
	manage complications.		
2.	Medical and surgical management of hypotony		3
	from overfiltration, bleb leak, choroidals, and		
	other causes.		
3.	Treat malignant glaucoma and manage		3
	complications.		
4.	Treat failing or leaking blebs at slit lamp and		4
	manage complications.		
5.	Perform advanced techniques for revisions of		3
	glaucoma surgery blebs (e.g., sliding flap, free		
	graft, amniotic membrane) and manage		
	complications.		_
6.	Perform cyclodestructive procedures and		4
-	manage complications.		
7.	Perform trabeculectomy revisions, glaucoma		3
	drainage device surgery, and manage		
0	complications.		2
	Describe and manage cyclodialysis cleft.		3
	Perform releasable suture techniques.		3
	. Perform choroidal drainage. . Perform Phaco trabeculectomy/combined		3
ΤT			3
10	surgery and manage surgical complications.		
ΤZ	. Perform laser trabeculoplasty and manage surgical complications.		
12	. Manage end stage and high-risk glaucoma.		
	. Perform combined implant/phaco/penetrating		
14	keratoplasty/vitrectomy.		
			1

## Neuro-Ophthalmology

1.	Describe the arterial circulation in detail and know the general	C2
	venous drainage along the entire anterior visual pathway	
	(e.g., optic disc, retrobulbar optic nerve, intracranial segment	
	of optic nerve, chiasm, and lateral geniculate body).	
2.	Describe evaluation, give differential diagnosis, and outline a	C2
	management plan of the most advanced and least common	





	optic neuropathies (e.g., chronic recurrent inflammatory optic neuritis, posterior ischemic optic neuropathy, neuromyelitis optica, autoimmune optic neuropathy, rare toxic optic	
2	neuropathies).	
3.	Describe the cortical visual syndromes and know the	<b>C</b> 2
	localization of the causative lesion (e.g., akinetopsia, prosopagnosia, simultagnosia).	C2
4.	Describe typical and atypical features, evaluation, and	
	management of rare eye movement disorders (e.g.,	C2
	differential diagnosis of monocular oscillations, localization of	
	lesion and purported mechanism of oculopalatal myoclonus).	
5.	Describe typical features, pathophysiology, evaluation, and	
	management of rare pupillary syndromes (e.g., tadpole pupil, paradoxical pupillary constriction).	C2
6.	Describe the advantages, disadvantages, indications, and	
	pitfalls in special perimetric methods (e.g., blue-yellow	C2
	perimetry, automated kinetic perimetry, motion perimetry,	
7	and microperimetry). Describe and differentiate among various kinds of unusual	
7.	positive visual phenomena and know their possible causes	C2
	(e.g., palinopsia, persistent photopsia).	
8	Know the differential diagnosis and evaluation for acute or	
0.	progressive homonymous hemianopsia in a patient with a	C2
	normal MRI.	
9.	Describe the various prion diseases and their management.	
10.	Describe the various mitochondrial syndromes that have	C2
	neuro-ophthalmic manifestations, and provide appropriate	C2
	genetic counseling for inherited neuroophthalmic diseases	
	(e.g., Kearns-Sayre and related syndromes, mitochondrial	
	encephalomyopathy, lactic acidosis, stroke-like episodes	
	[MELAS], neuropathy, ataxia, and retinitis pigmentosa	
	[NARP]).	
11.	Describe evaluation, give differential diagnosis, and outline a	
	management plan for patients with headache and facial pain	C1
	presenting as neuro-ophthalmic manifestations.	
12.	Describe the features, evaluation, and differential diagnosis of	
	dizziness and vertigo from neuro-ophthalmic problems.	C1



1.	Recognize pitfalls in interpretations of unusual results of pharmacologic tests used for diagnosis of	P2	3
	pupillary disorders.		
2.	Know techniques that reveal the most subtle		3
	manifestations of eye movement disorder		
	(e.g., slow medial rectus saccade as the only sign of		
	internuclear ophthalmoplegia, fundus photos for		
	excyclotorsion, head shaking test).		
3.	Perform and interpret the complete neurologic		3
	examination.		
4.	Be able to detect symptomatic lesions overlooked		2
	by the neuroradiologist (e.g., small lesion in optic		
	canal, carotid dissection).		
5.	Be able to perform specific maneuvers that		2
	definitively reveal nonorganic visual loss or overlay		
	(e.g., 4-diopter prism test, rocking mirror).		
6.	Perform and interpret spectral-domain OCT (e.g.,		2
	outer retinal disorders, detection of drusen).		
7.	Interpret indocyanine green angiography and		3
	autofluorescence imaging.		

## **Ophthalmic Pathology**

1.	Describe advanced ocular anatomy, and identify histology of the	C2
	minor structures of the eye and their uncommon variants (e.g.	
	congenital grouped pigmentation).	
2.	Describe the more complex pathophysiology of the disease	C2
	processes of the eye, and identify major histologic findings of	
	each (e.g., inflammatory pseudotumor, lymphoma, artifacts of	
	tissue processing, virus particles).	
3.	Describe the histology of the less common but potentially vision	C2
	or life-threatening ocular and adnexal diseases (e.g., healed giant	
	cell arteritis, mimics and masqueraders of inflammation and	
	neoplasm, less common benign and malignant neoplasms).	



4. Describe ancillary procedures for oncology (e.g., bone marrow	C2
aspiration, cerebrospinal fluid cytology).	

1.	Manage consultation between the clinician and ophthalmic pathologist regarding indications for special stains (e.g., Gram stain for bacteria, Congo red for amyloid; Gomori methenamine silver staining for fungi; Prussian blue for hemosiderosis; von Kossa for calcium; Oil Red O or Sudan Black for sebaceous	P2	3
	carcinoma) or processing (e.g., orientation of specimen, special handling).		
	Participate as an observer during the microscopic examination of active ophthalmology cases, including more advanced stains and techniques.		3
3.	Participate in subspecialty clinical pathological meetings (e.g., with corneal surgeons, infection specialists, tumor board).		4
4.	Handle appropriately gross or cytologic specimens in the ophthalmic pathology laboratory (e.g., vitreous biopsy, exenteration specimen).		3
5.	Prepare more advanced histologic specimens for review by the ophthalmic pathologist (e.g., special stains or fixation methods such as glutaraldehyde fixation for electron microscopy).		3
6.	Perform microscopic examination of a paraffin- embedded specimen and a frozen-section specimen without direct supervision; provide a relevant differential diagnosis; draft a report-preferably previewing slides in advance of the pathologist-to come up with a diagnosis and to suggest special stains and immunohistochemistry, without the influence of the ophthalmic pathologist; review the report and special stain orders with the ophthalmic pathologist.		
7.	Participate with the ophthalmic pathologist in tumor board and similar multidisciplinary meetings, presentations on recent advances, and journal clubs involving pathology.		4





8. Research requirement: Publish at least one paper based on basic, translational, or clinical research involving ophthalmic pathology. The purpose of the requirement is to further the trainee's in-depth knowledge of pathophysiology and laboratory techniques relating to ophthalmic pathology.

### **Oculoplastic Surgery and Orbit**

- A. Cognitive Skills
  - General

1.	Describe the clinical features, evaluation, and management of congenital syndromes, inflammation, trauma, ectropion, entropion, trichiasis, blepharoptosis, eyelid retraction, epiblepharon, dermatochalasis, blepharochalasis, eyelid tumors, blepharospasm, facial nerve palsy, eyebrow, midface and lower face function; and aesthetics, histology, and pathology of the facial skin.	C2
2.	Describe ocular surface pathology, including cicatricial processes affecting the bulbar and palpebral conjunctiva, management of corneal and conjunctival exposure, and relationship of the lids, midface, and brow to ocular exposure.	C2
3.	Describe the assessment of eyebrow position for brow ptosis and paralysis, and determine its relation to upper eyelid dermatochalasis.	C2
4.	Describe complex eyelid trauma.	C2
5.	Describe complex eyelid reconstruction (e.g., Hughes flap, free tarsal grafts, local flaps, skin grafts, Cutler-Beard procedure).	C2

#### Eyelid

#### Lacrimal

1. Describe the etiology, evaluation, and medical and surgical	C2
treatment of congenital tearing, acquired tearing, and trauma.	





1.	Describe the etiology, evaluation, and medical and surgical	C2
	treatment of orbital problems of children (e.g., congenital	
	anomalies, cellulitis, benign and malignant tumors, and orbital	
	inflammations).	
2.	Describe the etiology, evaluation, and medical and surgical	C2
	treatment of orbital disorder of adults, including orbital	
	cellulitis, thyroid orbitopathy, idiopathic orbital inflammation,	
	vasculitis, congenital tumors, vascular tumors, neural tumors,	
	lacrimal gland tumors, fibro-osseous tumors, histiocytic	
	diseases, lymphoid tumors, metastatic tumors, blunt and	
	penetrating trauma, orbital and facial fractures, anophthalmic	
	socket problems, and skull base disease.	
3.	Describe the types of and indications for various biomaterials	C2
	and orbital implants.	

### Nose

1.	Describe basic anatomy and physiology.	C2

### Sinuses

1. Describe basic anatomy and physiology.	C2	
Head and Neck as it Relates to the Orbit and Adnexa		
1. Describe basic anatomy and physiology.	C2	

### B. Technical skills

## Eyelid

1.	Describe indications for and perform medical	P2	4
	and surgical treatment of floppy eyelid		
	syndrome.		
2.	Perform more complicated eyelid procedures,		3
	including:		
	a. Levator advancement		
	b. Retractor reinsertion		
	c. Lower eyelid elevation		
	d. Upper eyelid recession		
	e. Eyebrow elevation		
	-		



3.	Perform complex ptosis repairs (e.g.,	3
	reoperations for height or contour	
	abnormalities).	
4.	Perform complex lower eyelid procedures (e.g.,	4
	retraction using a spacer, cicatricial entropion	
	using a mucous membrane graft).	3
5.	Perform midface surgery (e.g., midface lift for	
	cicatricial and paralytic ectropion).	3
6.	Perform advanced brow elevation techniques	
	(e.g., endoscopic, pretrichial, coronal).	
7.	Perform advanced eyelid reconstruction (e.g.,	3
	Hughes flap, Cutler-Beard procedure, tissue	
	transfer, flaps, and grafts).	2
8.	Perform cosmetic upper blepharoplasty.	2
9.	Perform cosmetic lower blepharoplasty.	3
10.	Excise benign and malignant tumors involving	
	the periorbital and adjacent regions.	

### Lacrimal

1.	Treat lacrimal system abnormalities, including:	P2	4
	a. Complex congenital disorders (e.g., canalicular stenosis)		
	<ul> <li>b. Complex trauma (i.e., requiring lacrimal intubation)</li> </ul>		
2.	Describe indications for and complications of, and perform intranasal endoscopic examination.		3
3.	Describe management of complex acquired disorders and their treatment (e.g., external and endoscopic dacryocystorhinostomy, conjunctivodacryocystorhinostomy with Jones tube).		3

### Orbital

1. perfor	m basic	orbital	skills	and	procedures,	P2	2
includ	ing:						
a. Socket reconstructions (e.g., tissue transfers,							
grafts, flaps, synthetic implants)							



b.	Fracture repair of bones involving the	
	periorbital region and orbit (e.g., orbital	
	floor, medial orbital wall, Le Fort,	
	zygomaticomaxillary complex [ZMC], naso-	
	orbito-ethmoid [NOE])	
C.	Orbitotomy for exploration, biopsy, and	
	tumor removal using anterior, lateral,	
	medial, and superior approaches; and orbital	
	reconstruction	
a.	Enucleation, evisceration, exenteration, and	
	secondary implants of the orbit	
e.	Complex or difficult socket-related problems	
	and complications (e.g., extrusion of	
	implants, contracted socket, anophthalmic	
	enophthalmos)	
f.	Optic nerve sheath fenestration	
	•	
g.	Orbital decompression for thyroid eye	
	disease	

### Nasal

1.	Describe nasal endoscopy as related to the P2	2
	management of lacrimal and periorbital processes.	
2.	Describe turbinectomy and nasal surgery as related	
	to the management of lacrimal and periorbital	
	processes.	

### Head and Neck

	P2	2
<ol> <li>Repair upper face and brow conditions, including brow ptosis repair.</li> </ol>		
<ol> <li>Use neuromodulators (e.g., botulinum toxin), dermal fillers, other technologies (e.g., laser) and chemical/pharmaceutical agents for the management of contour and skin quality abnormalities (i.e., functional and aesthetic).</li> </ol>		3

## Pediatric Ophthalmology and Strabismus



1.	Describe and perform the most advanced strabismus examination	C2
	techniques (e.g., complicated prism cover testing in multiple	
	cranial neuropathies, patients with nystagmus, dissociated vertical	
	deviation, double Maddox rod testing).	
2.	Describe clinical application of the most advanced sensory	C2
	adaptations (e.g., anomalous head position, anomalous retinal	
	correspondence, methods of distance stereopsis).	
3.	Describe etiologies of amblyopia (e.g., refraction noncompliance,	C2
	patching failures, and pharmacologic penalization).	
4.	Describe etiologies of esotropia (e.g., optical; postrefractive	C2
	surgical esotrophia [ET]; prism-induced ET decompensated	
	esophoria; postsurgical amd consecutive ET; sixth nerve palsy and	
	paresis; thyroid eye disease, following closed head injury; Chiari	
	malformation).	
5.	Describe the etiologies of exotropia (e.g., supranuclear, paralytic	C2
	pontine exotropia, consecutive).	
6.	Identify complex ROP (e.g., stages, treatment indications, retinal	C2
	detachment).	
7.	Describe causes and testing of optic atrophy in children.	C2
8.	Describe methods of ocular assessment of children with other	C2
	disabilities.	
9.	Describe ocular cysticercosis.	C2
10.	Describe screening strategies for childhood blindness at the	C2
	community level and intervention.	
11.	. Describe how to guide/refer parents of children with severe vision	C2
	impairment.	

<ol> <li>Perform more complex extraocular muscle surgery (e.g., vertical and horizontal muscle surgery, including superior oblique procedures, transpositions, and reoperations).</li> </ol>	3
<ol> <li>Perform preoperative assessment, intraoperative techniques, and describe postoperative complications for more complicated strabismus surgery (e.g., reoperations, stretched scar, slipped muscle, lost muscle).</li> </ol>	4
······································	3



<ul> <li>3. Perform adjustable sutures in more complicated cases (e.g., thyroid ophthalmopathy).</li> <li>4. Manage more complex complications of strabismus surgery (e.g., globe perforation, corneal dellen, inclusion cysts, endophthalmitis, and overcorrection undercorrection).</li> <li>5. Perform surgery of congenital cataract including posterior polar cataract (PPC), vitrectomy with/without intraocular lens implantation, persistent hyperplasia of the primary vitreous (PHPV)/persistent fetal vasculature (PFV), including biometric measurements to determine aphakia contact lens or intraocular lens.</li> <li>6. Perform glaucoma surgery in pediatric and congenital glaucoma.</li> <li>7. Perform corrective surgery in congenital eyelid anomalies like ptosis.</li> <li>8. Perform nasolacrimal surgery in children.</li> <li>9. Perform electromyography (EMG) guided or intraoperative injection of botulinum toxin for strabismus.</li> <li>4</li> <li>10. Diagnose ROP and refer for treatment.</li> <li>11. Perform more complex strabismus procedures (e.g., Faden sutures, posterior myopexy, Yokoyama muscle union, "Y" splitting).</li> </ul>			
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	11.	. Perform more complex strabismus procedures	
Yokoyama muscle union, "Y" splitting).		(e.g., Faden sutures, posterior myopexy,	3
		Yokoyama muscle union, "Y" splitting).	

### Vitreoretinal Diseases

1.	Diagnose, evaluate, treat (or refer) the most complex forms of	С3
	retinal vascular diseases and diagnose/manage risk factors (e.g.,	
	blood dyscrasia) and systemic complications.	
2.	Diagnose, evaluate, and treat inherited, congenital, and acquired	C3
	macular diseases.	
3.	Compare the current therapeutic retinal treatment strategies and	C2
	be able to discuss the future improvements of the therapeutic	
	armamentarium.	
4.	Evaluate and treat traumatic injuries to the retina, including	С3
	complex cases such as intraocular foreign body with	





	rhegmatogenous retinal detachment and traumatic macular	
	holes, and be able to manage complications to the other ocular	
	structures.	
5.	Diagnose, evaluate, and understand the genetic alterations and	C3
	the possible applications of gene therapy for hereditary diseases.	
6.	Develop surgical proficiency in different surgical techniques for	C2
	management of retinal detachment, including complex cases	
	(e.g., combined rhegmatogenous/tractional retinal	
	detachments).	

1.	<ul><li>Perform posterior photocoagulation in complicated retinal cases:</li><li>a. Retinal breaks with vitreous hemorrhage</li><li>b. Cases with intraocular tamponade (i.e., gas,</li></ul>	P2	3
	silicone oil)		
2.	Interpret and apply electrophysiology in clinical practice.		3
3.	Interpret and apply ocular imaging techniques in		
	clinical practice (e.g., B-scan echography) and in		
	more complex cases (e.g., choroidal osteoma).		4
4.	Perform detailed fundus drawings of the retina with		
	vitreoretinal relationships in the most complex		
	retinal cases (e.g., recurrent retinal detachment,		4
	retinoschisis with and without retinal detachment).		
5.	Perform laser therapy or cryotherapy of retinal		
-	holes and other more complex retinal pathology.		
6.	Perform scleral buckling in complex retinal detachment.		3
7.	Perform advanced pars plana vitrectomy.		3
			3

## **Uveitis and Ocular Inflammation**





1.	Describe the clinical features and differential diagnoses for less	C2
	common forms of uveitis (e.g., Whipple disease, Crohn disease,	
	bilateral acute depigmentation of the iris [BADI], diffuse unilateral	
	subacute neuroretinitis [DUSN], and onchocerciasis.	
2.	Describe the global epidemiology of uveitis and relate this	C2
	information to the diagnosis.	
3.	Describe the management of the more complex complications of	C2
	uveitis.	
4.	Describe indications for ultrasound biomicroscopy (e.g., assess	C2
	state of ciliary body in hypotony), laser flare photometry and	
	electrophysiology in the evaluation of uveitis.	
5.	Describe indications, contraindications, and complications for	C2
	immunosuppressive therapy in uveitis (e.g., use of	
	antimetabolites, cyclosporine, alkylating agents, biologic agents).	
6.	Describe indications, contraindications, and complications of	C2
	retinal laser photocoagulation in uveitis.	
7.	Describe indications, contraindications, and complications of	C2
	intravitreal injection of medications (e.g., corticosteroids, antiviral	
	therapy, antibiotics, anti-VEGF, anti-mitotic agents) and drug	
	delivery systems (e.g., for corticosteroid, ganciclovir).	

1.	Integrate history, clinical examination, and investigations in order to evaluate the less common uveitis entities.	P2	4
2.	Administer corticosteroids in the treatment of uveitis by various routes (e.g., topical, periocular, systemic, and intravitreal injection).		4
3.	Perform retinal laser photocoagulation for retinal vasculitis complicated by retinal capillary nonperfusion and associated retinal or optic disc neovascularization.		3
4.	Regulate perioperative management of the uveitic eye for cataract removal.		4
5.	Perform intravitreal injection of medications (e.g., corticosteroids, antiviral therapy, antibiotics, anti- VEGF, antimitotic agents) and drug delivery systems (e.g., for corticosteroid, ganciclovir).		4



6. Co-manage with another subspecialist as	4
appropriate:	
a. Biopsy of the vitreous, retina, or choroid to	
confirm/exclude vitreoretinal lymphoma or	
other tumors/infectious causes	
b. Immunosuppressive therapy in uveitis	
including biologics (with or without the aid	
of an immunologist) and monitor for side	
effects Intravitreal implants containing	
antiviral or corticosteroid medications.	
Ocular complications of uveitis (e.g.,	
macular edema, cataract, glaucoma, retinal	
detachment, band keratopathy, choroidal	
neovascularization, hypotony).	

## Ocular Oncology

1. Describe the applied surgical anatomy, histology, and embryology	C2
of the eye and ocular adnexa with relevance to ocular oncology.	
2. Describe the applied physiology of the eye and adnexa with	C2
relevance to ocular oncology.	
3. Describe the applied pathology of the following:	C2
3.1 Ocular tumors and pseudotumor	
3.1.1 Congenital/developmental	
a. Conjunctiva	
i. Dermoid	
ii. Dermo lipoma	
iii. Choristoma (simple and complex)	
b. Uvea	
i. Lisch nodules	
ii. Stromal iris cyst	
iii. Lacrimal gland choristoma	
c. Retina	
i. Multiple congenital hypertrophy of the retinal	
pigment epithelium	
ii. (CHRPE)	
iii. Astrocytic hamartoma	



- iv. Hemangioblastoma
- v. Cavernous angioma
- vi. Dominant exudative vitreoretinopathy
- vii. Norrie disease
- viii. Incontinentia pigmenti
- ix. Solitary CHRPE
- x. Grouped pigmentation
- xi. Arteriovenous malformation (racemose angioma)
- xii. Posterior primary hyperplastic vitreous (PPHV)
- xiii. Glioneuroma
- 3.1.2 Inflammatory (infectious, noninfectious)
  - a. Conjunctiva
    - i. Granuloma (e.g., syphilis, sarcoid)
  - b. Uvea
    - i. Granuloma (e.g., tuberculosis) Uveal effusion
    - ii. Posterior scleritis
  - c. Retina
    - i. Granuloma (e.g., toxocara)
- 3.1.3 Benign
  - a. Conjunctiva
    - i. Nevus
    - ii. Papilloma
    - iii. Oncocytoma
    - iv. Primary acquired melanosis
    - v. Reactive lymphoid hyperplasia
    - vi. Other
  - b. Uvea
    - i. Nevus/melanocytoma
    - ii. Hemangioma
    - iii. Osteoma
    - iv. Neurilemmoma
    - v. Neurofibroma
    - vi. Leiomyoma
    - vii. Mesectodermal leiomyoma
    - viii. Reactive lymphoid hyperplasia
    - ix. Bilateral diffuse uveal melanocytic proliferation
    - x. Other rare conditions
  - c. Retina
    - i. Retinoma/retinocytoma
    - ii. Adenoma
    - iii. Fuchs adenoma



- iv. Benign medulloepithelioma
- v. Other
- 3.1.4 Malignant
  - a. Conjunctiva
    - i. Melanoma
    - ii. Squamous cell carcinoma
    - iii. Sebaceous carcinoma
    - iv. Kaposi sarcoma
    - v. Lymphoma
    - vi. Extraocular tumor spread
    - vii. Metastasis
    - viii. Other
  - b. Uvea
    - i. Melanoma
    - ii. Lymphoma
    - iii. Intraocular tumor spread from conjunctiva
    - iv. Systemic lymphoma
    - v. Systemic leukemia
    - vi. Metastasis
    - vii. Other
  - c. Retina
    - i. Retinoblastoma
    - ii. Adenocarcinoma
    - iii. Malignant medulloepithelioma
    - iv. Lymphoma
    - v. Leukemia
    - vi. Metastasis
    - vii. Other
- 3.1.5 Traumatic
  - a. Conjunctiva
    - i. Implantation cyst
    - ii. Foreign body granuloma
    - iii. Pyogenic granuloma
  - b. Uvea
    - i. Implantation cyst
    - ii. Choroidal hemorrhage
    - iii. Miotic cyst
  - c. Retina
    - i. Retinopathy of prematurity
    - ii. Retinal detachment
    - iii. Massive reactive gliosis



216 Dece			
3.1.6 Degenerative			
a.	Conjunctiva		
	i. Lacrimal retention cyst		
b.	. Uvea		
	i. Disciform lesion		
	ii. Sclerochoroidal calcification		
	iii. Vortex vein ampulla		
с.	Retina		
	i. Vasoproliferative tumor		
3.1.7	Idiopathic		
a.	Conjunctiva		
	i. Lymphangiectasia cyst		
b.	. Uvea		
	i. Juvenile xanthogranuloma		
C.			
	ii. Coats disease		
	iii. Combined hamartoma of retina and retinal		
	pigment epithelium		
	iv. Iris cyst		
	v. Ciliary epithelial cyst		
3.1.8	Paraneoplastic disease		
a.	Bilateral diffuse uveal melanocytic proliferation		
b.	Carcinoma-associated retinopathy		
C.			
d.	Other		
	tibe the following pathological conditions: C2		
	Non-neoplastic tumors		
	i. Hamartoma		
	ii. Choristoma		
	iii. Granuloma		
	iv. Cyst		
	v. Hyperplasia		
	vi. Metaplasia		
h	Neoplastic tumors		
υ.	i. Benign		
	ii. Malignant		
C.			
c. d.			
	Seeding Metastasic		
f.	Metastasis		



	g.	latrogenic disease	
	h.	Radiation	
	i.	Pharmacology	
	j.	Surgery	
	k.	Phototherapy	
5.	Descri	be relevant pathological techniques, such as:	C2
	a.	Fixatives	
	b.	Frozen sections	
	с.	Histology	
	d.	Immunohistochemistry	
	e.	Flow cytometry	
	f.	Other	
6.	Descri	be the following genetic abnormalities and techniques:	C2
	a.	Germinal mutations relevant to oncology	
	b.	Somatic mutations in tumors	
	с.	Genetic techniques	
		i. Karyotyping	
		ii. Polymerase chain reaction	
		iii. Fluorescence in situ hybridization	
		iv. Multiplex ligation-dependent probe amplification	
		v. Gene expression profiling	
		vi. Comparative genomic hybridization	
		vii. Other	
7.		be the relevant staging and grading systems for ocular	C2
		s (with ability to use appropriate methods as necessary,	
	using	appropriate references sources):	
	a.	TNM Classification of Malignant Tumors cancer staging	
		system	
		i. Uveal melanoma	
		ii. Retinoblastoma	
		iii. Conjunctival melanoma	
		iv. Conjunctival carcinoma	
	h	v. Ocular adnexal lymphoma	
	υ.	International retinoblastoma staging system vi. Ocular adnexal lymphoma	
	c		
	c. d.	International retinoblastoma staging system Reese-Ellsworth staging system for retinoblastoma	
	-	Other staging systems (e.g., Collaborative Ocular	
	с.	Melanoma Study)	
8.	Descri	be the etiology of ocular tumors:	C2
0.	a.	Environmental factors	
		-	



