



Hypertension

Disorders of surgical importance

Disorders of surgical importance

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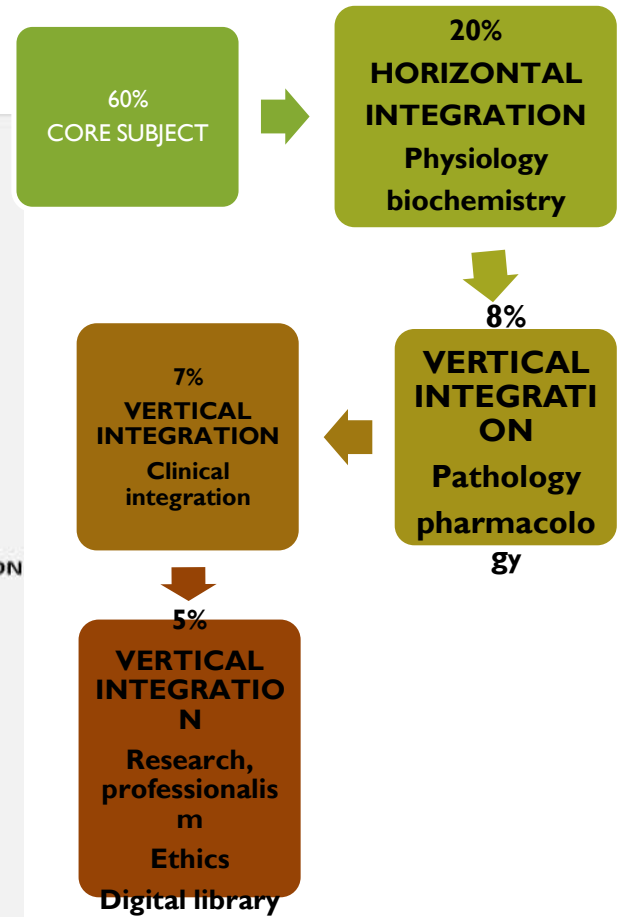