	11
b. Genetic factors	
c. Syndromes	
d. Malformations	
e. Other	
9. Describe the pathogenesis of ocular tumors:	C2
a. Secondary effects of uveal melanoma	
b. Secondary effects of retinoblastoma	
c. Secondary effects of other tumors (e.g., conjunctival	
tumors)	
10. Describe the epidemiology of ocular tumors:	C2
a. Principles of epidemiology	
11. Describe the principles of examination techniques:	C2
a. Inspection	
i. Slit-lamp examination	
ii. Gonioscopy and 3-mirror examination	
iii. Ophthalmoscopy	
b. Transillumination	
i. Transpupillary	
ii. Transscleral	
c. Color photography	
i. Standard ocular photography	
ii. Specialized cameras (e.g., RetCam, Optos)	
iii. Autofluorescence photography	
d. Angiography	
i. Fluorescein angiography	
ii. Indocyanine green angiography	
e. Ultrasonography	
i. A-scan ultrasonography	
ii. B-scan ultrasonography (including high frequency)	
iii. Doppler ultrasonography	
f. Magnetic resonance imaging	
g. Computerized tomography	
h. Positron emission tomography	
i. Biopsy	
ii. Aspiration	
iii. Incisional	
iv. Excisional	
v. Impression cytology	
i. Systemic investigation according to ocular tumor	
diagnosis	
i. History	
1. TIISCOLY	



ii. Clinical examination	
iii. Hematology and biochemistry	
iv. Radiography	
v. Ultrasonography	
vi. Computerized tomography	
vii. Magnetic resonance imaging	
viii. Genetic testing	
12. Describe the clinical features of each tumor type:	C2
a. Inspection/color photography	
<ul> <li>Investigational (i.e., angiography, echography)</li> </ul>	
13. List the differential diagnosis of each tumor and describe the	C2
investigational approach for each condition.	
14. Describe how the following therapeutic modalities and their	C2
effects are relevant to ocular tumors:	
a. Radiotherapy	
i. Radiation	
ii. Radioactive sources (e.g., iodine, ruthenium)	
iii. Types of radiation (e.g., gamma, beta, proton)	
iv. Biological effects	
b. Chemotherapy	
c. Phototherapy	
d. Cryotherapy	
e. Surgical resection	
15. Describe how the following statistics can be applied to ocular	C2
oncology:	
a. Statistical correlations	
i. Univariate	
ii. Multivariate	
b. Survival statistics	
i. Kaplan-Meier analysis	
ii. Cox analysis	
iii. Neural networks	
iv. Accelerated failure time	
c. Bias	
d. Power calculations	
e. Other relevant statistical methods	
B Technical skills	





1.	Perform or request the following examinations,	P2	4
	interpreting and documenting any findings,		
	demonstrating awareness of the indications,		
	contraindications, and limitations of each		
	investigation:		
	a. Slit-lamp examination of conjunctiva and		
	assessment of conjunctival fornices		
	b. Slit-lamp examination of anterior chamber		
	and gonioscopy		
	c. Binocular indirect ophthalmoscopy with		
	indentation		
	d. Transpupillary transillumination		
	e. A-scan and B-scan ultrasonography of		
	anterior and posterior eye		
	f. Color and autofluorescence photography		
	g. Fluorescein angiography		
	h. Indocyanine green angiography		
	i. Magnetic resonance imaging		
	j. Incisional and excisional conjunctival		
	tumor biopsy		
	k. Aspiration, incisional, or excisional biopsy		
	of intraocular tumor		
	I. Other relevant examinations and		
	investigations		
2.	Perform or refer for the following treatments for		
	conjunctival tumors, demonstrating awareness of		4
	the indications, contraindications, and		
	complications of each treatment:		
	a. Surgical excision		
	b. Cryotherapy		
	c. Brachytherapy		
	d. External beam radiotherapy, including		
	proton beam radiotherapy		
	e. Topical therapy (e.g., mitomycin C, 5-		
	fluorouracil, interferon)		
3.	Perform or refer for the following treatments for		
	intraocular tumors, demonstrating awareness of		
	the indications, contraindications, and		4
	complications of each treatment:		
	a. Radiotherapy		



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i. Brachytherapy (e.g., iodine,	
ruthenium, strontium, palladium,	
iridium)	
ii. External beam radiotherapy	
iii. Stereotactic radiotherapy	
iv. Charged particle radiotherapy	
(e.g., proton beam)	
v. Phototherapy	
vi. Photocoagulation	
vii. Transpupillary thermotherapy	
viii. Photodynamic therapy	
b. Surgical excision	
i. Iridectomy	
ii. Iridocyclectomy	
iii. Transscleral choroidectomy	
iv. Transretinal choroidectomy	
v. Enucleation	
vi. Exenteration	
c. Ocular pharmacological therapy by	
various routes (i.e., topical, intravitreal,	
ophthalmic artery infusion, subtenon,	
systemic)	
i. Chemotherapy and biological	
therapy	
ii. Antiangiogenic agents	
iii. Steroids	
4. Request the following investigations, interpreting	
and communicating the results to patients,	4
relatives, and health care workers, adjusting	
patient management accordingly:	
a. Histopathological assessment of tumor	
samples	
b. Genetic assessment of tumor samples	
c. Laboratory investigation of vitreous	
samples	
d. Other	
5. Estimate the prognosis and communicate the	
following implications with patients, relatives,	4
and health care workers, adjusting patient	
management accordingly:	
a. Visual acuity	



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b. Local tumor control	
c. Possible side effects and complications of	
therapy	
d. Ocular conservation	
e. Systemic manifestations of disease,	
including metastasis	
f. Systemic complications and side effects of	
therapy	
g. Survival probability and chances of	
disease-related mortality	
h. Heritability	
i. Use information technology and	
other aids to enhance	
prognostication	
6. Communicate the following to patients, relatives,	
and health care workers:	
a. Diagnosis, extent and severity of disease,	
including diagnostic uncertainty	4
b. Natural history without treatment	
c. Therapeutic options with advantages and	
limitations of each therapy, including	
methods available elsewhere	
d. Logistical implications of selected	
treatment	
e. Use information technology and other	
aids to support this process	
i. Websites	
ii. Printed leaflets	
iii. Audio recordings	
f. Other relevant materials	
7. Assist patients with selecting the most	
appropriate management, taking into account:	
a. Patient age, gender, culture, wishes,	
needs, and fears	4
b. Costs and logistics	
c. Availability of health care resources,	
locally and elsewhere	
8. Provide or organize appropriate psychological	
support, demonstrating empathy and an	
adequate awareness of the principles of this	4
aspect of care, such as:	<b>4</b>



a. Giving bad news	
b. Communicating with relatives	
c. Enabling long-term communication a	nd
support	
9. Develop and maintain a multidisciplinary team	of
health care professionals to provide patie	nt
focused care by activities, such as:	
a. Recruiting staff and coworkers	4
b. Developing service operating procedure	s.
c. Maintaining efficient and varied metho	ds
of communication and education	
i. Between multidisciplinary tea	Im
members (MDT)	
ii. Between MDT and oth	er
practitioners (e.g., pathologists)	
iii. Between MDT and patient	
10. Develop protocols and infrastructure for practic	ce-
based learning and improvement, including:	
a. Proformas and databases for storing data	ta
b. Protocols for extracting and analyzing da	ta <b>4</b>
c. Application of study designs and statistic	cal
methods	
d. Adherence to clinical governance	
i. Informed consent	
ii. Confidentiality	
iii. Ethical committee approval	

### Low Vision Rehabilitation

1. Describe the process of complex rehabilitation, including:	C2
a. Optical rehabilitation	
b. Non optical aids	
c. Eccentric fixation training and scotoma avoidance	
d. Orientation and mobility	
e. Activities of daily living	
f. Vision substitution (e.g., touch, hearing)	
g. Psychological care	
	C2





2.	Describe the role of all of the partners and team members in the	
	patient's care and in low vision rehabilitation (e.g.,	
	ophthalmologists, social workers, psychologists, rehabilitation	
	trainers).	C2
3.	Describe the main aims and projects of VISION 2020.	
4.	Describe the effects of low vision on the general health and on	C2
	the psychological wellbeing of the patient.	
5.	Describe the concept of artificial vision and implantation of	
	microchips for the treatment of patients with the most profound	C2
	visual impairments.	
6.	Describe a low-vision-friendly physical environment that	
	includes easy accessibility (e.g., ergonomics, special visual signs	
	in buildings/streets, talking elevators/traffic signs).	

1.	Identify basic low vision and other surgical and	P2	4
	medical interventions necessary to ensure the		
	best possible visual outcome.		
2.	Oversee and provide referrals to support the		4
	patient's psychological adjustment to life after		
	acute vision loss.		
3.	Educate patients on use of low vision equipment.		4
4.	Be well informed and instruct patients with low		
	vision of comprehensive rehabilitation resources		4
	in the region and in the country, including		
	offering provider contact details.		
5.	Interact with other professionals (e.g.,		
	psychologists, occupational therapists,		4
	vocational counselors, social workers) to		
	improve the daily life of patients with low vision.		

## Ethics and Professionalism in Ophthalmology

1. Apply advanced medical ethics in the ophthalmic practice:	C3
<ul> <li>a. Applicable informed consent documents (e.g., clinical research, off-label use disclosures)</li> </ul>	



b. Managemen	t (offering and rendering) of second opi	nions
-		
	and institutional responsibilities re	garoing
impaired phy	vsicians	
d. Responsibilit	y for postoperative care, including appr	opriate
transfer of ca	are to other physicians	
e. Managemen	t of conflicts of interest (clinical and non	clinical)
iii. Disclo	osures	
iv. Gifts	to physicians	
	advertising (and applicable laws)	
	opriate conduct as a medical-expert wit	tness in
litigat	ion	
2. Identify applicable in	nsurance coverage responsibilities in a p	practice C2
situation.		
3. Apply more advance	ed aspects of health care reimburseme	ent in a C3
clinical practice (e	g., denials of claims, hospital cont	racting.
electronic billing).		
	going ethical principles and knowledge i	n diract
	going ethical principles and knowledge i	nunect
patient care.		
5. Describe the respo	nsibility of ophthalmologists to shar	e their
knowledge of clinica	I arts and sciences for the benefit of p	atients, C2
the profession, and	society.	

## **Community Eye Health**

1.	Describe the principles of epidemiology, as applicable to community eye health.	C2
2.	Describe the principles of research methods, as applicable to community eye health.	C2
3.	Describe the principles of biostatistics, as applicable to community eye health.	C2
4.	Describe the principles of health economics, as applicable to community eye health.	C2
5.	Describe the principles of health systems strengthening, as applicable to community eye health.	C2
6.	Describe the principles of health education and health promotion, as applicable to community eye health.	C2
7.	Describe the principles of project and program management, as	C2
	applicable to community eye health.	





8. Describe the relevant WHO global programs (eg, millennium	C2
development goals, disability framework).	
9. Describe the relevance of the disability policy at a global level and	C2
within the health system.	
10. Describe the main concepts of habilitation, rehabilitation, and	C2
community-based rehabilitation for persons with visual disability	
and its integration within a health system.	

In addition to the technical skills listed for residency	P2	4
training, be able to:		
1. Plan and conduct research projects to inform the		
planning and implementation of district and		
national blindness prevention programs.		
2. Plan and conduct RAAB surveys.		
3. Plan and conduct RAT surveys.		
4. Plan, implement, and manage one-year district		
operational blindness prevention programs.		
5. Plan, implement, and manage national three-to-		
five-year strategic blindness prevention programs.		
6. Advocate for national policy implementation and		
community participation to strengthen national		
blindness prevention programs.		
7. Provide training in community eye health to		
different eye care cadres.		
8. Engage with public health practitioners to advocate		
for improvements in eye care services and the		
implementation of the disability framework.		
9. Assess the impact of disabilities and advocate the		
application of global disability policy at a local level.		



# **SECTION IV: TEACHING AND LEARNING STRATEGIES**







## Overview

Teaching and learning strategies in the MS Ophthalmology program are designed to create a comprehensive and engaging educational experience that meets the diverse needs of postgraduate residents. These strategies focus on developing clinical competence, surgical skills, critical thinking, and professional attitudes essential for successful ophthalmologists. A well-defined outline of these methods in the curriculum is crucial for standardizing resident training, ensuring consistency, and providing clear expectations for both educators and learners

•	npatient Services
•	Dutpatient/ambulatory Experiences
• (	Core Faculty Lectures (CFL)
• J	ournal Club Meeting (JC)
• (	Case-Based Learning CBL
• (	Grand Rounds (GR)
• (	Clinico-pathological Conferences CPC
• (	Clinical/Surgical Audit Based Learning
• F	Peer-Assisted Learning PAL
	Morbidity meeting (MM)
• 9	Skills Workshops
	Multidisciplinary Team-based Learning MDL
• •	Simulation Training
•	-learning and Online Resources

## **Outline of Different Teaching and Learning Strategies**

### **Inpatient Services**

Residents will rotate through various inpatient services to gain comprehensive experience.

### **Outpatient/Ambulatory Experiences**

Residents will demonstrate expertise in diagnosing and managing patients in acute care clinics and longitudinal clinics, gaining experience in various subspecialties

### **Core Faculty Lectures (CFL)**

Core faculty lectures will focus on monthly themes covering various specialty topics. Lectures will incorporate active learning techniques such as buzz groups.

### Journal Club Meeting (JC)



Residents will present and critically evaluate research articles, highlighting applicable results for clinical practice.

### **Case-Based Learning CBL**

Small groups will engage in case-based learning. This method emphasizes problem-solving skills and integrated knowledge.

### Grand Rounds (GR)

Weekly grand rounds will feature speakers from local, regional, and national training programs, presenting topics from the broad spectrum of topics.

### **Clinico-pathological Conferences CPC**

Using case methods, these conferences will involve discussing differential diagnosis, diagnostic data, and final diagnoses.

### **Clinical/Surgical Audit Based Learning**

Residents will participate in quality improvement processes by reviewing patient care against explicit criteria and implementing necessary changes.

### **Peer-Assisted Learning PAL**

Residents will engage in peer-assisted learning, providing opportunities for reinforcement, responsibility, self-confidence, and development of teaching and communication skills.

### Morbidity meeting (MM)

Adverse outcomes, not necessarily resulting in death, will be discussed and thoroughly reviewed.

### **Skills Workshops**

Conducting skills workshops on surgical techniques, diagnostic procedures, and equipment handling. Provide opportunities for students to practice and receive feedback on their skills.

### Multidisciplinary Team-based Learning MDL

Collaborating with other healthcare professionals to simulate a multidisciplinary team approach to patient care. Encourage students to understand the roles of different team members and practice effective communication and teamwork.

### **Simulation Training**

Utilizing simulation training tools and platforms to provide a realistic and safe environment for students to practice complex procedures like phacoemulsification. Incorporate simulation scenarios that mimic challenging clinical situations to enhance decision-making skills.

### E-learning and Online Resources/Digital Library

Integrated e-learning modules, online resources, and virtual simulations to supplement traditional teaching methods providing access to online databases, journals, and educational videos to support self-directed learning.

Encourage use of digital library available at RMU.



## **Teaching Schedule**

In addition to bedside teaching rounds, in the department there will be daily hourly sessions of formal teaching per week. The suggested time distribution of each session for department's teaching schedule as follows:

- Journal club Once a week
- Seminar once a week
- PG case discussion Twice a week
- Audit/Morbidity meeting Once a month
- Central session as per hospital schedule
- ➢ Workshop − once every 3 months

All sessions are supervised by faculty members. It is mandatory for all residents to attend the sessions except those posted in emergency.

All the teaching sessions are assessed by the faculty members at the end of session and marks are given out of 10 and kept in the office for internal assessment.

Attendance of the residents at various sessions has to be at compulsory.



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# **SECTION V: WORKSHOPS**

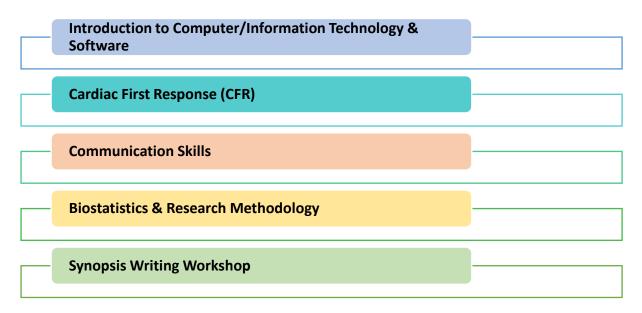


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## Framework of Workshops in MS Ophthalmology Program

The MS Ophthalmology Residency Program includes a comprehensive workshop series designed to equip residents with critical skills in research methodology, computer literacy, communication, medical ethics, and emergency response. These workshops are integrated across the four-year residency to ensure incremental skill development and practical application. Each workshop has specific learning objectives and topics that address various aspects essential for clinical practice, patient interaction, research acumen, and emergency preparedness.



### 1. Introduction to Computer/Information Technology & Software

- **Learning Objectives**: Develop fundamental IT skills for clinical and research applications, including basic word processing, data management, and presentation creation.
- **Topics Covered**: Hardware, software basics, file management, word processing, PowerPoint, Excel, email, internet navigation, and introductory data entry in statistical software (SPSS).

### 2. Biostatistics & Research Methodology

- Learning Objectives: Grasp basic biostatistics, understand the importance of research, develop a research question, and engage in scientific presentations.
- **Topics Covered**: Introduction to biostatistics, biomedical research principles, selecting research fields, ethics, writing and presenting papers, and literature search techniques

### 3. Communication Skills

- Learning Objectives: Enhance clinical communication skills, improve counseling techniques, and understand ethical responsibilities.
- **Topics Covered**: Non-medical interventions, crisis intervention, conflict resolution, breaking bad news, informed consent, patient confidentiality, and professional ethics.



### 4. Cardiac First Response (CFR)

- Learning Objectives: Train in emergency cardiac care, including BLS, AED usage, and ALS fundamentals.
- **Topics Covered**: Cardiac emergency recognition, BLS principles, AED usage, ALS basics, scenario-based training, team dynamics, and advanced resuscitation techniques

### 5. Synopsis Writing Workshop

- Learning Objectives: Develop and structure a research synopsis, from research question formulation to reference management.
- **Topics Covered**: Synopsis components, research question development, literature review, methodology, timeline and budget planning, peer review, editing, and finalizing the synopsis.

S.NO NAME OF THE WORKSHOP	LEARNING OBJECTIVES	TOPICS TO BE COVERED
<ol> <li>Biostatistics &amp; Research Methodology (4 days)</li> </ol>	<ul> <li>To understand the basics of Bio-Statistics</li> <li>To critique why research is important?</li> <li>To discuss the importance of Selecting a Field for Research</li> <li>To prepare oneself for Participationin National and International Research</li> <li>To prepare oneself for Participationin Pharmaceutical Company Research</li> <li>To interpret the importance of research ideas &amp; Criteria for a good research topic</li> <li>To discuss Ethics in Health Research</li> <li>To learn to write a Scientific Paper</li> <li>To learn to make a</li> </ul>	<ol> <li>Introduction to Bio-Statistics</li> <li>Introduction to Bio- Medical Research Whyresearch is important?</li> <li>What research to do?         <ol> <li>Selecting a Field for Research</li> <li>Drivers for Health Research</li> <li>Drivers for Health Research</li> <li>Participation in National and International Research</li> <li>Participation in Pharmaceutical Company Research</li> <li>Where do research ideas comefrom</li> <li>Criteria for a good research topicEthics in Health Research</li> </ol> </li> <li>Writing a Scientific Paper</li> <li>Making a Scientific Presentation &amp; Searching theLiterature</li> </ol>

### Learning Objectives of Workshops



Scientific	
Presentation	
• To learn to make a	
purposefulliterature	
search	



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			1.
2.	Introduction to	By the end of this workshop	2. Hardware and Software
	computer/Inform	student shouldbe able to:	Understand the main
	ation Technology	Appropriately	components of a computer,
	& Software(5	start up and shut	including input and output
	days)	down your	devices.
	uaysj	computer.	Understand the function of
		Navigate the	communicationdevices
		operating system and	such as smartphones and
		start applications.	tablets.
		Perform basic	
		functions of file	
			Operating Systems,
		management.	programs and apps.
		Perform basic	3. Windows
		functions in a word	Image: Turning on the computer and
		processor and	logging on.
		spreadsheet.	The Windows screen.
		Manage print	Running programs from the Start
		settings and print	Menu.
		documents.	Minimizing, maximizing,
		Receive and send email.	moving, resizing and closing
		<ul> <li>Use a web browser to</li> </ul>	windows.
		navigate theInternet.	Logging off and shutting down
		<ul> <li>work with windows,</li> </ul>	your computer.3.Working with
		toolbars, and	Programs
		command menus	<ul> <li>Running multiple programs.</li> </ul>
		<ul> <li>perform basic word</li> </ul>	Desktop icons and creating a
		processing and	desktop shortcut.
		graphic tasks	Imaging programs from the
		make a Power Point	taskbar.
		presentation	Closing programs.4.File
		explore Web browsing	Management
		basics	Managing Windows Explorer.
		<ul> <li>back up files</li> </ul>	Creating, moving, renaming and
		<ul> <li>save, copy, and organize</li> </ul>	deleting foldersand files.
		your work	Understandings file extensions.
		to enter data	<ul> <li>Viewing storage devices and</li> </ul>
		accurately in software	network connections.
		of Statistical Package	<ul><li>Managing</li></ul>
		for Social Sciences	USB flash drives.
			5.Word Processing
			<ul> <li>Creating documents in Microsoft</li> </ul>
			Word.
			<ul><li>Typing text, numbers and dates</li></ul>
			into a document.
			<ul><li>Easy formatting.</li></ul>
			Lasy ioi IIIattilig.



	and MEDICIN ST		di Medical University
		?	Checking the spelling in your
			document.
			Making and saving changes to your
			document.
		6.Power	Point
		Making	Power Point
		presenta	
		7.Spread	d sheets



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TO AT MEDICAL STREET	Rawalpindi Medical University
	<ul> <li>Understanding spreadsheet functionality.</li> <li>Creating spreadsheets in Microsoft Excel.</li> </ul>
	<ul><li>Typing text numbers and dates into a worksheet.</li><li>Easy formulas.</li></ul>
	<ul><li>Easy formatting.</li><li>Charting your data.</li><li>Making and saving changes to your</li></ul>
	workbook. • Printin
	<ul><li>g a worksheet.</li><li>8.Printing</li><li>Print preview.</li></ul>
	<ul> <li>Print settings.</li> <li>Managing</li> <li>the print queue.</li> </ul>
	<ul> <li>9.Using Email</li> <li>The Outlook mail screen elements.</li> </ul>
	<ul> <li>Composing and sending an email message.</li> <li>Manag</li> </ul>
	ing the Inbox. 10.Accessing
	<ul> <li>the Internet</li> <li>Going to a specific website and bookmarking.</li> </ul>
	<ul> <li>Understanding how to search/Google effectively.</li> <li>Copy and paste Internet</li> </ul>
	<ul><li>content into your</li><li>documents and emails.</li><li>Stopping and refreshing pages.</li></ul>
	<ul> <li>Demystifying the Cloud.</li> <li>Understanding social media platforms such asFacebook</li> </ul>
	<ul><li>and Twitter.</li><li>Computer security</li><li>best practices.</li></ul>
	11.Statistical Package for Social Sciences
	<ul> <li>general understanding for data entry</li> </ul>



			MEDICAC 4	Nawaipinai medicai oniversity
3.	communication	•	To learn to use	1. Use of Non-medicinal
	skills (3 days)		Non-medicinal Interventions in	Interventions in ClinicalPractice Communication Skills
		•	Communication Skills of Clinical Practice To discuss the importance ofcounseling To role play as a counselor	Counseling Informational Skills Crisis Intervention/Disaster Management Conflict Resolution

				RM	ر اولېنډى ميټيكل يونيورسټى
			MEDICAL ST	Rawalpin	ndi Medical University
		•	To learn to manage a conflict resolution To learn to break bad news To discuss the importance of Medical Ethics, Professionalism andDoctor-Patient Relationship Hippocratic Oath To learn to take an informed consent To illustrate the importance of confidentiality To summarize Ethical Dilemmas in a	6. 7. Doctor Oath 8. malefi 9. 10.	Adi Medical University Breaking Bad News Medical Ethics, Professionalism and r-PatientRelationship Hippocratic Four Pillars of Medical Ethics (Autonomy, Beneficence, Non- cence and Justice) Informed Consent and Confidentiality Ethical Dilemmas in a Doctor's Life
		Doctor			
4.	Synopses Writing		Introduction to	Introdu	iction to Synopsis Writing:
		Synops	is Writing and		
		Develo •	Understand the purpose and structure of a research synopsis.		<ul> <li>Definition and importance of a research synopsis.</li> <li>Key components of a synopsis: title, abstract, introduction, objectives, methodology, and timeline.</li> <li>Differences between a</li> </ul>
		•	Learn to develop a clear and concise research question.		synopsis, proposal, and full research paper.
		•	Literature Review	Develop	ping a Research Question:
		•	ethodology Master techniques for conducting a literature review. Understand how to design a robust research methodology		<ul> <li>Characteristics of a good research question: clarity, specificity, and feasibility.</li> <li>Techniques for formulating research questions: PICOT framework, FINER criteria.</li> <li>Refining and narrowing down research questions.</li> </ul>

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Writing the Synopsis	Conducting a Literature Review:
and Managing References	<ul> <li>Purpose and scope of a</li> </ul>
<ul> <li>Learn to write each section of a research synopsis.</li> <li>Understand the importance of proper citation and reference management.</li> </ul>	
	Designing the Research Methodology:
<ul> <li>Peer Review and Finalizing the Synopsis</li> <li>Learn the peer-review process and its importance.</li> <li>Finalize and polish the research synopsis.</li> </ul>	<ul> <li>Selection of appropriate study design based on the research</li> </ul>
	Writing the Synopsis:
	<ul> <li>Title and Abstract:</li> <li>Crafting a clear and informative title.</li> <li>Writing a concise abstract that summarizes the research.</li> </ul>
	<ul> <li>Introduction:         <ul> <li>Background and rationale for the study.</li> <li>Stating the research problem and objectives.</li> </ul> </li> </ul>
	<ul> <li>Methodology:         <ul> <li>Detailed description of the research design, data collection, and analysis.</li> <li>Timeline and Budget:</li> </ul> </li> </ul>



	A MEDICAL AS	Rawaipindi Medical University	, <u> </u>
		time resea ■ Estin justif budg o <b>References:</b> ■ Citing accu	nating and ying the research yet.
		Reference Management:	
		(e.g., EndNo Zotero).	inaging references te, Mendeley, ion styles (e.g., 'ancouver).
		Peer Review Process:	
		<ul> <li>research.</li> <li>How to prov feedback.</li> <li>Reviewing a synopses.</li> </ul>	of peer review in ride constructive nd critiquing peer og feedback to synopsis.
		Finalizing the Synopsis:	
		<ul> <li>techniques.</li> <li>Ensuring cla and concises</li> <li>Checking for adherence t</li> </ul>	proofreading rity, coherence, ness in writing. completeness and o guidelines. le final document on.



5.	Card	Introduction to	1.	Introduction to Cardiac Emergencies:
	iac	Cardiac Emergencies and		<ul> <li>Overview of cardiac</li> </ul>
	First	Basic Life Support (BLS)		emergencies: heart attack,
				cardiac arrest, angina, and
	Response	<ul> <li>Understand the types</li> </ul>		arrhythmias.
		and signs of cardiac		<ul> <li>Recognizing symptoms and risk</li> </ul>
		emergencies.		factors.
		Learn the		<ul> <li>The importance of timely</li> </ul>
		fundamentals of Basic		intervention and the concept
		Life Support (BLS).		of the "golden hour."
			2.	Basic Life Support (BLS):
				<ul> <li>Principles of BLS: ensuring</li> </ul>
		Automated External		scene safety, assessing
		Defibrillator (AED) Use and		responsiveness, and calling for
		Advanced Life Support (ALS)		help.
				<ul> <li>Steps of BLS: airway, breathing,</li> </ul>
		Gain proficiency in the		and circulation (ABC).
		use of an Automated		<ul> <li>Hands-on practice: chest</li> </ul>
		External Defibrillator		compressions, rescue breaths,
		(AED).	-	and using a barrier device.
		Understand the basics	3.	Automated External Defibrillator
		of Advanced Life		(AED):
		Support (ALS).		• Function and importance of an
				AED in cardiac emergencies.
		Scenario-Based		<ul> <li>Step-by-step instructions on</li> </ul>
		Training and Team		how to use an AED.
		Dynamics		<ul> <li>Safety precautions and travelactions actions</li> </ul>
				troubleshooting common
		Apply knowledge and		issues.
		skills in realistic,		<ul> <li>Hands-on practice with AED simulators.</li> </ul>
		scenario-based	1	Introduction to Advanced Life
		training.	4.	Support (ALS):
		Understand the     importance of		
		importance of effective team		<ul> <li>Overview of ALS and its components.</li> </ul>
		dynamics during a		<ul> <li>The role of medications and</li> </ul>
		cardiac emergency.		advanced airway management.
		calulae entergency.		<ul> <li>Introduction to ECG</li> </ul>
		Advanced Skills and		interpretation for identifying
		Final Assessment		cardiac rhythms.
				<ul> <li>Coordination and</li> </ul>
		Learn advanced skills		communication in a
		for managing cardiac		resuscitation team.
		emergencies.	5.	Scenario-Based Training:
		chiergeneics.		<ul> <li>Simulated cardiac emergencies</li> </ul>
				with real-time response.

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		•	Demonstrate		0	Role-playing various scenarios:
			competency through a			out-of-hospital cardiac arrest,
			final assessment.			in-hospital cardiac arrest, and
						post-resuscitation care.
					0	Debriefing and feedback
					0	sessions to identify strengths
						and areas for improvement.
				C	Teer	-
				6.		Dynamics in Cardiac
					Emer	gencies:
					0	Importance of teamwork and
						clear communication.
					0	Roles and responsibilities of
						team members during a
						resuscitation effort.
					0	Strategies for effective
					-	leadership and coordination.
					0	Hands-on practice with team
					0	drills and role assignments.
				7.	۸dva	nced Skills:
				7.	Auva	
					0	Advanced airway management:
						intubation and supraglottic
						airway devices.
					0	Intravenous (IV) access and
						medication administration.
					0	Post-resuscitation care:
						monitoring and stabilizing the
						patient.
					0	Review of ACLS algorithms and
						protocols.
				8.	Final	Assessment:
				0.	0 I IIIai	
					0	
						cardiac emergency scenarios to
						assess BLS, AED, and ALS skills.
					0	
						knowledge of cardiac
						emergency management, BLS,
						and ALS protocols.
					0	Feedback and discussion on
						performance.
					0	
						who meet competency
						standards.
L	1					



# **SECTION VI: ROTATIONS (ELECTIVES)**







### **Overview**

The MS Ophthalmology program incorporates rotations in various medical specialties to provide residents with a well-rounded clinical perspective, which is essential in delivering comprehensive patient care. Exposure to fields like emergency medicine, dermatology, radiology, pathology, oncology, neurology, plastic surgery, and community ophthalmology equips residents with multidisciplinary knowledge and skills. This approach enhances diagnostic accuracy, broadens the understanding of systemic conditions affecting the eye, and fosters holistic treatment approaches for patients with complex ocular and systemic comorbidities.



# **Rotation Framework**

### 1st Year

- Department: Emergency Medicine
- **Duration**: 1 Month
- **Importance**: Emergency Medicine rotation enables residents to gain experience in acute care management, including prompt evaluation and treatment of ophthalmic emergencies. This rotation is essential for developing skills in managing trauma, acute visual loss, and other time-sensitive conditions in ophthalmology.

## 3rd & 4th Year

Residents rotate through several subspecialties to integrate a comprehensive understanding of systemic factors influencing ocular health and to strengthen collaborative skills across specialties.



### 1. Dermatology

- **Duration**: 2 Weeks
- **Importance**: Dermatology provides insights into systemic skin conditions, such as rosacea, psoriasis, and lupus, which can manifest with ocular involvement. Knowledge of dermatologic conditions helps in diagnosing and managing periocular and anterior segment diseases with dermatological associations.

## 2. Radiology

- **Duration**: 2 Weeks
- **Importance**: Radiology rotation trains residents to interpret CT scans, MRIs, and ultrasounds relevant to the eye and orbit. Imaging skills are crucial for diagnosing orbital tumors, fractures, optic nerve disorders, and other pathologies requiring radiological evaluation.

## 3. Pathology

- **Duration**: 2 Weeks
- **Importance**: Pathology rotation familiarizes residents with histopathological aspects of ocular conditions. Understanding tissue pathology, especially in tumors, inflammatory diseases, and degenerative conditions, is invaluable for diagnosing and guiding treatment plans in complex cases.

### 4. Oncology

- **Duration**: 2 Weeks
- **Importance**: Oncology introduces residents to ocular and orbital cancers, such as retinoblastoma and melanoma. It emphasizes early detection, staging, and collaborative management, allowing ophthalmology residents to effectively coordinate care with oncologists and deliver informed counseling to patients.

#### 5. Neurology

- **Duration**: 2 Weeks
- **Importance**: Neurology rotation provides essential training in recognizing neuro-ophthalmic conditions like optic neuritis, cranial nerve palsies, and intracranial hypertension. Residents learn to correlate neurological symptoms with ocular findings, enhancing their ability to manage neuro-ophthalmic cases.

### 6. Plastic Surgery

- **Duration**: 2 Weeks
- **Importance**: Plastic surgery rotation enhances skills in handling periocular trauma, reconstructive surgery, and aesthetic procedures. This training is crucial for understanding anatomy and surgical techniques that support safe, effective eyelid and orbital procedures in ophthalmology.

## 7. Community Ophthalmology

- **Duration**: 4 Weeks
- **Importance**: Community Ophthalmology emphasizes preventive and public health aspects of eye care, such as vision screening, outreach programs, and awareness campaigns. This rotation develops skills in managing community-level eye care and understanding the socio-economic factors affecting eye health in diverse populations



# **Rotation planner**

Year	Departments	Duration
1 <sup>st</sup> Year	Emergency Medicine	1 Month
2 <sup>nd</sup> Year	No Rotation	No Rotation
3 <sup>rd</sup> & 4 <sup>th</sup> year	Dermatology	2 weeks
	Radiology	2 weeks
	Pathology	2 weeks
	Oncology	2 weeks
	Neurology	2 weeks
	Plastic Surgery	2 weeks
	Community Ophthalmology	4 weeks

# **Minimum Log Book Entries for Rotations**

(This minimum number is being provided for uniformity of the training and convenience for monitoring of the resident's performance by Quality Assurance Cell & University Research Training & Monitoring Cell of RMU but resident is encouraged to show performance above this minimum required number)



SR.NO	ENTRY	Minimum cases /Time duration
01	Case presentation	02 per rotation
02	Topic presentation	01 per rotation
03	Journal club	01 per rotation
04	Bed side teaching	2 per rotation
05	Large group teaching	01 per rotation
06	Emergency cases	05 per rotation
07	OPD	10 per rotation
08	Indoor (patients allotted)	10 per rotation
09	Directly observed procedures	5 per rotation
10	СРС	01 per rotation
11	Mortality & Morbidity meetings	01 per rotation



# **SECTION VII: RESEARCH**





**Resident research pathway** 

# **4 YEARS UNIVERSITY RESIDENTS RESEARCH PATHWAY**



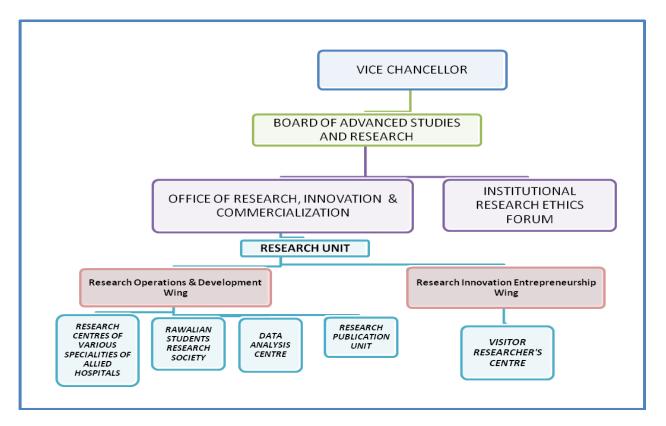
For residency program research work on synopsis and thesis writing starts from entry into university by getting registration ID number from the research unit. It has been structured in 06 monthly & annual time scale goals. Essential steps are included in eligibility criteria of yearly, midterm and final assessments. Compulsory workshops have been designed to train residents along the pathway of research conduction. The charts below show the structure and timeline description of the tasks required.



# **Outline of research curriculum**

Component	Year
Clinical Audit /Disease Statistical Review	Y1
Basic Research Methodology	Y1
Research lectures	Y1
Synopsis Writing	Y2
Referencing Manager	Y2
Research lectures	Y2
Advance Research Methodology	Y3
Data Entry & Analysis SPSS	Y3
Thesis writing workshop	Y4
Writing an Article / Publications	Y4
Research lectures	Y4

## Model of Research at Rawalpindi Medical University





## 2. Year 1

- Milestones:
  - **Research Registration ID** (1st Month): Registration establishes an official record of the resident's research participation and aligns with institutional requirements.
  - **Synopsis Topic Assignment & Submission to Research Unit** (1st 6 Months): Residents are assigned a research topic, ideally aligned with their clinical interests, which lays the foundation for their thesis.
  - **Single Disease Statistical Review or Research Paper** (Before End of Year 1): Residents complete a statistical review or publish a research paper, introducing them to data interpretation and critical analysis.
- Training Components:
  - **Basic Research Methodology**: A series of introductory lectures on research methodology, covering study designs, sample size calculation, and research ethics.
  - **Research Lectures**: Regular sessions to familiarize residents with foundational research principles.
  - **Synopsis Writing & Referencing Manager Training**: Workshops on writing a research synopsis and managing references using software like EndNote or Zotero.

## 3. Year 2

- Milestones:
  - **Submission of Synopsis** (1st 6 Months): Residents submit a detailed research proposal outlining their study objectives, methodology, and expected outcomes.
  - **Technical Committee Evaluation, IRF/ERB Approval, and BASAR Synopsis Approval**: These steps involve institutional and ethical approvals, ensuring the research project meets ethical standards and feasibility.
- Training Components:
  - Advanced Research Methodology: Building on the basics, this module covers complex statistical tests, bias minimization, and confounding variables.
  - **Research Lectures**: Continued educational sessions to reinforce methodological rigor and address any challenges in the research process.

## 4. Year 3

- Milestones:
  - **1 Disease Statistical Review or 1 Research Paper** (Optional): An optional review or paper to further enhance research skills.
  - **Data Collection** (1st 6 Months): Residents begin gathering data for their thesis under guidance, focusing on data quality and integrity.
  - **Data Analysis** (Last 6 Months): Residents learn to apply statistical techniques using software like SPSS, analyzing the data collected for meaningful insights.
- Training Components:
  - **Data Entry & Analysis with SPSS**: A workshop on using SPSS for data entry and performing essential statistical analyses, equipping residents with hands-on analytical skills.



- **Research Lectures**: Ongoing lectures addressing specific challenges in data management and analysis.
- 5. Year 4
  - Milestones:
    - **Thesis Writing** (1st 6 Months): Residents draft their thesis with structured guidance on writing style, formatting, and scientific rigor.
    - **BASAR Thesis Approval** (Last 6 Months): The thesis is submitted for approval, marking the completion of the primary research requirement.
    - **Thesis Completion Certificate by DME** (Last 6 Months): Upon approval, the Department of Medical Education certifies the thesis completion, fulfilling the academic research requirement.
  - Training Components:
    - **Thesis Writing Workshop**: A dedicated session on organizing research findings, structuring a thesis, and using appropriate academic language.
    - Writing an Article/Publications: Guidance on publishing research findings, including manuscript preparation, journal selection, and the peer-review process.
    - **Research Lectures**: Concluding lectures covering advanced topics in publication ethics and responding to reviewer feedback.

# **Research milestones**

MILESTONE	TIMELINE	
Research registration id	1 <sup>st</sup> Month	¥1
Synopsis topic assignment and submission to Research Unit	1 <sup>st</sup> 05 Month	Y1
Single disease statistical review or 1 paper in RJRMC	Before end of year 1	Y1
Submission of synopsis	1 <sup>st</sup> 05 Month	¥2
Technical committee evaluation	1 <sup>st</sup> 05 Month	¥2
IRF/ERB synopsis approval	1 <sup>st</sup> 05 Month	Y2
Basar synopsis approval	last 06 Month	¥2
1 disease statistical review or 1 research paper in RJRMC	optional	Y3
Data collection	1 <sup>st</sup> 06 Month	¥3
Data analysis	Last 05 Month	¥3
Thesis writing	1 <sup>st</sup> 05 Month	¥4
BASAR thesis approval	Last 05 Month	¥4
Thesis completion certif icate by DME	Last 06 Month	¥4



### **Research Work Assessment**

#### Submission of Synopsis and Thesis

The candidates shall prepare their synopsis as per guidelines provided by the Advanced Studies & Research Board, available on RMU website.

Synopsis of research project should be submitted and approved by the end of the 1<sup>st</sup> year of MS program.

The minimum duration between approval of synopsis and submission of thesis shall be one year, but the thesis cannot be submitted later than 8 years of enrolment.

Thesis shall be submitted by the candidate duly recommended by the Supervisor.

The research thesis must be compiled and bound in accordance with the Thesis Format Guidelines approved by the University and available on website.

The research thesis will be submitted along with the fee prescribed by the University.

#### **Thesis Assessment**

All candidates admitted in MS course shall appear in thesis evaluation component of the MTA after completion of 4<sup>th</sup> years of their training course.

Only those candidates shall be eligible for thesis evaluation who have passed Midterm Examination and Oral & Practical/ Clinical component of Exit Examination.

The examination shall include thesis evaluation with defense.

The Vice Chancellor shall appoint three external examiners for thesis evaluation, preferably from other universities and from abroad, out of the panel of examiners approved by the Advanced Studies & Research Board. The examiners shall be appointed from respective specialty.

The thesis shall be sent to the external examiners for evaluation, well in time before the date of defense examination and should be approved by all the examiners.

After the approval of thesis by the evaluators, the thesis defense examination shall be held within the University on such date as may be notified by the Controller of Examinations. The Controller of Examinations shall make appropriate arrangements for the conduct of thesis defense examination in consultation with the supervisor, who will co-ordinate the defense examination.

The thesis defense examination shall be conducted by two External Examiners who shall submit a report on the suitability of the candidate for the award of degree. The supervisor shall act as coordinator.

Candidates and faculty interested in further details relating to research, please refer to the document on Research curriculum (also available on RMU website)



# **SECTION VIII: TRAINING MILESTONES**







# **Charting the Road to Competence: Developmental Milestones**

Remember to celebrate for the milestones as you prepare for the road ahead----Nelson Mandela.

High-quality assessment of resident performance is needed to guide individual residents' development and ensure theirpreparedness to provide patient care. To facilitate this aim, reporting milestones are now required across all Ophthalmology residency programs. Milestones promote competency-based training in internal medicine. Residency program directors may use them to track the progress of trainees in the 6 general competencies including *patient care, Medical Knowledge, Practice-Based Learning and Improvement, Interpersonal and Communication Skills, Professionalism and Systems-Based Practice.* Mile stones inform decisions regarding promotion and readiness for independent practice. In addition, the milestones may guide curriculum development, suggest specific assessment strategies, provide benchmarks for resident self-directed assessment-seeking, assist remediation by facilitating identification of specific deficits, and provide a degree of national standardization in evaluation. Finally, by explicitly enumerating the profession's expectations for graduates, they may improve public accountability for residency training.

Table-1	Developmental Milestones for Ophthalmology Training— Patient Care		
Competency	Developmental MilestonesInforming Competencies	Approximate Time FrameTrainee Should Achieve Stage (months)	General Evaluation StrategiesAssessment Methods/ Tools
A. Clinical skills and reasoning	ļ	Historical data gathering	
<ul> <li>Manage patients using clinical skills of interviewing and physical examination</li> <li>Demonstrate</li> </ul>	<ol> <li>Acquire accurate and relevant history from the patient in an efficiently customized, prioritized, and hypothesis driven fashion</li> <li>Seek and obtain</li> </ol>	8	<ul> <li>Standardized patient</li> <li>Direct observation</li> </ul>
<ul> <li>Demonstrate competence in the performance of procedures</li> <li>Appropriately use</li> </ul>	appropriate, verified, and prioritized data from secondary sources (eg, family, records, pharmacy)		
laboratory and imaging techniques	3. Obtain relevant historical subtleties that inform and prioritize both differential diagnoses and diagnostic plans, including sensitive, complicated, and detailed information that may not often be volunteered by the patient	24	
	4. Role model gathering subtle and reliable information from the patient for junior members of the health care team	40	
	1. Perform an accurate physical examination that is appropriately targeted to the patient's complaints and medical conditions. Identify pertinent abnormalities using common maneuvers	Performing a physica 8	<ul> <li>Standardized patient direct observation</li> <li>Simulation</li> </ul>



2. Accurately track important changes in the physical examination over time in the outpatient and inpatient settings123. Demonstrate and teach how to elicit24	
important physical findings for junior members of the health care team	
4. Routinely identify40subtle orunusual physicalfindings that mayinfluence clinicaldecision making, usingadvanced maneuverswhere applicable	
Clinical reasoning	
available data, including	Chart-stimulated recall Direct
examination, and	observation Clinical vignettes
2. Develop prioritized 32 differential diagnoses, evidence- based diagnostic and therapeutic plan for common inpatient and ambulatory conditions	
3. Modify differential 32 diagnosis and care plan based on clinical course and data as appropriate	
4.Recognize disease 48 presentations that deviate from common patterns and that require complex decision making	
Invasive procedures	
	Simulation Direct



		MEDICAL ST	Rawalpindi Medical University
	procedures and		observation
	provide post-procedure		
	management for		
	common procedures		
B. Delivery of			
patient- <sup>*</sup> centered	1. Make appropriate		<ul> <li>Chart-stimulated</li> </ul>
clinical care	clinical decisions based		recall
<ul> <li>Manage</li> </ul>	on the results of		Standardized
patients	common diagnostic		tests
with	testing, including but	16	
progressive	not limited to routine	10	Clinical vignettes
responsibility	blood chemistries,		
. ,	hematologic studies,		
<ul> <li>Manage</li> </ul>	coagulation tests, ECG,		
patients	chest radiographs,		
across the	Auto-refraction, Cycloplegic refraction,		
spectrum of	FFA, A-scan, B-scan,		
clinical	Intra-ocular pressure,		
diseases seen	Keratometry, Perimetry		
in the	and Gonioscopy		
practice of	2. Make appropriate		
general	clinical decision based	24	
internal	on the results of more	24	
medicine	advanced diagnostic		
<ul> <li>Manage</li> </ul>	tests		
patients in a	Pat		
variety of			
health care	1. Recognize situations		Simulation
settings to	with a need for urgent	8	Chart-stimulated
include the	or emergent medical	0	recall
	care and/or surgical		
inpatient	care.		Multisource
ward, critical			feedback
care units,	2. Recognize when to	8	<ul> <li>Direct</li> </ul>
the	seek additional		observation
ambulatory	guidance		Chart audit
setting, and	3. Provide appropriate		
the	preventive care and	`	
emergency	•		
	teach patient regarding		
setting	teach patient regarding self-care		
setting <ul> <li>Manage</li> </ul>	teach patient regarding self-care 4. With supervision,		
-	teach patient regarding self-care 4. With supervision, manage patients with		
<ul> <li>Manage undifferentiat</li> </ul>	teach patient regarding self-care 4. With supervision, manage patients with common clinical	16	
<ul> <li>Manage undifferentiat ed acutely</li> </ul>	teach patient regarding self-care 4. With supervision, manage patients with common clinical disorders seen in the	16	
<ul> <li>Manage undifferentiat ed acutely and severely</li> </ul>	teach patient regarding self-care 4. With supervision, manage patients with common clinical disorders seen in the practice of inpatient	16	
<ul> <li>Manage undifferentiat ed acutely and severely ill patients</li> </ul>	teach patient regarding self-care 4. With supervision, manage patients with common clinical disorders seen in the	16	
<ul> <li>Manage undifferentiat ed acutely and severely ill patients</li> <li>Manage</li> </ul>	teach patient regarding self-care 4. With supervision, manage patients with common clinical disorders seen in the practice of inpatient department.	16	
<ul> <li>Manage undifferentiat ed acutely and severely ill patients</li> <li>Manage patients in</li> </ul>	teach patient regarding self-care 4. With supervision, manage patients with common clinical disorders seen in the practice of inpatient department. 5. With minimal	16	
<ul> <li>Manage undifferentiat ed acutely and severely ill patients</li> <li>Manage patients in the</li> </ul>	teach patient regarding self-care 4. With supervision, manage patients with common clinical disorders seen in the practice of inpatient department.		
<ul> <li>Manage undifferentiat ed acutely and severely ill patients</li> <li>Manage patients in the prevention,</li> </ul>	teach patient regarding self-care 4. With supervision, manage patients with common clinical disorders seen in the practice of inpatient department. 5. With minimal supervision, manage	16	
<ul> <li>Manage undifferentiat ed acutely and severely ill patients</li> <li>Manage patients in the</li> </ul>	<ul> <li>teach patient regarding self-care</li> <li>4. With supervision, manage patients with common clinical disorders seen in the practice of inpatient department.</li> <li>5. With minimal supervision, manage patients with common</li> </ul>		



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diagnosis, and treatment of gender- specific diseases			
<ul> <li>Image</li> <li>Patients as</li> <li>a</li> <li>onsultant</li> </ul>	6. Initiate management and stabilize patients with emergent conditions	16	
to other physician	7. Manage patients with conditions that require intensive care	48	
	<ol> <li>8. Independently manage patients with a broad spectrum of clinical disorders seen in the practice of ophthalmology.</li> </ol>	48	
	9. Manage complex or rare ophthalmological conditions	48	
	10. Customize care in the context of the patient's preferences and overall health	48	
		Consultative care	
	1. Provide specific, responsive consultation to other services	32	<ul> <li>Simulation</li> <li>Chart-stimulated recall</li> <li>Multiseurse</li> </ul>
	2. Provide ophthalmological consultation for patients with more complex clinical problems requiring detailed risk assessment	48	<ul> <li>Multisource feedback</li> <li>Direct observation</li> <li>Chart audit</li> </ul>

Table-2 D		nes for Ophthalmology Iowledge	Training— Medical
ompetency	Developmental MilestonesInforming Competencies	Approximate Time FrameTrainee Should Achieve Stage (months)	General Evaluation StrategiesAssessment Methods/ Tools
A. Core knowledge		Knowledge of co	re content
of ophthalmology • Demonstrate a level of expertise in	1. Understand the relevant pathophysiology and basic science for common conditions	8	<ul> <li>Direct observation</li> <li>Chart audit</li> <li>Chart-stimulated recall</li> </ul>
the knowledge of those areas appropriate for an internal	2. Demonstrate sufficient knowledge to diagnose and treat common conditions that require hospitalization	16	<ul> <li>Standardized tests</li> </ul>
medicine specialist • Demonstrate	3. Demonstrate sufficient knowledge to evaluate common conditions	24	
sufficient knowledge to treat ophthalmic conditions	4. Demonstrate sufficient knowledge to diagnose and treat undifferentiated and emergent conditions	24	
commonly managed by internists,	5. Demonstrate sufficient knowledge to provide preventive care	24	
provide basic preventive care, and recognize and provide initial	6. Demonstrate sufficient knowledge to identify and treat conditions that require intensive care	32	
management of emergency problems	7. Demonstrate sufficient knowledge to evaluate complex or rare conditions and multiple coexistent conditions	48	
	8. Understand the relevant pathophysiology and basic science for uncommon or complex conditions	48	
	9. Demonstrate sufficient knowledge of sociobehavioral sciences including but not limited to health care	48	



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B. Common modalities used in the practice of ophthalmolog y & Demonstrate sufficient knowledge to interpret basic clinical tests and	economics, medical ethics, and medical education Diagnostic tests 1. Understand indications for and basic interpretation of common diagnostic testing, including but not limited to routine blood chemistries, hematologic studies, coagulation tests, ECG, chest	16	<ul> <li>Chart-stimulated recall</li> <li>Standardized tests</li> <li>Clinical vignettes</li> </ul>
tests and images, use common pharmacothe rapy, and appropriately use and perform diagnostic and therapeutic procedures.			
	2. Understand indications for and has basic skills in interpreting more advanced diagnostic tests	24	
	<ol> <li>Understand prior probability and test performance characteristics</li> </ol>	24	



Table-3         Developmental Milestones for Ophthalmology Training—				
Practice-Based Learning and Improvement				
Competen	Developmental	Approximate	General Evaluation	
су	Milestones	Time Frame	StrategiesAssessment	
1	Informing	Trainee	Methods/ Tools	
	Competencies	Should		
	competencies	Achieve		
A Loorning		Stage (months)		
A. Learning and		e the quality of co	ire for a panel of	
improving via audit	patient	S		
of	1. Appreciate the	10	<ul> <li>Several</li> </ul>	
performa	responsibility to	16	elements of	
nce & Systemat	assess and improve care collectively for		quality	
ically	a panel of patients		improvement	
analyze practice	2. Perform or		project	
using	review audit of a	32	<ul> <li>Standardized</li> </ul>	
quality improve	panel of patients	52	tests	
ment	using standardized,			
methods, and	disease-specific,			
implemen	and evidence-			
t changes with the	based criteria 3. Reflect on audit			
goal of	compared with			
practice improve	local or national	22		
ment	benchmarks and	32		
	explore possible			
	explanations for			
	deficiencies,			
	including doctor-			
	related, system-			
	related, and patient related factors			
	4. Identify areas in	48		
	resident's own	40		
	practice and local			
	system that can be			
	changed to improve			
	effect of			
	the processes and			
	outcomes of care 5. Engage in a	48		
	quality	40		
	improvement			
	intervention			
B. Learning	<ul> <li>Ask answerabl</li> </ul>	e questions for eme	erging information needs	
and	1. Identify learning		<ul> <li>Evidence-based</li> </ul>	
improvemen t via	needs (clinical	16	medicine	
t via answering	questions) as they		evaluation	
8.10110110	emerge in patient			



			THEDICAL ST	Rawaipindi Medical Univers
	clinical questions	care activities		, instruments
	from patient			EBM mini-CEX
	scenarios	2. Classify and	32	Chart-stimulated
•	Locate,	precisely	52	recall
	appraise, and	articulate clinical		
	assimilate	questions		
	evidence	3. Develop a system	32	
	from	to track, pursue,		
	scientific	and reflect on clinical questions		
	studies	cliffical questions	Acquiras the he	stovidonco
	related to	4. A	Acquires the be	
	their	1. Access medical information		Evidence-based
	patients'	resources to answer	16	medicine
	health	clinical questions		<ul> <li>evaluation</li> </ul>
	problems;	and support decision		instruments
•	Use	making		• EBM, mini-CEX,
	information	<ol><li>Effectively and efficiently search</li></ol>	16	Chart-stimulated
	technology to	NLM database for	10	recall
	optimize	original		
	learning	clinical research		
		articles		
		3. Effectively and	22	
		efficiently search evidence- based	32	
		summary medical		
		information		
		resources		
		4. Appraise the quality of medical		
		information	48	
		resources and select		
		among them based		
		on the		
		characteristics of the clinical question		
			for validity and usefulne	SS
		1. With assistance,		
		appraise study		Evidence-based
		design, conduct, and	16	medicine
		statistical analysis in		<ul> <li>evaluation</li> </ul>
		clinical		instruments
		research papers		EBM mini-CEX
		2. With assistance, appraise clinical	32	<ul> <li>Chart-stimulated</li> </ul>
		guidelines		recall
		3. Independently		
		appraise study	48	
		design, conduct, and		
		statistical		
		analysis in clinical research papers		
		4. Independently,		
		appraise clinical		
		guideline	48	
		recommendations		



	for bias and cost- benefit considerations		
		evidence to decisio	n-making for individual
	<b>patients</b> 1. Determine if clinical evidence can be generalized to an individual patient	16	<ul> <li>Evidence-based medicine</li> <li>evaluation instruments EBM mini-CEX</li> </ul>
	2. Customize clinical evidence for an individual patient	32	<ul> <li>Chart-stimulated recall</li> </ul>
	<ol> <li>Communicate risks and benefits of alternatives to patients</li> </ol>	48	
	4. Integrate clinical evidence, clinical context, and patient	48	
	preferences into decision making		
C. Learnin g and		Improves vi	ia feedback
<ul> <li>improv ing via feedbac k and self- assess ment</li> <li>Identify strengths, deficiencies, and limits in one's</li> </ul>	1. Respond welcomingly and productively to feedback from all members of the health care team including faculty, peer residents, students, nurses, allied health workers, patients, and their	16	<ul> <li>Multisource feedback</li> <li>Self-evaluation forms with action plans</li> </ul>
knowledge	advocates 2. Actively seek	24	
<ul><li>and expertise</li><li>Set learning and</li></ul>	feedback from all members of the health care team	24	
improvement goals • Identify and	<ol> <li>Calibrate self- assessment with feedback and other external data</li> </ol>	32	
perform appropriate learning activities	4. Reflect on feedback in developing plans for improvement	32	
<ul> <li>Incorporate</li> </ul>		-	n self-assessment
formative evaluation feedback into	1. Maintain awareness of the situation in the moment, and	32	<ul> <li>Multisource feedback</li> <li>Reflective</li> </ul>



		A MEDICAL M	Rawaipinui medicai Univer
<ul><li>daily practice</li><li>Participate in</li></ul>	respond to meet situational needs		practice surveys
the education of patients, families, students, residents, and other health professionals	2. Reflect (in action) when surprised, applies new insights to future clinical scenarios, and reflects (on action) back on the process	48	
	Participates in	the education of a	all members of the
	, health care tea		,
	1. Actively participate in teaching conferences	16	<ul> <li>OSCE with standardized</li> </ul>
	2. Integrate teaching, feedback, and evaluation with supervision of interns' and students' patient care	32	<ul><li>learners Direct</li><li>observation</li><li>Peer evaluations</li></ul>
	3. Take a leadership role in the education of all members of the health care team.	48	

Table-4 Developmental Milestones for Ophthalmology Training— Interpersonal and Communication Skills			
Competen cy	Developmental Milestones Informing Competencies	Approxima te Time Frame Trainee Should Achieve Stage (months)	General Evaluation StrategiesAssessment Methods/ Tools
<u>A. Communicate</u>		Communicate e	ffectively
<ul> <li><i>effectively:</i></li> <li>Patients and family</li> </ul>	1. Provide timely and comprehensive verbal and written communication to patients/advocates	16	<ul> <li>Multisource feedback</li> <li>Patient surveys</li> <li>Direct</li> </ul>
Communicate effectively with patients, families, and	2. Effectively use verbal and nonverbal skills to create rapport with patients/families	16	observation <ul> <li>Mentored self- reflection</li> </ul>
the public, as appropriate, across a	3. Use communication skills to build a therapeutic relationship		
broad range of socioeconomi	<ol> <li>Engage patients /advocates in shared decision making for uncomplicated</li> </ol>	32	



		TO MEDICAL ST	Rawalpindi Medical Univers
c and cultural backgrounds	diagnostic and therapeutic scenarios		
	5. Use patient-centered education strategies	32	
	<ol> <li>Engage patients /advocates in shared decision making for difficult, ambiguous, or controversial scenarios</li> </ol>	48	
	7. Appropriately counsel patients about the risks and benefits of tests and procedures, highlighting cost awareness and resource allocation	48	
	8. Role model effective communication skills in challenging situations	48	
		Intercultural sens	sitivity
	1. Effectively use an interpreter to engage patients in the clinical setting, including patient education	8	<ul> <li>Multisource feedback</li> <li>Direct observation</li> </ul>
	2. Demonstrate sensitivity to differences in patients including but not limited to race, culture, gender, sexual orientation, socioeconomic status, literacy, and religious beliefs	16	• Mentored self- reflection
	3. Actively seek to understand patient differences and views and reflects this in respectful communication and shared decision-making with the patient and the healthcare team	40	
B. Physicians and		<b>Transitions</b> of	of care
other health care professionals	1. Effectively communicate with other caregivers in order to maintain	16	<ul> <li>Multisource feedback</li> <li>Direct</li> </ul>
<ul> <li>Communicat e effectively with</li> </ul>	appropriate continuity during transitions of care		observation <ul> <li>Sign-out</li> </ul>



	r	MEDICAL	Rawaipinui medicai onive
physicians,	2. Role model and	32	form ratings
other health	teach effective		Patient
professional	communication with		surveys
s, and	next caregivers during		301 7 6 9 3
health-	transitions of care		
related		Interprofessional team	
agencies Work			
effectively as a member			
or leader of			
a health care			
team or			
other			
professional			
	1 Deliver	0	
group	1. Deliver	8	Multisource
Act in a	appropriate,		feedback
consultat	succinct,		
ive role	hypothesis-driven		
to other	oral presentations		
physician	2. Effectively	16	
s and	communicat		
health	e plan of		
professio	care to all		
nals	members of		
	the health care team		
	3. Engage in	40	
	collaborative		
	communication		
	with all members		
	of the health care		
	team		
		Consul	
	1. Request	8	<ul> <li>Multisource</li> </ul>
	consultative		feedback
	services in		
	an effective		<ul> <li>Chart audit</li> </ul>
	manner		
	2. Clearly	16	
	communicate the		
	role of consultant		
	to the patient, in		
	support of the		
	primary care		
	relationship	40	
	3. Communicate	48	
	consultative		
	recommendations to		
	the referring team in		
	an effective manner		
C Madiaul vacanda		llealth re	
<u>C. Medical records</u>		Health re	
Maintain	1. Provide legible,	8	<ul> <li>Chart audit</li> </ul>
comprehensive	accurate,		
, timely, and	complete, and		
	timely written		
legible medical	communication		
records	communication that is congruent		
	communication		



2. Ensure succinct, relevant, and patient-specific written communication	32	

Table-5 Develo Professionalism	pmental Milestones	s for Ophthalmol	ogy Training—
Competen cy	Developmental Milestones Informing Competencies	Approxima te Time Frame Trainee Should Achieve	General Evaluation StrategiesAssessment Methods/ Tools
		Stage (months)	
A. <u>Physicianship</u> Demonstrate compassion, integrity, and respect for others responsiveness to	1. Document and report clinical information truthfully	Adhere to bas	ic ethical principles • Multisource feedback
patient needs that supersedes self-	2. Follow formal policies	1.5	
interest Accountability to patients, society, and the profession	3. Accept personal errors and honestly acknowledge them	8	
	4. Uphold ethical expectations of research and scholarly activity	48	
	Demo patie		ion and respect to
	1. Demonstrate empathy and compassion to all patients	4	<ul> <li>Multisource feedback</li> </ul>
	2. Demonstrate a commitment to relieve pain and suffering	4	
	3. Provide support (physical, psychological, social, and spiritual) for dying patients and their	32	



		MEDICAL ST	Rawaipinui Medicai Onive
	families		•
	4. Provide leadership for a team that respects patient dignity and autonomy	32	
-		Provide timely constructive feedback to colleagues	<b>,</b>
	1. Communicate constructive feedback to other members of the health care team	16	<ul> <li>Multisource feedback</li> <li>Mentored self- reflection</li> </ul>
	2. Recognize, respond to, and report impairment in colleagues or substandard care via peer review process	24	Direct     observation
	1. Respond promptly and appropriately to clinical responsibilities including but not limited to calls and pages	Maintain ac	Multisource feedback
	2. Carry out timely interactions with colleagues, patients, and their designated caregivers	8 Recognize conflicts of	finterest
	1. Recognize and manage obvious conflicts of interest, such as caring for family members and professional associates as patients	8	<ul> <li>Multisource feedback</li> <li>Mentored self- reflection</li> <li>Clinical vignettes</li> </ul>
	2. Maintain ethical relationships	40	



	A MEDICAL A	Rawaipindi Medical Unive
with industry		
3. Recognize and manage subtler conflicts of interest	40	
1. Dress and behave appropriately	1.5	<ul> <li>Multisource feedback</li> <li>Discussion</li> </ul>
2. Maintain appropriate professional relationships with patients,	1.5	Direct observation
families, and		
staff 3. Ensure prompt completion of clinical, administrative, and curricular	8	
tasks 4. Recognize and address personal, psychological, and physical limitations that may affect professional performance	16	
5. Recognize the scope of his/her abilities and ask for supervision and assistance appropriately	16	
6. Serve as a professional role model for more junior colleagues (eg, medical students, interns)	40	
7. Recognize the need to assist colleagues in the provision of duties	40	
	Practice indi	vidual patient advocacy
1. Recognize when it is	8	Multisource feedback



		TH MEDICAL ST	Rawalpindi Medical Univer
	advocate for individual patient needs		Direct observation
	2. Effectively advocate for individual patient needs	40	
		Comply with	public health policies
	1. Recognize and take responsibility for situations where public health supersedes individual health (eg, reportable infectious diseases)	32	<ul> <li>Multisource feedback</li> </ul>
B. <u>Patient-</u>	Respect t	he dignity, culture,	beliefs, values, and
<u>centeredness</u>	opinions o	of the patient	
<ul> <li>Respect for patient privacy and autonomy Sensitivity and</li> </ul>	1. Treat patients with dignity, civility and respect, regardlessof race, culture, gender, ethnicity, age, or socioeconomic status	1.5	<ul> <li>Multisource feedback</li> <li>Direct observation</li> </ul>
responsiveness to a diverse patient population, including but	2. Recognize and manage conflict when patient values	40	
not limited to diversity in	differ from their own		
gender,		onfidentiality	
age, culture, race, religion, disabilities, andsexual orientation	1. Maintai n patient confid entiali ty	1.5	<ul> <li>Multisource feedback</li> <li>Chart audits</li> </ul>
	2. Educate and hold others accountable for patient confidential ity	24	
	·		Recognize and address
	1. Recognize that disparitiesexist in health care among populations and that they may impact care of the patient	16	<ul> <li>disparities in health care</li> <li>Multisource feedback</li> <li>Direct observation</li> <li>Mentored self- reflection</li> </ul>



	MEDICAL ST	
2. Embrace physicians' role in assisting the public and policy makers in understandin g and addressing causes of disparity in disease and suffering	40	
3. Advocates for appropriate allocation of limited health care resources.	40	

# Table-6 Developmental Milestones for Ophthalmology Training— Systems **Based Practice**

Competency	Developmental MilestonesInforming Competencies	Approximat e Time Frame Trainee Should Achieve Stage (months)	Metho	ation egies sment ods/ Tools
A. <u>Work effectively with</u> other care providers		ectively within m	ultiple	health
<u>andsettings</u>	delivery sy	<i>stems</i>		
<ul> <li>Work effectively in various health care delivery settings and systems relevant to their clinical practice</li> <li>Coordinate patientcare within the health care system</li> </ul>	<ol> <li>Understand unique roles and services provided by local health care delivery systems.</li> <li>Manage and coordinate care and care transitions across multiple delivery systems, including ambulatory, subacute, acute, rehabilitation, and skilled nursing.</li> </ol>	16 32	5	Multisourc e feedback Chart- stimulated recall Direct observation
system relevant to their clinical specialty 2 Work in interprofession		48 ffectively within fessional team	an	



		ST AT MEDICAL ST ST	Rawalpindi Medical Univers
al teams to enhance patient safety and improve patient care quality Work in teams and effectively transmit necessary clinical information to ensure safe and proper care of patients, including the transition of care between settings	<ol> <li>Appreciate roles of a variety of health care providers, including but not limited to consultants, therapists, nurses, home care workers, pharmacists, and social workers.</li> <li>Work effectively as a member within the interprofession al team to ensure safe patient care.</li> <li>Consider alternative solutions provided by other teammates</li> <li>Demonstrate how to manage the team by using the skills and coordinating the activities of interprofessional team</li> </ol>	8 8 16 48	<ul> <li>Multisourc e feedback</li> <li>Chart- stimulated recall</li> <li>Direct observation</li> </ul>
De las ser las has lik	members.		
B. <u>Improving health</u> <u>caredelivery</u>			nd advocates for
<ul> <li>Advocate for quality patient careand optimal patient care systems</li> <li>Participate in</li> </ul>	system improv 1. Recognize health system forcesthat increase the risk for error including barriers to optimal patient care	16	<ul> <li>Multisourc e feedback</li> <li>Quality improvemen t project</li> </ul>
identifying systemerrors and implementin g potential systems solutions	2. Identify, reflect on, and learnfrom critical incidents such as near misses and preventable medical errors	16	
<ul> <li>Recognize and function effectively</li> </ul>	<ol> <li>Dialogue with care team members to identify risk for and prevention of medical error</li> </ol>	32	
in high- quality care system	4. Understand mechanisms for analysis and correction of systems errors	32	
	5. Demonstrate ability to understand and	48	



		MEDICAL	TLAWC		leuical onive
	quality improvement intervention.		•		
	6. Partner with other health care professionals to identify, propose improvement opportunities within the system.	48			
C. <u>Cost-effective care for</u> patients and	Identifies forces that impa	ct the cost of	health	care and	l advocates
populations	for cost-effective care				
& Incorporate	1. Reflect awareness of common	16		?	Standardize
considerations of cost	socioeconomic barriers				d
awareness and risk-	that impact				examination
benefitanalysis in	patient care. 2. Understand how	16		[?]	s Direct
patient and/or	cost-benefit analysis	10		Ľ	observation
population- based	is applied to patient			?	Chart-
care asappropriate	care(ie, via principles of screening tests and			Ŀ	stimulated
	the development of				
	clinical guidelines)				recall
	<ol> <li>Identify the role of various health care</li> </ol>	32			
	stakeholders				
	including providers,				
	suppliers, financiers, purchasers, and				
	consumers and their				
	varied impact on the cost of and access				
	to health care.				
	4. Understand coding and	32			
	reimbursement principles.				
		Practices co	st-effe	ctive car	е
	1. Identify costs for	8		•	Chart-
	common diagnostic or therapeutic				stimulated
	tests.				recall
	2. Minimize	8			
	unnecessary care including tests,				
	procedures,				
	therapies, and				
	ambulatory or				
	3. Demonstrate the	24			
	incorporation of cost-	24			
	awareness principles				
	into standard clinical judgments and				
	Jaabinents and				
	4. Demonstrate the	48			
	incorporationof cost-				
	awareness principles				
	into				



# **SECTION IX: ASSESSMENT AND EVALUATION**



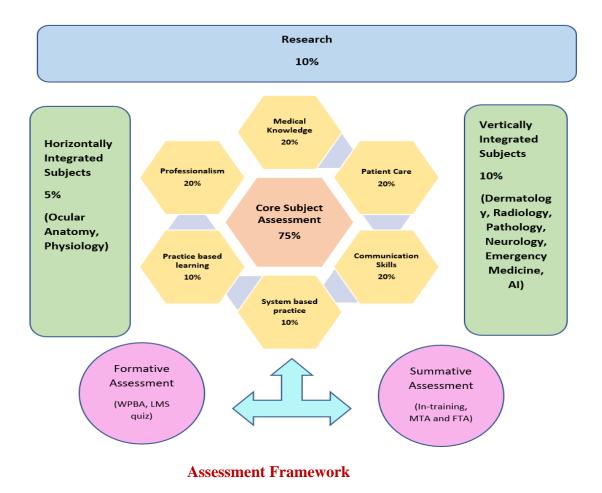




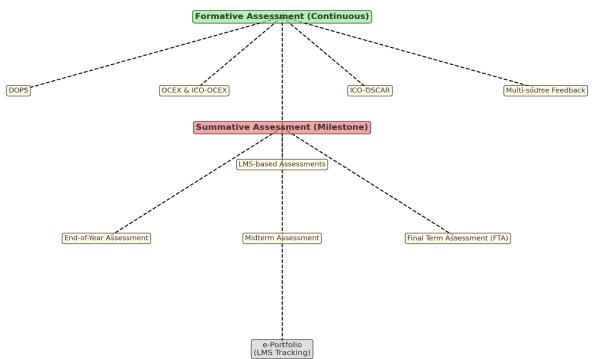
Assessment is essential to ensure that residents in the MS Ophthalmology program are progressing effectively through key competencies required for clinical and academic excellence. It enables both trainees and faculty to identify strengths and areas for improvement, fostering a structured approach to developing diagnostic, procedural, and patient management skills. Through regular formative assessments, feedback, and summative evaluations, residents gain a clear understanding of their performance. This approach not only promotes lifelong learning and professional development but also guarantees that residents meet the high standards expected in patient care, research, and medical ethics.

## Framework of Assessment for MS Ophthalmology Residency Program

The MS Ophthalmology Residency Program at Rawalpindi Medical University uses a structured assessment framework to ensure residents' development in medical knowledge, procedural skills, professionalism, and patient care. Formative assessments, including WPBA and LMS quizzes, provide continuous feedback recorded in each resident's e-log book and portfolio. Summative assessments at key milestones—end-of-year, midterm, and final exams—evaluate the resident's comprehensive knowledge and skills. This approach promotes a well-rounded, independent ophthalmologist committed to high standards of patient care and lifelong learning.







## **Assessment Hours Allocation**

Assessment hours based on the given 8:1 ratio of teaching hours to assessment hours:

**Total Teaching Hours** for the program = 2,112 hours. The formula for assessment hours is:

Assessment Hours=2,112/8=264 hours

Therefore, the MS Ophthalmology training program would include 264 assessment hours over the four years.

#### **Formative Assessments**

Formative assessments are used continuously to monitor progress and provide feedback. These include Workplace-Based Assessments (WPBAs), LMS-Based Assessments, log book and portfolio entries, and other regularly scheduled assessments.



Formative Assessment Hours Allocation:

Type of Formative Assessment	Frequency	Time per Assessment	Hours nor	Total Hours over 4 Years
Assessments (e.g. ULEX	Biannual (2 per year)	1 hour each	2 hours	8 hours
LMS-Based Assessments (Quizzes)	Twice a month	30 minutes each	12 hours	48 hours
Log Book and Portfolio Review	Quarterly	1 hour each	4 hours	16 hours
Case based discussion		30 minutes each	12 hours	48 hours

*Total Formative Assessment Hours*: **8** + **48** + **16** + **48** = **120 hours** 

#### Summative Assessments

Summative assessments are used to evaluate knowledge, skills, and competency at specific stages. These include in-training assessments, and the midterm and final term assessments

#### **Summative Assessment Hours Allocation:**

Type of Summative Assessment	Frequency	Time per Assessment	Total Hours per Year	Total Hours over 4 Years
In-Training Assessment (Year 1)	Once at year- end	5 hours	5 hours	5 hours
In-Training Assessment (Year 3)	Once at year- end	5 hours	5 hours	5 hours
Midterm Assessment	Annually	8 hours each	8 hours	8 hours
Final Term Assessment	Annually	10 hours	10 hours	10 hours

*Total Summative Assessment Hours:* 4 + 4 + 8 + 10 = 28 hours



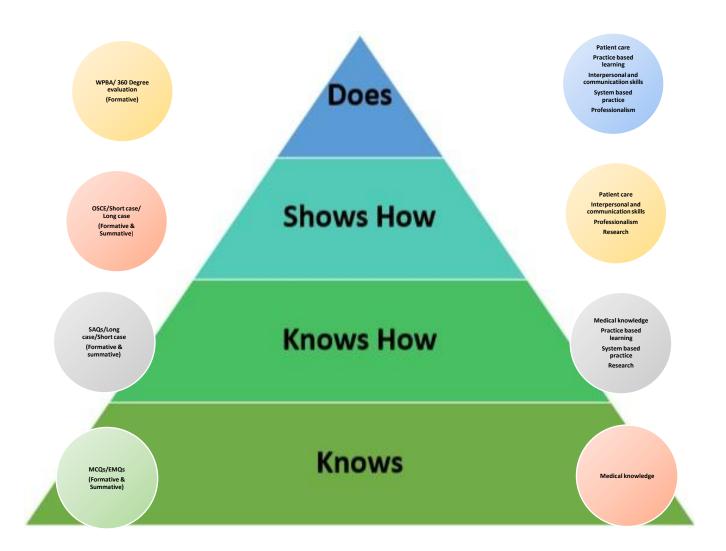
Assessment Type	Frequency (suggested)	Time per Session	Additional Hours per Year	Total Additional Hours over 4 Years
Enhanced Practical OSCEs	Biannual	1 hour each	2 hours	8 hours
Mock Examinations (Comprehensive)	Annual	6 hours each	6 hours	24 hours
Additional Case-Based Discussions	Monthly	20 minutes each	4 hours	16 hours
Focused Skills Workshops	Quarterly	2 hours each	8 hours	32 hours
WPBA sessions (External reviewers)	Biannual	30 minutes each	1 hours	4 hours
Additional Log Book & Portfolio Reviews	Quarterly	1 hour each	8 hours	32 hours

# **Proposed Additional Allocation of Remaining Hours**

Total Additional Hours: 116 hours



# Assessment framework based on Miller's Pyramid and ACGME core competencies for the MS Ophthalmology program



### Weightage of Core Competencies in Assessment

Core Competency	Weightage
Medical knowledge	40% both (106 hours)
Patient care	
Interpersonal and communication skills	40% both (106 hours)
Professionalism	
Practice based learning	10% both (26 hours)
System based practice	
Research	10% (26 hours)



## Formative Assessment

#### 1. Workplace-Based Assessments

Purpose: Continuous evaluation of resident performance, skills, and knowledge in real-world settings.

#### Assessment Tools:

Multi-Source Feedback: Collect feedback from various healthcare professionals.

OCEX/mini-CEX

ICO-OCEX

ICO-OSCAR

Direct Observation of Procedural Skills (DOPS)

The workplace-based assessment methods include feedback opportunities as an integral part of the assessment process. All assessments will be documented in the trainee's e-logbook and portfolio.

#### Schedule of WPBA

Frequency: Conducted biannually (internal and external evaluations).

#### 2. LMS-Based Assessments:

Regularly scheduled online quizzes based on core content and designed to evaluate theoretical knowledge and understanding of ophthalmology concepts.

#### Schedule of LMS-based Assessments

**Frequency**: Conducted fortnightly (15<sup>th</sup> and 30<sup>th</sup> of every month).

## 3. Log Book and Portfolio:

Maintain a record of clinical experiences and skills throughout the program.

#### 4. Annual Evaluation:

Assess resident performance based on:

Competence in skills.

Completion of assignments.

Attitude and behavior.

Participation in journal clubs, lectures, presentations, and clinico-pathologic conferences

## Summative Assessment

## 1. In-Training Assessment for First Year

All candidates admitted in MS Ophthalmology course shall appear in an examination at end of the first calendar year.



**Components:** SEQs (Structured Essay Questions) and OSCE (Objective Structured Clinical Examination).

Pass Percentage: 60%.

#### 2. In-Training Assessment for Third Year

All candidates admitted in MS Ophthalmology course shall appear in an examination at end of the third calendar year.

Components: SEQs and Clinical OSCE.

Pass Percentage: 60%.

## 3.Midterm Assessment MTA

All candidates admitted in MS Ophthalmology course shall appear in Midterm examination at the end of second calendar year.

The examination shall be held on biannual basis.

The candidate who fails to pass the examination in 3 consecutive attempts availed or unavailed, shall be dropped from the course.

Subjects to be examined shall be Basic Ophthalmic Medical Sciences (Anatomy, Physiology, Biochemistry, Pathology, Pharmacology), Optics & Refraction, Behavioral Sciences, Biostatistics & Research Methodology and Community Ophthalmology.

Only those candidates, who pass in theory papers, will be eligible to appear in the TOACS.

The candidates, who have passed written examination but failed in TOACS, will re-appear only in TOACS.

The maximum number of attempts to re-appear in TOACS alone shall be three, after which the candidate shall have to appear in both written and TOACS as a whole.

To be eligible to appear in midterm assessment the candidate must submit;

Duly filled, prescribed Admission Form to the Controller of Examinations duly recommended by the Principal/Head of the Institution in which he/she is enrolled.

A certificate by the Principal/Head of the Institution, that the candidate has attended at least 75% of the lectures, seminars, practical/clinical demonstrations.

Examination fee as prescribed by the University.

To be declared successful in midterm examination the candidate must secure 60% marks in each paper

#### Frequency: Biannual

## **Components:**

Paper I: MCQs (Optics & Refraction) - 75 Marks

Paper II: MCQs - 75 Marks

TOACS: 150 Marks (15 interactive stations)



## **Eligibility:**

75% attendance, submitted forms, and examination fee.

Pass Percentage: 60% in each paper.

### 4. Final Term Assessment (FTA)

All candidates admitted in MS Ophthalmology course shall appear in FTA at the end of structured training program (end of 4<sup>th</sup> calendar year), and having passed MTA. However, a candidate holding FCPS Ophthalmology / FRCS Ophthalmology / Diplomat American Board shall be exempted from MTA and shall be directly admitted to FTA, subject to fulfillment of requirements for the examination.

The examination shall be held on biannual basis.

To be eligible to appear in FTA the candidate must submit;

duly filled, prescribed Admission Form to the Controller of Examinations duly recommended by the Principal/Head of the Institution in which he/she is enrolled;

a certificate by the Principal/Head of the Institution, that the candidate has attended at least 75% of the lectures, seminars, practical/clinical demonstrations;

Original Log Book complete in all respect and duly signed by the Supervisor (for Oral & practical/clinical Examination);

certificate of having passed the midterm examination;

certificates of all the mandatory rotations;

Examination fee as prescribed by the University.

Only those candidates, who pass in theory papers, will be eligible to appear in the Oral & Practical/ Clinical Examination.

The candidates, who have passed written examination but failed in Clinical Examination, will re-appear only in three consecutives. Clinical examination after which the candidate shall have to appear in both written and clinical examinations as a whole.

The candidate with 80% or above marks shall be deemed to have passed with distinction.

## Log Book/Assignments:

Throughout the length of the course, the work record of the candidate shall be entered on the Log Book.

The Supervisor shall certify every year that the Log Book is being maintained and signed regularly.

The performance of the candidate shall be evaluated on annual basis, e.g., 25 marks for each year in four years MS Ophthalmology course. The internal assessment shall reflect the performance of the candidate on following parameters:

Year wise record of the competence of skills.



Year wise record of the assignments.

Year wise record of the evaluation regarding attitude & behavior.

Year wise record of journal club / lectures / presentations / clinico-pathologic conferences attended & / or made by the candidate.

Eligibility: Passed midterm assessment, submitted required documentation.

#### **Components:**

Written Exam: 200 marks

Paper I: MCQs - 100 Marks

**Paper II**: MCQs – 100 Marks

Clinical Examination: 450 marks

OSCE – 150 marks (15 stations)

Long Case – 100 marks

Short Cases - 200 marks (50 marks each)

Pass Requirements: 60% in each component and 50% in each sub-component.

**Re-Assessment Policies:** 

Re-Appears: Candidates who fail any component will have specified re-examination attempts.

**Distinction:** Candidates scoring 80% or above in the final examination will be deemed to have passed with distinction.

## Table Of Specifications

## 1st Year In-House Training TOS

Year One In-Training Assessment- Total marks 200						
Exam Component	No. of Questions/Stations	Marks Distribution	Total Marks	Passing Marks		
Written (MCQ)	100	1 mark each	100	60		
Clinical - OSCE	10	10 marks each	100	60		
AV-OSPE	10	10 marks each	100	60		



Written component Table of Specifications

- **O** Section (I) Clinical Ophthalmology
- O No. of items (50)
  - O Level of questions (according to Bloom's taxonomy)
  - O C1
  - O C2
  - О СЗ

S. No	Торіс	Impact	Frequency	I × F	Weight	No of items	Diagnosis	Investigation	Treatment	Basic knowledge
1	Eyelids	3	2	6	0.05	2.56=2	1	0	1	0
2	Lacrimal drainage system	3	2	6	0.05	2.56=2	1	0	1	0
3	Orbit	3	2	6	0.05	2.56=2	1	1	0	0
4	Dry eye disorders	2	2 4		0.034	1.70=2	1	1	0	0
5	Conjunctiva	2	2	4	0.034	1.70=2	1	0	1	0
6	Cornea	2	3	6	0.05	2.56=2	1	1	0	0
7	Corneal and Refractive surgery	2	2	4	0.034	1.70=2	1	0	1	0
8	Episclera and sclera	2	2	4	0.034	1.70=2	1	0	1	0
9	Lens	3	3	9	0.076	3.84=4	1	1	1	1
10	Glaucoma	3	3	9	0.076	3.84=4	1	1	1	1
11	Uveitis	2	3	6	0.05	2.56=2	1	1	0	0
12	Ocular Tumors	2	1	2	0.017	0.85=1	1	0	0	0
13	Retinal Vascular disease	3	3	9	0.076	3.84=4	1	1	1	1
14	Acquired Macular Disorders	2	2	4	0.034	1.70=2	1	1	0	0
15	Hereditary Fundus Dystrophies	2	3	6	0.05	2.56=2	1	1	0	0



							MEDICH	tumuipinui mee		- 9
16	Retinal Detachment	2	3	6	0.05	2.56=2	1	0	1	0
17	Strabismus	2	3	6	0.05	2.56=2	1	0	1	0
18	Vitreous Opacities	1	2	2	0.017	0.85=1	1	0	0	0
19	Neuro- Ophthalmology	2	3	6	0.05	2.56=2	1	1	0	0
20	Ocular side effects of systemic medications	1	1	1	0.008	0.407=0	0	0	0	0
21	Ocular Trauma	2	3	6	0.05	2.56=2	1	0	1	0

- O Discipline Ophthalmology
- O Level of exam (First Year In-house)
- **O** Section (II) Optics and Refraction
- O No. of items (50)
  - O Level of questions (according to Bloom's taxonomy)
  - O C1
  - O C2
  - О СЗ

S. No	Торіс	Impact	Frequency	I × F	Weight	No of items	Diagnosis	Investigation	Treatment	Basic knowledge
1	Physical Optics	3	2	6	0.103	5.17=5	1	0	0	4
2	Geometrical Optics	3	2	6	0.103	5.17=5	1	0	0	4
3	Optics of human eye	3	2	6	0.103	5.17=5	1	0	0	4
4	Practical clinical refraction	2	3	6	0.103	5.17=5	1	0	2	2
5	Contact lenses	2	2	4	0.068	3.45=3	1	0	1	1
6	Visual rehabilitation	2	3	6	0.103	5.17=5	1	0	1	3
7	Intraocular lenses	2	2	4	0.068	3.45=3	1	0	1	1
8	Lasers	2	2	4	0.068	3.45=3	1	0	1	1



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9	Optical	3	3	9	0.155	7.75=8	3	0	1	4
	considerations									
	in refractive									
	surgery									
1	0 Optical	3	3	9	0.155	7.75=8	2	0	1	5
	instruments									

# **Clinical Component Table of Specifications**

OSCE/AV-OSPE stations- 10 Total marks- 100 Passing marks- 60 Time allowed per station- 5min Interactive stations- 5

# **Topic Wise Distribution of Ophthalmology OSCE/AV-OSPE Stations**

Station No.	Station Description & Topics	Skill to be assessed
1.	Clinical Methods <ul> <li>Pupil Evaluation</li> <li>Extraocular movements</li> </ul>	To assess the candidate's ability to perform the given examination task on a patient/simulated subject.
2.	<ul> <li>Clinical Methods</li> <li>Slit lamp examination techniques</li> <li>Tonometry</li> <li>Direct ophthalmoscopy</li> </ul>	To assess the candidate's ability to perform the given examination task on a patient/simulated subject.
3.	<ul><li>Investigations</li><li>Fundus fluorescein angiography</li></ul>	Images/videos will be shown to the candidate with relevant clinical scenarios to assess the ability to interpret findings and discuss the management plan
4.	Investigations <ul> <li>Visual fields</li> </ul>	Images/videos will be shown to the candidate with relevant clinical scenarios to assess the ability to interpret findings and discuss the management plan
5.	Ophthalmic Radiology • B-scan	Images/videos will be shown to the candidate with relevant clinical scenarios to assess the ability to interpret findings, make a diagnosis, and discuss the management/ complications.



6.	Clinical Problem Solutions <ul> <li>Cataract</li> <li>Glaucoma</li> <li>Cornea</li> <li>Medical Retina</li> <li>Surgical Retina</li> </ul>	A clinical scenario will be given to the candidate to assess the ability to interpret the findings, make a diagnosis, and discuss the management/ complications.
7.	Retinoscopy	To assess the ability of the candidate to perform steps of procedure on patient/simulated subject.
8.	Transposition	To assess the ability of the candidate to write a refractive prescription and transpose the refractive prescription of the retinoscopy findings
9.	Instruments. <ul> <li>Cross-cylinder</li> <li>Maddox Rods</li> <li>Focimeter</li> <li>Maddox Wings</li> <li>Low vision aids</li> </ul>	To assess the candidate's ability to demonstrate the use of a given item on a patient/simulated subject. To assess the ability of candidate to identify the equipment and its uses
10.	Biometry • A-scan • Keratometry	To assess the ability of the candidate to perform steps of the asked procedure on the patient/simulated subject and calculate the IOL number using the appropriate formula on the data obtained.

Station No.	Station Description & Topics	Skill to be assessed
1.	Cataract/Glaucoma/Strabismus/Neuro- Ophthalmology	To assess the ability of the candidate to interpret findings and discuss the management plan amicably.
2.	Trauma/Keratitis/Conjunctivitis/Uveitis	To assess the candidate's ability to interpret findings, make diagnosis and discuss the management plan.
3.	Ophthalmic pathology	To assess the candidate's ability to interpret findings, make diagnosis and discuss the management plan



4.	Investigations <ul> <li>Fundus fluorescein angiography</li> <li>OCT</li> </ul>	Images/videos will be shown to the candidate with relevant clinical scenarios to assess the ability to interpret findings and discuss the management plan			
5.	Surgical retina	A clinical scenario will be given to the candidate to assess the ability to interpret the findings, make a diagnosis, and discuss the management/complications.			
6.	Ophthalmic Emergencies <ul> <li>Acute Congestive Glaucoma</li> <li>Chemical burns,</li> <li>Trauma</li> </ul>	To assess the ability of the candidate to amicably approach the emergency and perform steps of management.			
7.	Investigations <ul> <li>Corneal Topography/Visual</li> <li>Fields</li> </ul>	Images/videos will be shown to the candidate to assess the ability to interpret the findings, make a diagnosis, and discuss the management/ complications.			
8.	Medical retina	A clinical scenario will be given to the candidate to assess the ability to interpret the findings, make a diagnosis, and discuss the management/complications.			
9.	Transposition	To assess the ability of the candidate to write a refractive prescription and transpose the refractive prescription of the retinoscopy findings			
10.	Instruments. <ul> <li>Cross-cylinder</li> <li>Maddox Rods</li> <li>Focimeter</li> <li>Maddox Wings</li> <li>Low vision aids</li> </ul>	To assess the candidate's knowledge of principle and the use of a given item. To assess the ability of candidate to identify the equipment and its uses			



# Midterm Assessment TOS Calgary Model

# PAPER II (Clinical Ophthalmology)

**O** No. of items (75)

S. No	Торіс	Presentation	Impact	Frequency	I × F	Weight	No of items	Diagnosis	Investigation	Treatment	Basic knowledge
1	Eyelids	Anatomy & physiology	3	2	6	0.017	1.21=1	0	0	0	1
		Benign nodules, cysts and malignant tumors	1	2	2	0.005	0.40=0	0	0	0	0
		Bacterial and viral infections, Allergic disorders and blepharitis	1	3	3	0.008	0.6=1	0	0	1	0
		Ptosis, entropion and ectropion	2	3	6	0.017	0.75=1	0	0	1	0
2	Lacrimal drainage	Anatomy and physiology	3	2	6	0.017	1.21=1	0	0	0	1
	system	Congenital and acquired obstruction	1	3	3	0.008	0.6=1	0	1	0	0
		Chronic canaliculitis	1	1	1	0.0027	0.20=0	0	0	0	0
		Dacryocystitis	2	3	6	0.017	1.21=1	1	0	0	0
3	Orbit	Anatomy	3	2	6	0.016	1.21=1	0	0	0	1
		Thyroid eye disease	2	2	4	0.010	0.75=1	0	1	0	0
		Infections and Inflammatory disease	2	3	6	0.017	1.21=1	1	0	0	0

	Vascular malformations and tumors	2	2	4	0.010	0.75=1	0	1	0	0
	Cystic lesions	1	3	3	0.008	0.6=1	1	0	0	0



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4	Dry eye disorders	Anatomy and physiology of tear film	2	2	4	0.010	0.75=1	0	0	0	1
		Dry eye disorders	1	3	3	0.008	0.6=1	0	1	0	0
5	Conjunctiva	Anatomy and physiology	2	2	4	0.010	0.75=1	0	0	0	1
		Bacterial, viral and allergic conjunctivitis	1	3	3	0.008	0.6=1	0	0	1	0
	Cornea	Cicatrizing conjunctivitis and degenerations	1	2	2	0.005	0.40=0	0	0	0	0
6	Cornea	Anatomy and physiology	2	2	4	0.010	0.75=1	0	0	0	1
		Bacterial, viral and fungal keratitis	2	3	6	0.017	1.21=1	0	0	1	0
		Interstitial and protozoan keratitis	2	2	4	0.010	0.75=1	1	0	0	0
		Rosacea, Neurotrophic and exposure keratitis, Bacterial hypersensitivity mediated corneal disease	2	2	4	0.010	0.75=1	1	0	0	0
		Corneal ectasias	2	2	4	0.008	0.6=1	0	0	1	0

	Corneal	1	1	1	0.0027	0.20=0	0	0	0	0
	dystrophies									
	and									
	degenerations									



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		Peripheral corneal ulceration and Metabolic keratopathies	1	1	1	0.0028	0.40=0	0	0	0	0
7	Corneal	Keratoplasty	2	2	4	0.0011	0.8=1	0	0	1	0
	and refractive surgery	Refractive surgeries	2	2	4	0.010	0.75=1	0	0	1	0
8	Episclera	Anatomy	1	1	1	0.0027	0.20=0	0	0	0	0
	and sclera	Episcleritis and scleritis	2	2	4	0.011	0.8=1	1	0	0	0
		Scleral discoloration	1	1	1	0.0027	0.20=0	0	0	0	0
9	Lens	Anatomy and physiology	2	2	4	0.010	0.75=1	0	0	0	1
	с С	Congenital cataract	2	2	4	0.011	0.8=1	0	0	1	0
		Acquired cataract	3	3	9	0.025	1.8=2	0	1	1	0
		Ectopia lentis	2	2	4	0.010	0.75=1	1	0	0	0
10	Glaucoma	Anatomy and physiology	2	2	4	0.010	0.75=1	0	0	0	1
		Tonometry, gonioscopy and perimetry	2	2	4	0.010	0.75=1	0	1	0	0
	F c g F a	Primary openangle glaucoma	3	3	9	0.025	1.8=2	0	1	1	0
		Primary angleclosure glaucoma	2	2	4	0.011	1.21=1	0	0	1	0
		Ocular hypertension and	1	2	2	0.005	0.40=0	0	0	0	0

Normal tension glaucoma									
Pseudo exfoliation and Inflammatory glaucoma	2	2	4	0.010	0.75=1	1	0	0	0



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		Pigment dispersion glaucoma and Iridocorneal endothelial syndrome	2	2	4	0.0028	0.4=1	0	0	1	0
		Neovascular and lens-related glaucoma	3	2	6	0.017	1.21=1	0	0	1	0
		Traumatic glaucoma and Glaucoma in intraocular tumors	1	1	1	0.0028	0.40=0	0	0	0	0
		Primary congenital glaucoma	2	2	4	0.010	0.75=1	1	0	0	0
		Phacomatosis	1	1	1	0.0028	0.40=0	0	0	0	0
11	1 Uveitis	Anatomy and physiology	2	2	4	0.010	0.75=1	0	0	0	1
		Anterior uveitis	2	3	6	0.017	1.21=1	1	0	0	0
		Intermediate uveitis	2	2	4	0.010	0.75=1	0	0	1	0
		Uveitis in spondyloarthropat hies and juvenile arthritis	1	3	3	0.008	0.6=1	0	0	1	0
		Sarcoidosis and Bechet syndrome	1	2	2	0.005	0.40=0	0	0	0	0

		Vogt-koyanagi Harada syndrome	1	1	1	0.0028	0.40=0	0	0	0	0
		Bacterial and viral uveitis	2	2	4	0.010	0.75=1	1	0	0	0
		Fungal and parasitic uveitis, Uveitis in bowel and renal disease	1	1	1	0.0027	0.20=0	0	0	0	0
12	Ocular tumors	Benign and malignant conjunctival tumors	1	2	2	0.005	0.40=0	0	0	0	0



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		Iris cysts and tumors, Ciliary body tumors	1	1	1	0.0027	0.20=0	0	0	0	0
		Tumors of the retina and choroid	2	2	4	0.010	0.75=1	1	0	0	0
		Primary intraocular lymphoma	1	1	1	0.005	0.40=0	0	0	0	0
		Paraneoplastic syndromes	1	1	1	0.0027	0.20=0	0	0	0	0
13	8 Retinal vascular disease	Diabetic retinopathy	3	3	9	0.025	1.8=2	0	1	1	0
		Retinal venous occlusive disease	3	2	6	0.017	1.21=1	0	0	1	0
		Retinal arterial occlusive disease	3	1	3	0.008	0.6=1	0	0	1	0
		Ocular ischemic	1	1	1	0.0027	0.40=0	0	0	0	0
		syndrome and Sickle cell retinopathy									
		reunopatity									

		Hypertensive disease	2	2	4	0.010	0.75=1	0	0	1	0
		Retinopathy of prematurity	2	2	4	0.010	0.75=1	1	0	0	0
		Retinal artery macro aneurysm and Primary retinal telangiectasis	1	1	1	0.0027	0.20=0	0	0	0	0
1		Eales disease	2	2	4	0.010	0.75=1	1	0	0	0
		Radiation retinopathy and Purtscher retinopathy	1	1	1	0.0027	0.20=0	0	0	0	0



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		Takayasu Disease	1	1	1	0.0027	0.2=0	0	0	0	0
14	Acquired macular disorders	Imaging in macular disease	2	2	4	0.010	0.75=1	0	1	0	0
		Age related macular degeneration	3	2	6	0.017	1.2=1	1	0	0	0
		Age-related macular hole	2	3	6	0.017	1.2=1	0	1	0	0
		Central serous retinopathy and Cystoid macular edema	2	2	4	0.011	0.8=1	1	0	0	0
		Macular epiretinal membrane and Polypoidal	1	1	1	0.0027	0.40=0	0	0	0	0

		choroidal vasculopathy									
		Degenerative myopia	1	3	3	0.008	0.6=1	0	0	1	0
		Angioid streaks and Vitreomacular traction syndrome	1	1	1	0.0027	0.40=0	0	0	0	0
		Idiopathic choroidal neovascularization	1	1	1	0.0027	0.2=0	0	0	0	0
		Solar retinopathy	1	1	1	0.0027	0.20=0	0	0	0	0
15	5 Hereditary Fundus dystrophies	Macular dystrophies and Choroidal dystrophies	1	3	3	0.008	0.6=1	1	0	0	0
		Generalized photoreceptor dystrophies	1	3	3	0.008	0.6=1	0	1	0	0
		Albinism and Cherry red spot at macula	1	1	1	0.0027	0.2=0	0	0	0	0



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16	Retinal detachment	Rhegmatogenous	3	3	9	0.024	1.82=2	0	1	1	0
		retinal									
		detachment									
		Tractional retinal detachment	2	2	4	0.010	0.75=1	0	0	1	0
		Exudative retinal detachment	1	3	3	0.008	0.6=1	0	0	1	0
		Pars plana vitrectomy	2	2	4	0.011	1.1=1	0	0	1	0
17	Strabismus	Amblyopia	1	2	2	0.0056	0.4=1	0	0	1	0

		Heterophoria and vergence abnormalities	1	2	2	0.0056	0.4=0	0	0	0	0
		Esotropia	1	3	3	0.008	0.6=1	0	0	1	0
		Exotropia	1	3	3	0.008	0.6=1	0	0	1	0
		Congenital cranial dysinnervation disorders	1	1	1	0.0027	0.20=0	0	0	0	0
		Monocular elevation deficiency and Brown syndrome	1	1	1	0.0028	0.20=0	0	0	0	0
		Alphabet patterns	1	2	2	0.0056	0.4=0	0	0	0	0
18	Vitreous opactities	Vitreous hemorrhage	2	3	6	0.017	1.21=1	0	0	1	0
19	Neuroophthalmol	Neuroimaging	2	2	4	0.010	0.75=1	0	1	0	0
	ogy	Optic nerve, pupil, chiasma, retro chiasmal pathways	2	3	6	0.017	1.21=1	1	0	0	0
		Ocular motor nerves	2	3	6	0.017	1.21=1	1	0	0	0
		Supranuclear disorders of ocular motility	1	1	1	0.0027	0.20=0	0	0	0	0
		Nystagmus and Facial spasm	1	1	1	0.0027	0.20=0	0	0	0	0



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		Ocular myopathies and Neurofibromatosis	1	3	3	0.008	0.6=1	1	0	0	0
		Migraine and Neuralgias	1	1	1	0.0027	0.2=0	0	0	0	0
20	Ocular side	Cornea	1	1	1	0.0027	0.2=0	0	0	0	0
	effects of systemic	Ciliary effusion and lens	1	1	1	0.0027	0.2=0	0	0	0	0
	medications	Optic nerve	2	3	6	0.017	1.21=1	1	0	0	0
21	Ocular trauma	Eyelid trauma	1	3	3	0.008	0.6=1	0	0	1	0
		Orbital trauma	2	2	4	0.010	0.75=1	1	0	0	0
		Trauma to globe	3	2	6	0.017	1.21=1	0	0	1	0
		Chemical injuries	2	3	6	0.017	0.017=1	0	0	1	0

Level of questions (according to Bloom's taxonomy)

C1= 9 , C2 = 27 , C3 = 39

## **MIDTERM EXAMINATION MTA**

# PAPER I (Refraction)

**O** No. of items (75)

S. No	Торіс	Presentation	Impact	Frequency	I × F	Weight	No of items	Diagnosis	Investigation	Treatment	Basic knowledge
1	Physical optics	Interference and coherence	2	2	4	0.011	0.8=1	0	0	0	1
		Polarization	2	2	4	0.011	0.8=1	1	0	0	0
		Diffraction	2	2	4	0.011	0.8=1	1	0	0	0
		Scattering	2	2	4	0.011	0.8=1	1	0	0	0
		Reflection	2	3	6	0.016	1.2=1	1	0	0	0
		Illumination	2	2	4	0.011	0.8=1	0	0	0	1
		Transmission and absorption	2	2	4	0.011	0.8=1	0	0	0	1
		Laser fundamentals	3	2	6	0.016	1.2=1	0	0	0	1
2	Geometrical optics	Pinhole imaging	3	3	9	0.025	1.8=2	1	0	0	1



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Imaging with lenses and mirrors	3	3	9	0.025	1.8=2	1	0	0	1
Image and object characteristics	3	3	9	0.025	1.8=2	1	0	0	1
Light propagation	2	2	4	0.011	0.8=1	0	0	0	1
Ophthalmic lenses	3	3	9	0.025	1.8=2	1	0	0	1
Ophthalmic prisms	3	3	9	0.025	1.8=2	1	0	0	1
Mirrors	2	2	4	0.011	0.8=1	1	0	0	0
Optical aberrations	3	3	6	0.016	1.2=1	0	0	0	1

3	Optics of human eye	Human eye as an optical system	3	2	6	0.016	1.2=1	0	0	0	1
		Schematic eyes	3	2	6	0.016	1.2=1	0	0	0	1
		Visual acuity	3	3	9	0.025	1.8=2	1	0	0	1
		Contrast sensitivity	3	2	6	0.016	1.2=1	0	0	0	1
		Refractive states of eye	3	3	9	0.025	1.8=2	1	0	0	1
4	Practical Clinical	Objective refraction	3	3	9	0.025	1.8=2	1	0	0	1
	refraction	Subjective refraction	3	3	9	0.025	1.8=2	1	0	0	1
		Cycloplegic and non-cycloplegic refraction	3	2	6	0.016	1.2=1	0	0	0	1
		Spherical and cylindrical correcting lenses	3	2	6	0.016	1.2=1	0	0	0	1
		Prescribing for children	3	2	6	0.016	1.2=1	0	0	1	0
		Clinical accommodative problems	3	2	6	0.016	1.2=1	1	0	0	0



			Read Internet Medical Children							/		
			Prescribing multifocal lenses	3	2	6	0.016	1.2=1	0	0	1	0
			Prescribing special lenses	3	1	3	0.008	0.6=1	0	0	1	0
	5	Contact lenses	Contact lens optics	3	2	6	0.016	1.2=1	0	0	0	1
			Contact lens materials	3	2	6	0.016	1.2=1	0	0	0	1

		Contact lens fitting	3	3	9	0.025	1.8=2	1	0	0	1
		Therapeutic lens usage	3	2	6	0.016	1.2=1	0	0	1	0
6	Visual rehabilitatio	Low vision and legal blindness	3	2	6	0.016	1.2=1	1	0	0	0
	n	Optics of low vision aids, convex lens	3	2	6	0.016	1.2=1	0	0	0	1
7	Intraocular lenses	Optical considerations for IOLs	3	2	6	0.016	1.2=1	0	0	0	1
		Lens related visual disturbances	З	3	9	0.025	1.8=2	1	0	0	1
		Multifocal IOLs	3	1	3	0.008	0.6=1	0	0	0	1
8	Lasers	Types of lasers	3	2	6	0.016	1.2=1	0	0	0	1
		Laser modes	2	2	4	0.011	0.8=1	0	0	1	0
		Effects of laser energy on tissues	3	2	6	0.016	1.2=1	0	0	0	1
9	Optical consideratio ns in	Angle kappa, corneal shape pupil size	2	1	2	0.006	0.4=0	0	0	0	0
	refractive surgery	Surgical correction of myopia, hypermetropia and presbyopia	3	3	9	0.025	1.8=2	1	0	1	0
		Surgical corrections of astigmatism	3	3	9	0.025	1.8=2	0	1	1	0



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		Vitreoretinal surgery	3	2	6	0.016	1.2=1	0	0	0	1
10	Optical instruments	Direct and indirect ophthalmoscope	3	3	9	0.025	1.8=2	1	0	0	1
		Retinoscope	3	3	9	0.025	1.8=2	1	0	0	1
		Instrument used to study corneal curvature	3	2	6	0.016	1.2=1	0	1	0	0
		Computerized analysis of corneal topography	2	2	4	0.011	0.8=1	0	1	0	0
		Slit lamp biomicroscope	3	3	9	0.025	1.8=2	1	0	0	1
		Lensometer	3	3	9	0.025	1.8=2	0	1	0	1
		Applanation tonometer	3	3	9	0.018	1.8=2	1	0	0	1
		Pachymeter	3	2	6	0.016	1.2=1	0	1	0	0
		Operating microscope	2	2	4	0.011	0.8=1	0	0	0	1
		Specular microscope	1	1	1	0.003	0.2=0	0	0	0	0
		Keratometer	3	3	9	0.025	1.8=2	0	1	0	1
		Macular function testing equipment	3	2	6	0.016	1.2=1	0	1	0	0
		Wavefront aberrometer	1	1	1	0.003	0.2=0	0	0	0	0
		Optical coherence tomography	3	2	6	0.016	1.2=1	0	1	0	0

Level of questions (according to Bloom's taxonomy)

C1 =38 , C2 =21 , C3 =16

# **TOS for OSCE in Mid-Training Assessment**

- 1. Total number of stations 15 (All Interactive)
- 2. Time Allocation for each station -5 Minutes
- 3. Marks Allocation for each station 10 Minutes



# **Topic Wise Distribution of Ophthalmology OSCE Stations**

Station No.	Station Description & Topics	Skill to be assessed
1.	Counseling (Scenario Based) <ul> <li>Retinoblastoma</li> <li>Diabetic Retinopathy</li> <li>Cataract</li> <li>Glaucoma</li> </ul>	In a given scenario the candidate's ability to counsel the family about diagnosis, its implications, and management options.
2.	Retinoscopy	To assess the ability of the candidate to perform steps of procedure on patient/simulated subject.
3.	Transposition	To assess the ability of the candidate to write a refractive prescription and transpose the refractive prescription of the retinoscopy findings
4.	Instruments. • Cross-cylinder • Maddox Rods • Focimeter • Maddox Wings • Low vision aids	To assess the candidate's ability to demonstrate the use of a given item on a patient/simulated subject. To assess the ability of candidate to identify the equipment and its uses
5.	Clinical Methods <ul> <li>Pupil Evaluation</li> <li>Extraocular movements</li> <li>Cover- Uncover Test</li> </ul>	To assess the candidate's ability to perform the given examination task on a patient/simulated subject.
б.	Clinical Methods <ul> <li>Optic Nerve function test</li> <li>Visual fields</li> <li>Macular function test</li> </ul>	To assess the candidate's ability to perform the given examination task on a patient/simulated subject.
7.	Clinical Methods <ul> <li>Slit Lamp Examination</li> <li>Different illumination techniques</li> <li>Tests for dry eye</li> </ul>	To assess the ability of the candidate to perform steps of the asked procedure on the patient/simulated subject.
8.	Surgical Skills	To assess the ability of the



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	<ul> <li>Eyelid Laceration Repair</li> <li>Cystotome formation</li> <li>Partial thickness Incision for extracapsular cataract extraction</li> <li>Full-thickness incision for extracapsular cataract extraction</li> <li>Sutures</li> <li>Ophthalmic drops</li> <li>Intravitreal Injections</li> </ul>	candidate to perform steps of the asked procedure on a goat eye/provided specimen.
9.	<ul><li>Investigations</li><li>Fundus fluorescein angiography</li><li>Visual fields</li></ul>	Images/videos will be shown to the candidate with relevant clinical scenarios to assess the ability to interpret findings and discuss a management plan
10.	Ophthalmic Radiology • B-scan • CT Scan • MRI	Images/videos will be shown to the candidate with relevant clinical scenario to assess the ability to interpret findings, make a diagnosis and discuss management with complication.
11.	Clinical Problem Solutions <ul> <li>Anterior Segment</li> </ul>	The candidate will be required to assess the ability to examine the patient interpret the findings, make a diagnosis and discuss management with complication.
12.	Clinical Problem Solutions <ul> <li>Posterior Segment</li> </ul>	The candidate will be required to assess the ability to examine the patient interpret the findings, make a diagnosis and discuss management with complication.
13.	<ul> <li>Ophthalmic Emergencies</li> <li>Acute Congestive Glaucoma</li> <li>Central Retinal artery Occlusion</li> <li>Chemical burns,</li> <li>Trauma</li> </ul>	To assess the ability of the candidate to amicably approach the emergency and perform steps of management.



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14.	Biometry • A-scan • Keratometer	To assess the ability of the candidate to perform steps of the asked procedure on the patient/simulated subject and calculate the IOL number using the appropriate formula on the data obtained.
15.	General Surgery Or Emergency Medicine	To assess the ability of candidate to perform steps of asked procedure on patient/simulated subject/mannequin



Third Year In-Training Assessment- Total marks 200										
Exam Component	No. of Questions/stations	Marks Distribution	Total Marks	Passing Marks						
Written (MCQs)	100	1 mark each	100	60						
Clinical - OSCE	10	10 marks each	100	60						
AV-OSPE	10	10 marks each	100	60						

## **THEORY TABLE OF SPECIFICATIONS**

- **O** Section (I) Clinical Ophthalmology
- O No. of items (50)
  - O Level of questions (according to Bloom's taxonomy)
  - O C2
  - О СЗ

S. No	Торіс	Impact	Frequency	I × F	Weight	No of items	Diagnosis	Investigation	Treatment	Basic knowledge
1	Eyelids	3	2	6	0.103	5.17=5	2	1	2	0
2	Lacrimal drainage system	3	2	6	0.103	5.17=5	2	1	2	0
3	Orbit	3	2	6	0.103	5.17=5	2	1	2	0
4	Dry eye disorders	2	2	4	0.068	3.45=3	1	1	1	0
5	Conjunctiva	2	3	6	0.103	5.17=5	2	1	2	0
6	Cornea	2	3	6	0.103	5.17=5	2	1	2	0
7	Corneal and Refractive surgery	2	2	4	0.068	3.45=3	2	0	1	0



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8	Episclera and sclera	2	2	4	0.068	3.45=3	1	1	1	0
9	Lens	3	3	9	0.155	7.75=8	3	2	3	0
10	Glaucoma	3	3	9	0.155	7.75=8	3	2	3	0

- O Discipline Ophthalmology
- O Level of exam (Third Year In-house)
- **O** Section (II) Clinical Ophthalmology
- O No. of items (50)
  - O Level of questions (according to Bloom's taxonomy)
  - O C2
  - О СЗ

S.	Торіс	Impact	Frequency	I × F	Weight	No of	Diagnosis	Investigation	Treatment	Basic
No						items				knowledge
1	Uveitis	2	3	6	0.096	4.83=5	2	1	2	4
2	Ocular Tumors	2	1	2	0.032	1.612=2	1	0	1	0
3	Retinal Vascular disease	3	3	9	0.145	7.25=7	3	1	3	0
4	Acquired Macular Disorders	2	2	4	0.064	3.22=3	1	1	1	0
5	Hereditary Fundus Dystrophies	2	3	6	0.096	4.83=5	3	1	1	0
6	Retinal Detachment	3	3	9	0.145	7.25=7	3	1	3	0
7	Strabismus	2	3	6	0.096	4.83=5	2	1	2	0
8	Vitreous Opacities	2	2	4	0.064	3.22=3	2	0	1	0
9	Neuro- Ophthalmology	2	3	6	0.096	4.83=5	3	1	1	0
10	Ocular side effects of systemic medications	1	1	1	0.016	0.806=1	1	0	0	0
11	Ocular Trauma	3	3	9	0.145	7.25=7	3	1	3	0



#### **CLINICAL COMPONENT TABLE OF SPECIFICATIONS**

OSCE stations- 10 Total marks- 100 Passing marks- 60 Time allowed per station- 5min Interactive stations- 5

#### **Topic Wise Distribution of Ophthalmology OSCE/AV-OSPE Stations**

No of Stations	Station Description & Topics	Competence to be assessed				
3	Ophthalmic Investigations Fundus fluorescein angiography, OCT, Visual fields, Corneal topography, Hess chart	To assess the ability of the candidate to interpret the investigation and answer the questions (Critical thinking & Problem solving)				
1	Clinical Methods <ul> <li>Pupil Evaluation</li> <li>Extraocular movements</li> <li>Squint assessment</li> <li>Visual fields by confrontation</li> </ul>	To assess the candidate's ability to perform the given examination task on a patient/simulated subject.				
1	Clinical Methods <ul> <li>Slit lamp examination</li> <li>techniques</li> <li>Tonometry</li> <li>Indirect ophthalmoscopy</li> <li>Gonioscopy</li> </ul>	To assess the candidate's ability to perform the given examination task on a patient/simulated subject.				
1	Ophthalmic procedures	Images/videos will be shown to the candidate with relevant clinical scenarios to assess the ability to interpret findings and discuss the management plan (Critical thinking & Problem solving)				
1	Ophthalmic Radiology • B-scan • A-scan • CT-Scan/MRI	Images/videos will be shown to the candidate with relevant clinical scenarios to assess the ability to interpret findings, make a diagnosis, and discuss the management/				



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		complications.
1	Clinical Problem Solutions <ul> <li>Cataract</li> <li>Glaucoma</li> <li>Cornea</li> <li>Uveitis</li> <li>Strabismus</li> </ul>	A clinical scenario will be given to the candidate to assess the ability to interpret the findings, make a diagnosis, and discuss the management/ complications. (Clinical reasoning and problem solving)
1	Clinical Problem Solutions <ul> <li>Medical retina</li> </ul>	A clinical scenario will be given to the candidate to assess the ability to interpret the findings, make a diagnosis, and discuss the management/complications. (Clinical reasoning and problem solving)
1	Clinical Problem Solutions <ul> <li>Surgical retina</li> </ul>	A clinical scenario will be given to the candidate to assess the ability to interpret the findings, make a diagnosis, and discuss the management/complications. (Clinical reasoning and problem solving)

AV-OSPE stations- 10 Total marks- 100 Passing marks- 60 Time allowed per station- 5min

#### **Topic Wise Distribution of Ophthalmology AV-OSPE Stations**

Station No:	Station Description & Topics	Competence to be assessed
1	Ophthalmic Investigations Fundus fluorescein angiography, OCT	To assess the ability of the candidate to interpret the investigation and answer the questions (Critical thinking & Problem solving)
2	Ophthalmic Investigations Visual fields	To assess the ability of the candidate to interpret the investigation and answer the questions (Critical thinking & Problem solving)



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3	Ophthalmic Investigations Corneal topography	To assess the ability of the candidate to interpret the investigation and answer the questions (Critical thinking & Problem solving)
4	Ophthalmic Investigations Hess chart	To assess the ability of the candidate to interpret the investigation and answer the questions (Critical thinking & Problem solving)
5	Ophthalmic Radiology • B-scan • A-scan • CT-Scan/MRI	Images/videos will be shown to the candidate with relevant clinical scenarios to assess the ability to interpret findings, make a diagnosis, and discuss the management/ complications.
6	Clinical Problem Solutions • Cataract • Glaucoma	A clinical scenario will be given to the candidate to assess the ability to interpret the findings, make a diagnosis, and discuss the management/ complications. (Clinical reasoning and problem solving)
7	Clinical Problem Solutions <ul> <li>Medical retina</li> </ul>	A clinical scenario will be given to the candidate to assess the ability to interpret the findings, make a diagnosis, and discuss the management/complications. (Clinical reasoning and problem solving)
8	Clinical Problem Solutions <ul> <li>Surgical retina</li> </ul>	A clinical scenario will be given to the candidate to assess the ability to interpret the findings, make a diagnosis, and discuss the management/complications. (Clinical reasoning and problem solving)
9	Clinical Problem Solutions <ul> <li>Cornea</li> <li>Uveitis</li> </ul>	Images/videos will be shown to the candidate with relevant clinical scenarios to assess the ability to interpret findings and discuss the



		management plan (Critical thinking
		& Problem solving)
10	Clinical Problem Solutions <ul> <li>Strabismus</li> <li>Neuro-Ophthalmology</li> </ul>	Images/videos will be shown to the candidate with relevant clinical scenarios to assess the ability to interpret findings and discuss the management plan (Critical thinking & Problem solving)

### Final Term Assessment TOS Calgary Model

- O Paper (I)
- O No. of items (100)

S. No	Торіс	Presentation	Impact	Frequency	I × F	Weight	No of items	Diagnosis	Investigation	Treatment	Basic knowledge
1	Eyelids	Benign nodules and cysts	1	3	3	0.024	2.4=2	1	0	1	0
		Malignant tumors	2	2	4	0.03	3=3	1	1	1	0
		Bacterial and viral infections, Allergic disorders and blepharitis	1	3	3	0.024	2.4=2	1	0	1	0
		Ptosis, entropion and ectropion	2	3	6	0.048	4.8=5	02	0	2	1
2	Lacrimal drainage system	Congenital and acquired obstruction	1	3	S	0.024	2.40=2	0	1	1	0
	•	Chronic canaliculitis	1	1	1	0.007	0.7=1	1	0	0	0
		Dacryocystitis	2	3	6	0.048	4.8=5	0	2	3	0
3	Orbit	Thyroid eye disease	3	2	6	0.048	4.8=5	0	2	2	1
		Infections and Inflammatory disease	2	2	4	0.03	3=3	1	1	1	0
		Vascular malformations and tumors	1	1	1	0.007	0.7=1	0	0	1	0

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	Cystic lesions	1	1	1	0.007	0.7=1	0	0	1	0	
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4	Dry eye	Tear film	2	3	6	0.048	4.8=5	1	2	2	0
	disorders	Dry eye disorders	1	3	3	0.024	2.4=2	0	1	1	0
5	Conjunctiva	Bacterial, viral and allergic conjunctivitis	1	3	3	0.024	2.40=2	0	1	1	0
		Cicatrizing conjunctivitis and degenerations	1	2	2	0.016	1.60=2	0	1	1	0
6	Cornea	Bacterial, viral and fungal keratitis	2	3	6	0.048	4.8=5	1	1	3	0
		Interstitial and protozoan keratitis	2	1	2	0.016	1.6=2	1	0	1	0
		Rosacea, Neurotrophic and exposure keratitis, Bacterial hypersensitivity mediated corneal disease	2	1	2	0.016	1.6=2	1	0	1	0
		Corneal ectasias	1	3	3	0.024	2.4=2	0	1	1	0
		Corneal dystrophies and degenerations	2	1	2	0.016	1.6=2	1	1	0	0
		Peripheral corneal ulceration and Metabolic keratopathies	1	1	1	0.007	0.7=1	0	0	1	0
7	Corneal and refractive surgery	Keratoplasty	2	2	4	0.03	3=3	0	1	2	0

	Refractive	2	2	4	0.03	3=3	0	1	2	0
	surgeries									



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8	Episclera and	Episcleritis and scleritis	1	2	2	0.016	1.6=2	1	0	1	0
	sclera	Scleral discoloration	1	1	1	0.007	0.7=1	1	0	0	0
9	Lens	Congenital cataract	2	2	4	0.03	3=3	1	0	2	0
		Acquired cataract	2	3	6	0.048	4.8=5	1	1	3	0
		Ectopia lentis	2	2	4	0.030	3.0=3	1	0	2	0
10	Glaucoma	Tonometry, gonioscopy and perimetry	2	2	4	0.030	3.0=3	1	2	0	0
		Primary openangle glaucoma	2	3	6	0.048	4.8=5	0	2	3	0
		Primary angleclosure glaucoma	2	2	4	0.03	3=3	1	1	1	0
		Ocular hypertension and Normal tension glaucoma	1	2	2	0.016	1.6=2	1	0	1	0
		Pseudo exfoliation and Inflammatory glaucoma	2	2	4	0.030	3.0=3	1	0	2	0
		Pigment dispersion glaucoma and Iridocorneal endothelial syndrome	1	2	2	0.016	1.60=2	1	0	1	0
		Neovascular and lens-related glaucoma	2	2	4	0.03	3.0=3	2	0	1	0
		Traumatic glaucoma and Glaucoma in intraocular tumors	1	2	2	0.016	1.6=2	1	0	1	0
		Primary congenital glaucoma	2	1	2	0.016	1.6=2	1	0	1	0



- Level of questions (according to Bloom's taxonomy)
- C2 26
- C3 74
  - **O** Discipline Ophthalmology
  - O Level of exam (FTA)
  - O Paper (II)
  - **O** No. of items (100)

S. No	Торіс	Presentation	Impact	Frequency	I × F	Weight	No of items	Diagnosis	Investigation	Treatment	Basic knowledge
1	Uveitis	Anterior uveitis	2	3	6	0.042	4.2=4	0	2	2	0
		Intermediate uveitis	2	1	2	0.014	1.4=1	0	0	1	0
		Uveitis in spondyloarthropat hies and juvenile arthritis	1	1	1	0.007	0.7=1	0	0	1	0
		Sarcoidosis and Bechet syndrome	1	1	1	0.007	0.7=1	0	0	1	0
		Vogt-koyanagi Harada syndrome	1	1	1	0.007	0.7=1	0	0	1	0
		Bacterial and viral uveitis	1	1	1	0.007	0.7=1	0	0	1	0
		Fungal and parasitic uveitis, Uveitis in bowel and renal disease	1	1	1	0.007	0.7=1	1	0	0	0
2	Ocular tumors	Benign and malignant conjunctival tumors	1	2	2	0.014	1.4=1	0	0	1	0
		Iris cysts and tumors, Ciliary body tumors	1	1	1	0.007	0.7=1	1	0	0	0

	Tumors of the retina and choroid	1	1	1	0.007	0.7=1	0	0	1	0
	Primary intraocular lymphoma	1	1	1	0.007	0.7=1	0	0	1	0



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							MEDICP				-,
		Paraneoplastic syndromes	1	1	1	0.007	0.7=1	1	0	0	0
3	Retinal vascular	Diabetic retinopathy	3	2	6	0.042	4.2=4	0	2	2	0
	disease	Retinal venous occlusive disease	3	2	6	0.042	4.2=4	0	2	2	0
		Retinal arterial occlusive disease	2	1	2	0.014	1.4=1	0	1	0	0
		Ocular ischemic syndrome and Sickle cell retinopathy	1	1	1	0.007	0.7=1	1	0	0	0
		Hypertensive disease	2	3	6	0.042	4.2=4	0	2	2	0
		Retinopathy of prematurity	3	1	3	0.021	2.1=2	1	0	1	0
		Retinal artery macro aneurysm and Primary retinal telangiectasis	1	1	1	0.007	0.7=1	0	0	1	0
		Eales disease	1	1	1	0.007	0.7=1	0	0	1	0
		Radiation retinopathy and Purtscher retinopathy	1	1	1	0.007	0.7=1	1	0	0	0

		Valsalva retinopathy	1	1	1	0.007	0.7=1	0	0	1	0
4	Acquired macular	Imaging in macular disease	3	2	6	0.042	4.2=4	0	2	2	0
	disorders	Age related macular degeneration	2	2	4	0.028	2.8=3	0	1	2	0
		Age-related macular hole	2	1	2	0.014	1.4=1	0	1	0	0



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							MEDICAT				
		Central serous retinopathy and Cystoid macular edema	1	2	2	0.014	1.4=1	0	0	1	0
		Macular epiretinal membrane and Polypoidal choroidal vasculopathy	1	1	1	0.007	0.7=1	0	0	1	0
		Degenerative myopia	1	2	2	0.014	1.4=1	0	0	1	0
		Angioid streaks and Vitreomacular traction syndrome	1	1	1	0.007	0.7=1	0	0	1	0
		Idiopathic choroidal neovascularization	1	1	1	0.007	0.7=1	0	0	1	0
		Solar retinopathy	1	1	1	0.007	0.7=1	0	0	1	0
5	Hereditary Fundus dystrophies	Macular dystrophies and Choroidal	1	2	2	0.014	1.4=1	1	0	0	0
		dystrophies									

		Generalized photoreceptor dystrophies	1	2	2	0.014	1.4=1	0	1	0	0
		Albinism and Cherry red spot at macula	1	1	1	0.007	0.7=1	1	0	0	0
6	Retinal detachment	Rhegmatogenous retinal detachment	3	2	6	0.042	4.2=4	0	1	3	0
		Tractional retinal detachment	2	2	4	0.028	2.8=3	1	0	2	0
		Exudative retinal detachment	1	1	1	0.007	0.7=1	0	0	1	0
		Pars plana vitrectomy	3	2	6	0.042	4.2=4	0	1	3	0
7	Strabismus	Amblyopia	1	3	3	0.021	2.1=2	0	1	1	0
		Heterophoria and vergence abnormalities	1	2	2	0.014	1.4=1	0	0	1	0
		Esotropia	1	2	2	0.014	1.4=1	0	0	1	0
		Exotropia	1	1	1	0.007	0.7=1	0	0	1	0



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							MEDICAL	amaipina	weutcal U	interesty	
		Congenital cranial dysinnervation disorders	1	1	1	0.007	0.7=1	1	0	0	0
		Monocular elevation deficiency and Brown syndrome	1	1	1	0.007	0.7=1	0	0	1	0
		Alphabet patterns	1	1	1	0.007	0.7=1	0	0	1	0
8	Vitreous opactities	Vitreous hemorrhage	2	2	4	0.028	2.8=3	1	0	2	0
9		Neuroimaging	2	2	4	0.028	2.8=3	0	2	1	0
	Neuroophthalmol ogy	Optic nerve, pupil, chiasma, retro chiasmal pathways	2	3	6	0.042	4.2=4	1	1	2	0
		Ocular motor nerves	2	1	2	0.014	1.4=1	1	0	0	0
		Supranuclear disorders of ocular motility	1	1	1	0.007	0.7=1	1	0	0	0
		Nystagmus and Facial spasm	1	1	1	0.007	0.7=1	0	0	1	0
		Ocular myopathies and Neurofibromatosis	1	2	2	0.014	1.4=1	0	0	1	0
		Migraine and Neuralgias	1	1	1	0.007	0.7=1	0	0	1	0
10	Ocular side	Cornea	1	1	1	0.007	0.7=1	1	0	0	0
	effects of systemic	Ciliary effusion and lens	1	1	1	0.007	0.7=1	1	0	0	0
	medications	Optic nerve	1	1	1	0.007	0.7=1	1	0	0	0
11	Ocular trauma	Eyelid trauma	1	2	2	0.014	1.4=1	0	0	1	0
		Orbital trauma	2	2	4	0.028	2.8=3	0	1	2	0
		Trauma to globe	2	2	4	0.028	2.8=3	0	1	2	0
		Chemical injuries	2	2	4	0.028	2.8=3	2	0	1	0

Level of questions (according to Bloom's taxonomy)

C2 25

C3 75



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## Table of Specification for OSCE FTA

Serial #	Station Type	Clinical Competence	Task
1	Observed	Interpretation of investigation and	MRI interpretation,
	(Interactive)	clinical reasoning	diagnosis, management and
			complications
2	Observed	History taking and clinical	Patient of sudden /gradual
	(Interactive)	reasoning	painful/painless loss of
			vision (Real or SP)
3	Observed	Communication skills	Counselling of a patient of
	(Interactive)		DR, Glaucoma,
			retinoblastoma, retinitis
			pigmentosa, ROP (Real or
			SP)
4	Observed	Performance of task	Examination of patient of
	(Interactive)		squint, pupils, EOM, Visual
			fields, ptosis, proptosis
5	Observed	Interpretation of investigation and	OCT interpretation,
	(Interactive)	problem solving	diagnosis and treatment
6	Observed	Interpretation of investigation and	Fundus picture
	(Interactive)	problem solving	interpretation, diagnosis and
			treatment
7	Observed	Interpretation of investigations and	CT-scan interpretation and
	(Interactive)	critical thinking	treatment options of
			diagnosis
8	Observed	Critical thinking and problem	Hess chart interpretation
	(Interactive)	solving based on investigation	and critical thinking
9	Observed	Video interpretation and critical	Glaucoma, Cataract, RD
	(Interactive)	thinking	videos interpretation and
			treatment options of disease
10	Observed	Interpretation of investigation and	Corneal topography
	(Interactive)	clinical reasoning	interpretation and
			justifications for
			management
11	Observed	Interpretation of investigation and	FFA interpretation,
	(Interactive)	problem solving	diagnosis and treatment
12	Observed	Interpretation of investigation and	B-scan interpretation,
	(Interactive)	problem solving	diagnosis and treatment
13	Observed	Interpretation of investigation and	Visual fields interpretation,
	(Interactive)	critical thinking	diagnosis, management and
			complications
14	Observed (Interactive)		Slit lamp examination
		Performance of task and problem	techniques, Tonometry,
		solving	Indirect ophthalmoscopy,
			Gonioscopy
15	Observed	Performance of task	Ophthalmic procedures



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(Interactive)		Intravitreal injection, Lid
		stitching, corneal laceration
		repair, fortified antibiotic
		prep, probing and syringing

#### **Research Assessment**

#### Submission of Synopsis and Thesis

- 1. The candidates shall prepare their synopsis as per guidelines provided by the Advanced Studies & Research Board, available on RMU website.
- Synopsis of research project should be submitted and approved by the end of the 1<sup>st</sup> year of MS program.
- 3. The minimum duration between approval of synopsis and submission of thesis shall be one year, but the thesis cannot be submitted later than 8 years of enrolment.
- 4. Thesis shall be submitted by the candidate duly recommended by the Supervisor.
- 5. The research thesis must be compiled and bound in accordance with the Thesis Format Guidelines approved by the University and available on website.
- 6. The research thesis will be submitted along with the fee prescribed by the University.

#### **Thesis Defense**

- All candidates admitted in MS course shall appear in thesis evaluation component of the MTA after completion of 4<sup>th</sup> years of their training course.
- 2. Only those candidates shall be eligible for thesis evaluation who have passed Midterm Examination and Oral & Practical/ Clinical component of Exit Examination.
- 3. The examination shall include thesis evaluation with defense.
- 4. The Vice Chancellor shall appoint three external examiners for thesis evaluation, preferably from other universities and from abroad, out of the panel of examiners approved by the Advanced Studies & Research Board. The examiners shall be appointed from respective specialty.
- 5. The thesis shall be sent to the external examiners for evaluation, well in time before the date of defense examination and should be approved by all the examiners.
- 6. After the approval of thesis by the evaluators, the thesis defense examination shall be held within the University on such date as may be notified by the Controller of Examinations. The Controller of Examinations shall make appropriate arrangements for the conduct of thesis defense examination in consultation with the supervisor, who will co-ordinate the defense examination.
- 7. The thesis defense examination shall be conducted by two External Examiners who shall submit a report on the suitability of the candidate for the award of degree. The supervisor shall act as coordinator.



## **SECTION X: ENTRUSTABLE PROFESSIONAL ACTIVITY** (EPA)





#### Overview

Entrustable Professional Activities (EPAs) for a four-year ophthalmology residency program are essential in defining the specific tasks residents should be able to perform independently by the end of their training. These EPAs are aligned with clinical core competencies and are designed to ensure that residents progressively develop their skills and knowledge throughout their residency.

#### Levels of EPA

1)Be present and observe or Assist

2)Direct pro-active Supervision: The supervisor is physically present with the resident and the patient.

3) Indirect re-active Supervision is broken down into two levels: Direct Supervision Immediately Available: The supervisor is physically within the hospital or other site of patient care and is immediately available to provide direct supervision. Direct Supervision not readily Available: The supervisor is not physically present within the hospital or other site of patient care, but is immediately available by means of telephonic and/or electronic modalities, and is available to provide direct supervision.

4) Can supervise other junior residents

				TRUTH	· ·			
	RY-1		RY-2		RY-3		RY-4	
EPA (Clinical Competencies)	EPA Level	No	EPA Level	No	EPA Level	No	EPA Level	No
Conducting a Comprehensive Patient Evaluation and History Taking for Common Ophthalmic Presentations	2	20	2-3	20	4	20	4	20
Performing Essential Ophthalmic Examinations.	2	20	2-3	20	3-4	20	4	20
Recognizing Life- and Vision- Threatening Findings in Basic Ophthalmic Diagnostic Investigations	2	20	2-3	20	3-4	20	4	20
Developing an initial management plan for patients with an acute ophthalmic condition or presentation	2	20	2-3	20	3-4	20	4	20
Performing basic Ophthalmic surgical Procedures	2	20	2-3	20	3-4	20	4	20
Effectively Communicating Clinical Findings and Management Plans to Patients	2	20	2-3	20	3-4	20	4	20
Performing Cataract Extraction Surgeries	2	20	2-3	10	3-4	20	4	20
Performing anterior segment laser procedures	2	10	2-3	10	3-4	20	4	20
Performing Retinal Procedures, Including Laser Treatments and Injections	2	20	2-3	20	3-4	20	4	20
Performing orbit, oculoplastic and strabismus surgical procedures	2	10	2-3	10	3	10	3-4	10
Managing Ocular Trauma and Its Complications	2	20	2-3	20	3-4	20	4	20

## EPA of MS Ophthalmology Training Program



## **SECTION XI: LOGBOOK**





#### **Introduction to Logbook**

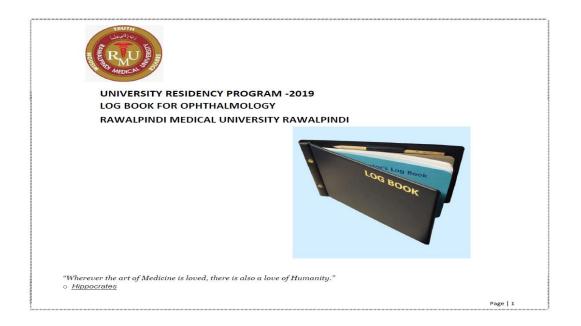
It is a structured book in which certain types of educational activities and patient related information is recorded, usually by hand. Logbooks are used all over the world from undergraduate to postgraduate training, in human, veterinary and dental medicine, nursing schools and pharmacy, either in paper or electronic format.

Logbooks provide a clear setting of learning objectives and give trainees and clinical teachers a quick overview of the requirements of training and an idea of the learning progress. Logbooks are especially useful if different sites are involved in the training to set a (minimum) standard of training. Logbooks assist supervisors and trainees to see at one glance which learning objectives have not yet been accomplished and to set a learning plan. The analysis of logbooks can reveal weak points of training and can evaluate whether trainees have fulfilled the minimum requirements of training.

Logbooks facilitate communication between the trainee and clinical teacher. Logbooks help to structure and standardize learning in clinical settings. In contrast to portfolios, which focus on students' documentation and self-reflection of their learning activities, logbooks set clear learning objectives and help to structure the learning process in clinical settings and to ease communication between trainee and clinical teacher. To implement logbooks in clinical training successfully, logbooks have to be an integrated part of the curriculum and the daily routine on the ward. Continuous measures of quality management are necessary.

#### Reference

Brauns KS, Narciss E, Schneyinck C, Böhme K, Brüstle P, Holzmann UM, etal. Twelve tips for successfully implementing logbooks in clinical training. Med Teach. 2016 Jun 2; 38(6): 564–569.





## Logbook Entry Map

Procedures	First Year	Second Year	Third Year	Fourth Year
Visual Acuity	10	10	05	05
Refraction	10	10	05	05
<b>Optic Nerve function test</b>	10	10	05	05
Slit Lamp Examination	10	10	15	15
Tonometry	10	10	15	15
Visual Fields Examination	10	10	00	00
Cover Uncover test	10	05	05	00
Direct Ophthalmoscopy	05	05	05	05
Indirect Fundoscopy	04	05	05	05
Biometry	10	05	05	00
B-scan ultrasonography	05	05	05	05
Pre-op management of eye	05	05	00	00
surgeries				
Post op management of eye surgeries	05	05	00	00
Facial block	05	05	00	00
Retrobulbar block	02	02	03	03
Gonioscopy	02	00	05	03
Corneal ROS	02	02	03	03
Orthoptic Assessment	02	00	04	03
Visual Field Interpretation	02	00	03	05
OCT Interpretation	02	05	06	05
FFA interpretation	02	00	03	05
Prosis Examination	02	05	00	00
Proptosis Examination	01	02	03	00
Intravitreal antibiotics	02	02	02	03
Chalazion I&D	02	05	02	00
Pterygium excision	02	03	05	00
Nd: YAG Laser	00	00	00	05
capsulotomy	00	00	00	05
Nd: YAG Laser iridotomy	00	00	00	03
Argon Laser	00	00	00	03
photocoagulation	00	00	02	05
Intravitreal anti VEGF	02	02	03	03
ECCE	03	03	06	07
Phacoemulsification	00	04	06	07
Trabeculectomy	00	00	02	03
Squint surgery	00	00	02	03
Retinal detachment surgery	00	00	02	03
Oculoplastic procedures	00	00	02	03
Trauma repair	00	00	03	02
Ptosis surgery	02	02	03	03
Orbital surgery	00	01	02	02
Orbital surgery Orbital tumors	00	01	02	02
	00	01	01	01
Keratoplasty Dermoid cyst	00	01	02	02
Scleral buckling	01	02	03	00
MPTCPC	00	00	02	03
Probing & Syringing	00	02	02	01
	03	02	00	00
Dacryocystorhinostomy	02	02	01	
Pars Plana Vitrectomy				03
Total	150	150	150	150



## **SECTION XII: PORTFOLIO**

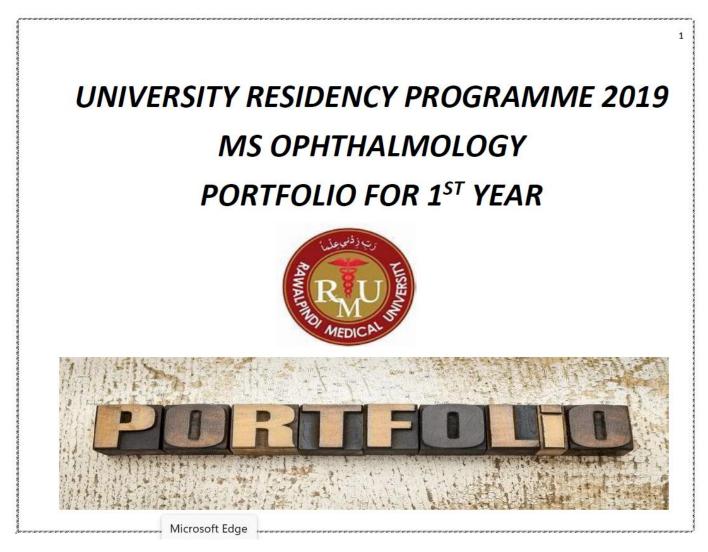




#### **Introduction to Portfolio**

#### What is a portfolio?

A collection of a learner's various documents and assessments throughout residency that reflect their professional development over time. May include referral letters and procedure logs (Rider et al., 2007). Portfolios also frequently include self-assessments, learning plans, and reflective essays (Epstein, 2007).



#### Portfolio Components and Expectations by Training Year

#### Year 1 Portfolio: Foundational Year

Focus: Developing foundational medical knowledge, basic clinical skills, and understanding professional behavior.

#### **Key Components**:

#### Academic Activities:

Attendance in didactic lectures, journal clubs, and seminars (≥80% attendance required).

Performance in LMS-based assessments and mid-module/end-module examinations.

Case-based discussions and participation in SDLs.



#### **Clinical Skills:**

Log of cases observed and assisted (e.g., cataract surgeries, slit-lamp exams).

Initial DOPS evaluations for basic procedures (e.g., retinoscopy, IOP measurement).

#### **Research Activities:**

Participation in research methodology workshops.

Selection of a thesis topic and development of a research proposal.

#### **Professionalism and Ethics:**

Documentation of adherence to ethical standards and professional behavior.

Feedback from supervisors and peers on interpersonal skills and teamwork.

#### **Reflection and Feedback:**

Resident's reflections on learning experiences and goals for improvement.

#### Year 2 Portfolio: Intermediate Skills Development

Focus: Gaining competency in intermediate diagnostic and therapeutic procedures and starting independent patient care.

#### **Key Components:**

#### Academic Activities:

Performance in block exams, OSCEs, and viva voce assessments.

Continued participation in journal clubs and case presentations.

#### **Clinical Skills:**

Log of independently performed diagnostic procedures (e.g., slit-lamp exams, gonioscopy).

Progress in minor surgical skills (e.g., chalazion excision, pterygium surgery).

DOPS and mini-CEX evaluations reflecting growth in patient care competencies.

#### **Research Activities:**

Submission of a literature review related to the thesis topic.

Regular updates on research progress and data collection.

#### **Professionalism and Ethics:**

Multi-source feedback (MSF) evaluations from faculty, peers, and patients.

Documentation of ethical challenges encountered and how they were managed.

#### **Reflection and Feedback:**

Self-assessment of skills and learning, with updated learning objectives.



#### Year 3 Portfolio: Advanced Clinical Competence

Focus: Mastering advanced clinical and surgical skills, managing complex cases, and conducting research independently.

#### **Key Components:**

#### **Academic Activities:**

Continued high performance in assessments, including SEQs and mock exams.

Leadership in journal clubs, seminars, and interdepartmental conferences.

#### **Clinical Skills:**

Log of advanced diagnostic and surgical procedures performed independently (e.g., phacoemulsification, trabeculectomy).

Evaluations through ICO-OCEX, ICO-OSCAR, and mini-CEX tools.

Detailed documentation of cases managed independently.

#### **Research Activities:**

Thesis completion, submission, and preparation for defense.

Presentation of findings at conferences or surgical audits.

#### **Professionalism and Ethics:**

Multi-source feedback from patients and staff assessing communication and professionalism.

Documentation of contributions to team-based practice and patient safety initiatives.

#### **Reflection and Feedback:**

Critical analysis of strengths and areas needing improvement.

#### Year 4 Portfolio: Competency and Leadership

Focus: Transitioning to independent practice, teaching juniors, and finalizing research contributions.

#### **Key Components:**

#### Academic Activities:

Demonstrating mastery in final assessments and case presentations.

Active mentoring of junior residents during clinical and academic sessions.

#### **Clinical Skills:**

Log of all advanced surgeries performed as a primary surgeon.

Final WPBA evaluations, focusing on readiness for independent practice.

#### **Research Activities:**

Successful thesis defense and publication of research findings in peer-reviewed journals.

Participation in surgical audits and contribution to program improvements.



#### **Professionalism and Ethics:**

Leadership roles in clinical teams and patient care.

Multi-source feedback evaluating readiness for independent, ethical practice.

#### **Reflection and Feedback:**

Comprehensive self-reflection on the residency journey, with feedback from faculty.

#### Assessment and Feedback

#### **Portfolio Grading**:

Each portfolio is graded on a structured rubric covering:

Completeness of documentation (20%)

Quality of work in each competency area (40%)

Feedback and reflection quality (20%)

Adherence to timelines and participation (20%)

#### **Feedback Sessions:**

Residents receive individualized feedback after each review to guide future improvements and ensure alignment with program milestones.

#### **Promotion Criteria**:

Satisfactory portfolio review is required for promotion to the next year of training. Deficiencies must be addressed in a remediation plan.

#### What to be included in a portfolio?

Resident may include the following components in his or her portfolio:

- 1. Curriculum Vitae (CV)
- 2. Personal Publications
- 3. Research abstracts presented at professional conferences
- 4. Presentations at teaching units/departmental meetings and teaching sessions
- 5. Patient (case) presentations
- 6. Log of clinical procedures
- 7. Copies of written feedback received (direct observations, field notes, daily evaluations)
- 8. Quality improvement project plan and report of results
- 9. Summaries of ethical dilemmas (and how they were handled)
- 10. Chart notes of particular interest
- 11. Photographs and logs of medical procedures performed
- 12. Consult/referral letters of particular interest
- 13. Monthly faculty evaluations
- 14. 360-degree evaluations
- 15. Copies of written instructions for patients and families
- 16. Case presentations, lectures, logs of medical students mentored



## **SECTION XIII: REFERENCES**







#### **Teaching Methods**

Kolb, D. Experiential Learning. Englewood Cliffs, NJ: Prentice Hall. 1984

Maudsley G. Do we all mean the same thing by "PBL"? Academic Medicine 1999; 74:178-85

Hill W. Learning Thru Discussion 2nd edition. London: Sage Publications. 1977.

Cook D. Web-based learning: pros, cons and controversies. Clinical Medicine 2007; 7(1):37-42.

Greenhalgh T. Computer assisted learning in undergraduate medical education. BMJ 2001; 322:40-4.

Chumley-Jones HS *et al* Web-based learning: Sound educational method or Hype? A review of the evaluationliterature. Academic Medicine 2002;77(10):S86-S93.

Schon D. Educating the reflective practitioner. San Francisco: Jossey Bass. 1984

Lockyer J *et al* Knowledge translation: the role and practice of reflection. Journal of Continuing Education. 2004;24:50-56

#### **Assessment methods**

Van der Vleuten, CPM and Swanson, D. Assessment of clinical skills with standardized patients: State of theart. *Teach Learn Med.* 1990; 2: 58-76.

Haladyna TM. *Developing and validating multiple-choice test items*. Hillsdale, New Jersey: L. Erlbaum Associates.1994.



Case SM, Swanson DB. *Constructing written test questions for the basic and* Philadelphia, PA:National Board of Medical Examiners, 1996 (<u>www.nbme.org</u>) clinical sciences.

Case SM, Swanson DB. *Constructing written test questions for the basic and clinical sciences*. Philadelphia, PA:National Board of Medical Examiners, 1996 (<u>www.nbme.org</u>)

Center for Creative Leadership, Greensboro, North Carolina (<u>http://www.ccl.org</u>).

Challis M. AMEE medical education guide no. 11 (revised): Portfolio-based learning and assessment in medicaleducation. *Med Teach*. 1999; 21: 370-86.

Gray, J. Global rating scales in residency education. Acad Med. 1996; 71: S55-63.

Haladyna TM. *Developing and validating multiple-choice test items*. Hillsdale, New Jersey: L. Erlbaum Associates.1994.

Kaplan SH, Ware JE. The patient's role in health care and quality assessment. In: Goldfield N and Nash D (eds). *Providing quality care (2<sup>nd</sup> ed): Future Challenge*. Ann Arbor, MI: Health Administration Press, 1995: 25-52.

Matthews DA, Feinstein AR. A new instrument for patients' ratings of physician performance in the hospital setting. *J Gen Intern Med.* 1989:4:14-22.

Mancall EL, Bashook PG. (eds.) *Assessing clinical reasoning: the oral examination and alternative methods*. Evanston, Illinois: American Board of Medical Specialties, 1995.

Munger, BS. Oral examinations. In Mancall EL, Bashook PG. (editors) *Recertification: new evaluation methods and strategies*. Evanston, Illinois: American Board of Medical Specialties, 1995: 39-42.

Noel G, Herbers JE, Caplow M et al. How well do Internal Medicine faculty members evaluate the clinical skills ofresidents? *Ann Int Med.* 1992; 117: 757-65.

Norman, Geoffrey. *Evaluation Methods: A resource handbook*. Hamilton, Ontario, Canada: Program for EducationalDevelopment, McMaster University, 1995: 71-77.Tekian A, McGuire CH, et al (eds.) *Innovative simulations for assessing professional competence*. Chicago, Illinois:University of Illinois at Chicago, Dept. Med. Educ. 1999



Tugwell P, Dok, C. Medical record review. In: Neufeld V and Norman G (ed). *Assessing clinical competence*. NewYork: Springer Publishing Company, 1985: 142-82.

Van der Vleuten, CPM and Swanson, D. Assessment of clinical skills with standardized patients: State of the art. *Teach Learn Med.* 1990; 2: 58-76.

Watts J, Feldman WB. Assessment of technical skills. In: Neufeld V and Norman G (ed). *Assessing clinical competence*. New York: Springer Publishing Company, 1985, 259-74.

Winckel CP, Reznick RK, Cohen R, Taylor B. Reliability and construct validity of a structured technical skillsassessment form. *Am J Surg.* 1994; 167: 423-27.

#### Milestones

https://www.acgme.org/Portals/0/PDFs/Milestones/Ophthalmology Milestones.pdf

http://education.med.ufl.edu/files/2010/10/Ophthalmology Milestones.pdf

http://www.upstate.edu/medresidency/current/competencies.php



## **SECTION XIV: APPENDICES**





#### **List of Appendices**

Workplace Based Assessments-Multi source feedback profoma- 360º evaluation Appendix "A"

Proforma for feedback by Nurse for core competencies of the resident "Appendix B"

Proforma for patient Medication Record "Appendix C"

Workplace Based Assessments- guidelines for assessment of Generic & specialty specific Competencies - ----- Appendix "D"

Supervisor's Annual Review Report Appendix "E"

Supervisors evaluation Proforma for continuous internal assessments Appendix "F"

Evaluation of resident by the faculty Appendix "G"

Evaluation of faculty by the resident Appendix "H"

- Evaluation of program by the faculty Appendix "I"
- Evaluation of program by the resident Appendix "J"
- Guidelines for program evaluation-----Appendix "K"
- Evaluation of Project Director by the residents Appendix "L"
- Registration and Enrollment-----Appendix 'M'





## **MENTOR / SUPERVISOR EVALUATION OF TRAINEE**

Resident's Name:	
Evaluator's Name(s):	
Hospital Name:	
Date of Evaluation:	
Traditional Track (10% Clinic)	Primary Care Track (20% Clinic)

1	Unsatisfactory
2	Below Average
3	Average
4	Good
5	Superior

Please circle the appropriate number for each item using the scale above.

	Patient Care		s	cal	е	
1.	Demonstrates sound clinical judgment	1	2	3	4	5
2.	Presents patient information case concisely without significant omissions or digressions	1	2	3	4	5
3.	Able to integrate the history and physical findings with the clinical data and identify all of the patient's major problems using a logical thought process	1	2	3	4	5
4.	Develops a logical sequence in planning for diagnostic tests and procedures and Formulates an appropriate treatment plan to deal with the patient's major problems	1	2	3	4	5
5.	Able to perform commonly used office procedures	1	2	3	4	5
6.	Follows age appropriate preventative medicine guidelines in patient care	1	2	3	4	5
	Medical Knowledge		s	ical	е	
1.	Uses current terminology	1	2	3	4	5
2.	Understands the meaning of the patient's abnormal findings	1	2	3	4	5
3.	Utilizes the appropriate techniques of physical examination	1	2	3	4	5
4.	Develops a pertinent and appropriate differential diagnosis for each patient	1	2	3	4	5
5.	Demonstrates a solid base of knowledge of ambulatory medicine	1	2	3	4	5
6.	Can discuss and apply the applicable basic and clinically supportive sciences	1	2	3	4	5
	Professionalism		s	cal	е	
1.	Demonstrates consideration for the patient's comfort and modesty	1	2	3	4	5
2.	Arrives to clinic on time and follows clinic policies and procedures	1	2	3	4	5
3.	Works effectively with clinic staff and other health professionals	1	2	3	4	5
4.	Able to gain the patient's cooperation and respect	1	2	3	4	5
5.	Demonstrates compassion and empathy for the patient	1	2	3	4	5
6.	Demonstrates sensitivity to patient's culture, age, gender, and disabilities	1	2	3	4	5
7.	Discusses end-of-life issues (DPOA, advanced directives, etc.) when appropriate	1	2	3	4	5





# 2

## Patient Medical Record / Chart Evaluation Proforma

Name of Resident

Location of Care or Interaction (OPD/Ward/Emergency/Endoscopy Department)

S#		Poor	Fair	Good	V. Good	Excellent
1.	Basic Data on Front Page Recorded	0	0	0	0	0
2	Presenting Complaints written in chronological order	0	0	0	0	0
3.	Presenting Complaints Evaluation Done	0	0	0	0	0
4.	Systemic review Documented	0	0	0	0	0
5.	All Components of History Documented	0	0	0	0	0
6.	Complete General Physical Examination done	0	0	0	0	0
7.	Examination of all systems documented	0	0	0	0	0
8.	Differential Diagnosis framed	0	0	0	0	0
9.	Relevant and required investigations documented	0	0	0	0	0
10.	Management Plan framed	0	0	0	0	0
11.	Notes are properly written and eligible	0	0	0	0	0
12.	Progress notes written in organized manner	0	0	0	0	0
13.	Daily progress is written	0	0	0	0	0
14.	Chart is organized no loose paper	0	0	0	0	0
15.	Investigations properly pasted	0	0	0	0	0
16.	Abnormal findings in investigations encircled.	0	0	0	0	0
17.	Procedures done on patient documented properly	0	0	0	0	0
18.	Medicine written in capital letter	0	0	0	0	0
19.	I/v fluids orders are proper with rate of infusion mentioned	0	0	0	0	0
20.	All columns of chart complete	0	0	0	0	0







-

Preview Form

#### **RESIDENT EVALUATION BY NURSE / STAFF**

Please take a few minutes to complete this evaluation form. All information is confidential and will be used constructively. You need not answer all the questions

Name of Resident\*

#### Location of care or interaction: (OPD/Ward/Emergency/Endoscopy Department)

Your position (Nurse, Ward Servant, Endoscopy Attendant) CH DOOD

		Poor	Fair	Good	V Good	Excellent	Insufficient Contact
1.	Resident is Honest and Trustworthy	0	0	0	0	0	0
2.	Resident treats patients and families with courtesy, compassion and respect	0	0	0	0	0	0
3.	Resident treats me and other member of the team with courtesy and respect	0	0	0	0	0	0
4.	Resident shows regard for my opinions	0	0	0	0	0	0
5.	Resident maintains a professional manner and appearance	0	0	0	0	0	0
INTE	RPERSONAL AND COMMUNICATIONS SKILLS	2	2	2 77			
6.	Resident communicates well with patients, families, and members of the healthcare team	0	0	0	0	0	0
7.	Resident provides legible and timely documentation	0	0	0	0	0	0
8.	Resident respect differences in religion, culture age, gender sexual orientation and disability	0	0	0	0	0	0
SYST	EMS BASED PRACTICE	\$ <u></u>	( )	9 - 3	(	2 <u>0</u>	9.
9.	Resident works effectively with nurses and other professionals to improve patient care.	0	0	0	0	0	0
PATI	ENT CARE			10 - 74	2	3	
10.	Resident respects patient preferences	0	0	0	0	0	0
11.	Resident is reasonable accessible to patients	0	0	0	0	0	0
12.	Resident take care of patient comfort and dignity during procedures.	0	0	0	0	0	0
PRA	TICE BASED LEARNING AND IMPROVEMENT	1	1	1		1	
13.	Resident facilitates the learning of students and other professionals	0	0	0	0	0	0
CON	IMENTS	<u>k</u> .		1: a.			5
14.	Please describe any praises or concerns or information about specific incidents	0	0	0	0	0	0

Poor: 0, Fair: 1, Good: 2, V. Good: 3, Excellent: 4







## **Patient Evaluation of Trainee**

Trainee Name:	
Date of Evaluation:	

1	Strongly Disagree
2	Disagree
3	Neutral
4	Agree
5	Strongly Agree

Please circle the appropriate number for each item using this scale. Please provide any relevant comments on the back of this form.

	This Trainee:	Scale							
1.	Introduces him/herself and greets me in a way that makes me feel comfortable.	1	2	3	4	5			
2.	Manages his/her time well and is respectful of my time.	1	2	3	4	5			
3.	Is truthful, upfront, and does not keep things from me that I believe I should know. ڈاکٹر صاحب نے میر سے مرض کی صورتحال یوری سچائی سے بیان کی۔	1	2	3	4	5			
4.	Talks to me in a way that I can understand, while also being respectful.	1	2	3	4	5			
5.	Understands how my health affects me, based on his/her understanding of the details of my life. د اکٹر صاحب نے میر ے علاج میں میں میں محت پر دانی زندگی کو مذظر رکھا۔	1	2	3	4	5			
6.	Takes time to explain my treatment options, including benefits and risks.	1	2	3	4	5			

Total Score \_\_\_\_/30







## **Resident/Fellow Evaluation of Faculty Teaching**

Evaluator:

Evaluation of: \_\_\_\_\_

Date:\_\_\_\_\_

Evaluation information entered here will be anonymous and made available only in aggregated form.

S#		Strongly Disagree	Disagree Moderately	Disagree Slightly	Agree Slightly	Agree Moderately	Strongly Agree
	*	PATI	ENT CARE				
1.	Teaches current scientific evidence for daily patient management*						
2.	Explains rationale behind clinical judgements/decisions*						
3.	Teaches clear diagnostic algorithms*						
4.	Teaches clear treatment algorithms*						
	PATIENT CARE	- OPERAT	IVE AND PE	ROCEDUR	AL SKILI	LS	
5.	Teaches operative/procedural skills during cases*						
6.	Allows learners to perform operative/procedural skills when appropriate*						
1	de la companya de la	MEDICAL	. KNOWLEE	GE	16	<i>8</i> .	5
7.	Teaches relevant pathophysiology needed to evaluate patient medical conditions*						
8.	Teaches how/when to use-order- perform procedures/tests*						
9.	Teaching content adds significantly to my medical knowledge						
10.	Teaches the use of literature / evidence based medicine to support clinical decisions/teaching points*						







#### FINAL Evaluation Scoring Sheet

Name of Reside			Na	me of Su	pervi	sor			Y	ear of T	raining	9		
Date	Faculty #1 (165)	Faculty #2 (165)	Faculty #3 (165)	Average Score		Duration Specialt Hospital	y	sessm	ent					
Medical Patient Care (30)					/30		Unit							
Medical Knowledge	(30)				/30									
Professionalism	(35)				/35							1.000		
Interpersonal and Communication Skills	(20)				_/20	(30)	(30)	(30)	ord (80)	ord (80)	ord (80)	(56)	(56)	(56)
System Based Practice	(35)				/35	1#1	t#2	t#3	al Rec na #1	al Reci na #2	il Reci na #3	Ξ	2	2
Practice Based Learning and Improvement	(15)				/15	Patient #1	Patient # 2	Patient # 3	Medical Record Performa #1 (	Medical Record Performa #2 (	Medical Record Performa #3	Staff #	Staff #2	Staff #3
Overall Rating														
Average:					/165		_	/30		53	/80		_	/56
												Gran	d To	tal
											-		!:	331

	NDI MEDICAL UNIVER	SITY			
Logbook	complete	incomplete			
Portfolio	complete	incomplete			
Leave /absentees:			-		
Comments					
·					
Supervisor Name (1)	Superv	risor Name (2)		Head of Unit	
Sign & Stamp	Sign & Sign	Stamp		Sign & Stamp	



RU

**RAWALPINDI MEDICAL UNIVERSITY** 

7	

# **RESIDENT SELF-ASSESSMENT PROFORMA**

Resident Name

Date

ear of Training _		Hosp	ital Name				U	nit _					
NA Not Applicable		<u> </u>	<b>2</b>			3					0 4	1	_
		t Applicable I rarely demonstrates I do this Sometimes I do this mos		st of the time of the time)			I do this all the tin (>75% of time)						
1.		o acquire accurate and re an efficient, prioritized ar			NA		1	۵	2		3		4
2.	prioritized	to seek and obtain ap d data from secondary nd pharmacy)		d o	NA		1		2		3	۵	
3.	10 C	to perform accurate p appropriately targeted t s.			NA		1		2		3	٦	
4.	interview	to synthesize all availa , physical exam, and pi ch patient's central clin	reliminary lab data to		NA		1		2		3		4
5.	evidence	to develop prioritized based diagnostic and t conditions in Internal N	herapeutic plans for		NA		1		2		3		
6.		to recognize situations ent medical care, inclue s.		nt 🗆	NA		1		2		3	٦	
7.	I am able guidance.	to recognize when to	seek additional		NA		1		2		3		2
8.	I am able	to provide appropriate	e preventive care.		NA		1		2		3		
9.	disorders with mini	to manage patients w in the practice of outp mal supervision.	atient internal medicin	e	NA		1		2		3		
10.		rformed several invasiv ted them in my New In			NA		1		2		3		
11.	treat com	trate sufficient knowled mon conditions that re	quire hospitalization.		NA		1		2		3	۵	
12.	CONTRACTOR CONTRACTOR	and the indications for ation of common diagn			NA				2		3		
13.	my medic level of tr		it should be for my		NA		1		2	0	3	٦	
14.	I am able	to identify clinical que	stions as they emerge		NA		1		2		3		





# **Rawalpindi Medical University**

# 8

#### DIRECT OBSERVATION OF PROCEDURAL SKILLS (DOPS)

Please complete the questions using a cross	Please use black ink and CAPITAL LETTERS
Doctor's Name:	
PMDC Number:	

Clinical setting:	A&E	OPD In-	patient Acu	te Admission	Other			
Procedure number								
Assessors position: Consul	tant SpSR	SpR S	pecialty doctor	Nurse	Other			
Number of previous DOPS	observed by	0	1 2	3	4 5-	9 >	>9	
assessor with any trainee						٦		
Number of times procedure	0 1-4	5-9 >10	Difficul	ty of	Low	Average	High	
performed by traince:			proced				Ē	
Please grade the	Well below	Below	1 proces	Meets	Above	Well above		
following areas	expectations	Expectation	Borderline	Expectations	Expectations	expectations	U/C*	
tonowing areas		s						
	1	2	3	4	5	6		
I Demonstrate understanding of						_		
indications, relevant anatomy, technique of procedure								
2 Obtains informed consent							1	
3 Demonstrates appropriate			<u></u>	<u> </u>			1	
preparation pre-procedure								
4 Appropriate analgesia or								
preparation pre-procedure								
5 Technical ability safe sedation	<u> </u>	-0-		<u> </u>	<u> </u>	<u> </u>	+	
6 Aseptic technique     7 Seeks help where appropriate		<u> </u>	<u> </u>			<u> </u>	-9-	
Seeks help where appropriate     Post procedure management	<u>+U</u>		<u>  U</u>	·				
9 Communication skills	<u> −−</u>   −−−							
10 Consideration of		<u>                                      </u>		1	1-11-	<u>  4</u>	1-12-	
Patient/professionalism								
11 Overall ability to perform								
procedure								
	<ul> <li>U/C Please mark this if you have not observed the behaviour and therefore feel unable to comment.</li> <li>Please use this space to record areas of strength or any suggested development</li> </ul>							
Please use	this space to re	ecord areas o	f strength or	any suggested	development	112 .		
A wething around allow good 2			- Cum	gestions for deve	lanment	inc		
Anything especially good?			Sug	gestions for deve	elopment.			
						х.		
Have you had training in the use of	Have you had training in the use of this assessment tool? Face to face Have read guidelines Web/ CD-Rom							
1								
2					Time taken	for observatio	m:	
					(in minute	s)		
Assessors signature:	Date (mm/	vv)			Time taken f	or feedback		
		,,,,			//m			
					fla attantest			
Assessor's Name:								
		•)						
*if appropriate Please	<ul> <li>if appropriate Please note failure of return of all completed forms to your administrator is a probity issue</li> </ul>							
Acknowledgement: Adapted with permission of the American Board of internal Medicine								

SpSR - Specialty Senior Registrar

SnR - Specialty Registrar



# Workplace Based Assessments - Guidelines for Supervisors for Assessment of Generic & SpecialtySpecific Competency

The Candidates of all MD programs will be trained and assessed in the following five generic competencies and also specialty specificcompetencies.

#### A. Generic Competencies:

#### i. <u>Patient</u> <u>Care.</u>

- Patient Care competency will include skills of history taking, examination, diagnosis, counseling Plan care through ward teachingdepartmental conferences, morbidity and mortality meetings core curriculum lectures and training in procedures and operations.
- b. The candidate shall learn patient care through ward teaching departmental conferences, morbidity and mortality meetings, carecurriculum lectures and training in procedures and operations.
- c. The Candidate will be assessed by the supervisor during presentation of cases on clinical ward rounds, scenario based discussions onpatients management multisource feedback evaluation, Direct observation of Procedures (DOPS) and operating room assessments
- d. These methods of assessments will have equal weightage.

#### ii. Medical knowledge and Research

- a. The candidate will learn basic factual knowledge of illnesses relevant to the specialty through lectures/discussions on topics selectedfrom the syllabus, small group tutorials and bed side rounds
- b. The medical knowledge/skill will be assessed by the teacher during
- c. The candidate will be trained in designing research project, data collection data analysis and presentation of results by the supervisor.
- d. The acquisition of research skill will be assessed as per regulations governing thesis evaluation and its acceptance.

#### iii. Practice and System Based Learning

- a. This competency will be learnt from journal clubs, review of literature policies and guidelines, audit projects medical error investigation, root cause analysis and awareness of health care facilities,.
- b. The assessment methods will include case studies, personation in mobility and mortality review meetings and presentation of auditprojects if any.
- c. These methods of assessment shall have equal weight-age
- iv. Communication Skills



- a. These will be learn it from role models, supervisor and workshops.
- b. They will be assessed by direct observation of the candidate whilst interacting with the patients, relatives, colleagues and withmultisource feedback evaluation.

#### v. <u>Professionalism as per Hippocratic oath</u>

- a. This competency is learnt from supervisor acting as a role model ethical case conferences and lectures on ethical issues such asconfidentially informed consent end of life decisions, conflict of interest, harassment and use of human subjects in research.
- b. The assessment of residents will be through multisource feedback evaluation according to preforms of evaluation and its scoring method.

#### B. Specialty Specific Competences.

- i. The candidates will be trained in operative and procedural skills according to a quarterly based schedule.
- ii. The level of procedural Competency will be according to a competency table to be developed by each specialty
- iii. The following key will be used for assessing operative and procedural competencies:

#### a. Level 1 Observer status

- b. The candidate physically present and observing the supervisor and senior colleagues
- c. Level 2 Assistant

#### status

The candidate assisting procedures and operations

#### d. Level 3 Performed under

#### supervision

The candidate operating or performing aprocedure under direct supervision

#### e. Level 4 Performed

#### independently

The candidate operating or performing aprocedure without any supervision

#### vi. Procedure Based Assessments (PBA)

- a. Procedural competency will assess the skill of consent taking, preoperative preparation and planning, intraoperative general and specific tasks and postoperative management
- b. Procedure Based assessments will be carried out during teaching and training of each procedure.
- c. The assessors may be supervisors, consultant colleagues and senior residents.
- d. The standardized forms will be filled in by the assessor after direct observation.
- e. The resident's evaluation will be graded as satisfactory, deficient requiring further training and not assessed at all.
- f. Assessment report will be submitted
- g. A satisfactory score will be required to be eligible for taking final examination.



#### Supervisor's Annual Review Report.

This report will consist of the following components: -

- I. Verification and validation of Log Book of operations & procedures according to the expected number of operations and proceduresperformed (as per levels of competence) determined by relevant board of studies.
- II. A 90% attendance in academic activities is expected. The academic activities will include: Lectures, Workshops other than mandatoryworkshops, journal Clubs Morbidity & Mortality Review Meetings and Other presentations.
- III. Assessment report of presentations and lectures
- IV. Compliance Report to meet timeline for completion of research project.
- V. Compliance report on personal Development Plan.
- VI. Multisource Feedback Report, on relationship with colleagues, patients.
- VII. Supervisor will produce an annual report based on assessments as per proforma in appendix-G and submit it to the ExaminationDepartment.
- VIII. 75% score will be required to pass the Continuous Internal Assessment on annual review.





#### Appendix "L"

#### Program Evaluation Committee (PEC)

#### **Background**

The purpose of this committee is to conduct and document a formal, systematic evaluation of the program & curriculum on an annualbasis.

#### <u>Membership</u>

The chair and membership of the committee are appointed by the Program Director. The membership of the committee consists of atleast two members of the program faculty, and at least one resident/subspecialty resident.

#### Meeting Frequency

The committee meets, at a minimum, annually.

#### **Responsibilities of the PEC**

- The PEC actively participates in planning, developing, implementing and evaluating the educational activities of the program.
- The PEC reviews and makes recommendations for revision of competency-based goals and objectives.
- Addresses areas of non-compliance with the standards; and reviews the program annually using written evaluations of faculty, residents, and others.

#### **Required Documentation of PEC Activities**

The PEC provides the GMEC with a written Annual Program Evaluation (APE) in the format that is appended to this document. Thisdocument details a written plan of action to document initiatives to improve performance based on monitoring of activities described below.

The APE document provides evidence that the PEC is monitoring the following areas, at a minimum:



- 1. Resident performance
- 2. Faculty development
- 3. Graduate performance, including performance of program graduates on the certifying examination
- 4. Assessment of program quality through:

. <u>A n n u a l confidential and formal feedback</u> from residents and faculty about the program quality;

#### b. <u>Assessment of improvements needed based on</u> <u>program evaluation feedback</u> from faculty, residents, and others

- 5. Continuation of progress made on prior year's action plan
- 6. Prepare and submit a written plan of action to
  - a. document initiatives to improve performance in one of more of the areas identified,
  - b. Delineate how they will be measured and monitored
  - c. Document continuation of progress made on the prior year's action plan



#### Template for Documentation of Annual Program Evaluation and Improvement

Date of annual program evaluation meeting:

#### Attendees:

- i. Program Director:
- ii. Program Coordinator:
- iii. Associate/Assistant PD:
- iv. Faculty Members:
- v. Residents:

	Reviewe d √	Discussion, Followup, Action Plan
1. Current Program Requirements & Institutional Requirements		
2. Most recent Internal Review Summary to ensure all recommendations are addressed		
3. Review Curriculum		
<ul> <li>a. effective mechanism in place to distribute Goals &amp; Objectives (G&amp;O) to residents and faculty</li> </ul>		
b. overall program educational goals		
c. up-to-date competency-based G&O for each assignment		
d. up-to-date competency-based G&O for each level of training		
e. G&O contain delineation of resident responsibilities for		
patient care, progressive responsibility for patient		
management, and supervision of residents		
4. Evaluation System		



a. Resident formative evaluation meets or exceeds program requirement	
<ul> <li>Resident summative evaluation meets or exceeds program requirement</li> </ul>	
c. Faculty evaluation meets or exceeds program requirement	
d. program evaluation meets or exceeds program requirement.	
5. Didactic Curriculum	
a. includes recognizing the signs of fatigue and sleep deprivation	
b. the didactic curriculum meets program requirements	
c. the didactic curriculum meets residents needs	
6. Clinical Curriculum – the effectiveness of in-patient and ambulatory	
teaching experience (structure, case mix, meets	
resident's needs)	
7. Volume and variety of patients and procedures (case log data) meets requirements and residents' needs	
8. Summary of written program evaluations completed by both faculty and	
residents	
9. Resident supervision complies with Program Requirement	
10. Recruiting results	
11. Duty hour monitoring results	
12. Track all research and scholarly activities of faculty and	
residents/fellows	
13. Educational outcomes: is the program achieving its educational	
objectives? What aggregate data (residents as a group)can be used to	
show the program is achieving its objectives? Board scores, in-service	
training exam scores, graduate surveys, employer surveys, etc.	



15. Clinical outcomes – specialty-specific metrics aligned with	
dept./division QI initiatives, disease outcomes, patientsafety	
initiatives (describe resident involvement), QI projects	
(describe resident involvement)	

Note:

If deficiencies are found during this process, the program should prepare a written plan of action to document initiatives to improve performance in the areasthat have been identified. The action plan should be reviewed and approved by the teaching faculty and documented in meeting minutes.

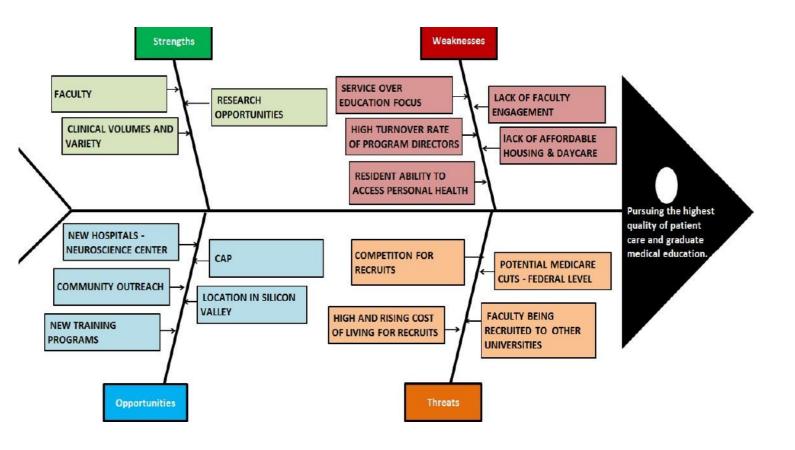


# Annual Program Evaluation (APE)

### **SWOT Analysis**

- S: Strengths
- W: Weaknesses
- **O:** Opportunities
- I T: Threats

#### SOWT Analysis (Fishbone – Ishikawa Diagram)





# Action Plan

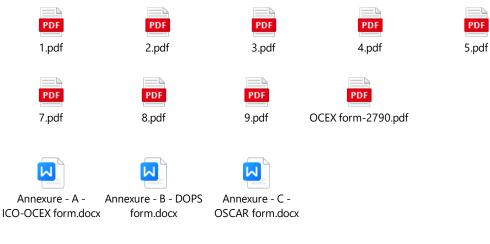
Item	Strategy	Resources	Timeline	Evaluation				
Preservation								
	Goals							
	(Strengths)							
	Elimi	nation						
		bals						
	(Weak	nesses)						
	Achie	vement						
	Go	bals						
	(Opportunities)							
Avoidance Goals								
(Threats)								



# 1. SECTION –X

**Miscellaneous attached documents** 









# **Registration and Enrolment**

ENROLMENT DETAILS	
Program of Admission	
Session	
Registration / Training Number	
Name of Candidate	-
Father's Name	_
Date of Birth / / CNIC No	
Present Address	
Permanent Address	
E-mail Address	
Cell Phone	



Date of Start of Training

Date of Completion of Training

Name of Supervisor

Designation of Supervisor

Qualification of Supervisor

Title of department / Unit



#### **Faculty Contributors**

SR. NO.	NAME & DESIGNATION	SIGNATURE
1.	Prof. Dr. Fuad Ahmad Khan Niazi Head Of ophthalmology Department RMU & Allied Hospitals, Rawalpindi	
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5.	Prof. Muhammad Umar (Hilal-e-Imtiaz, Sitara-e-Imtiaz) (MBBS, MCPS, FCPS, FACG, FRCP (Lon), FRCP (Glasg), AGAF Vice Chancellor Rawalpindi Medical University & Allied Hospitals	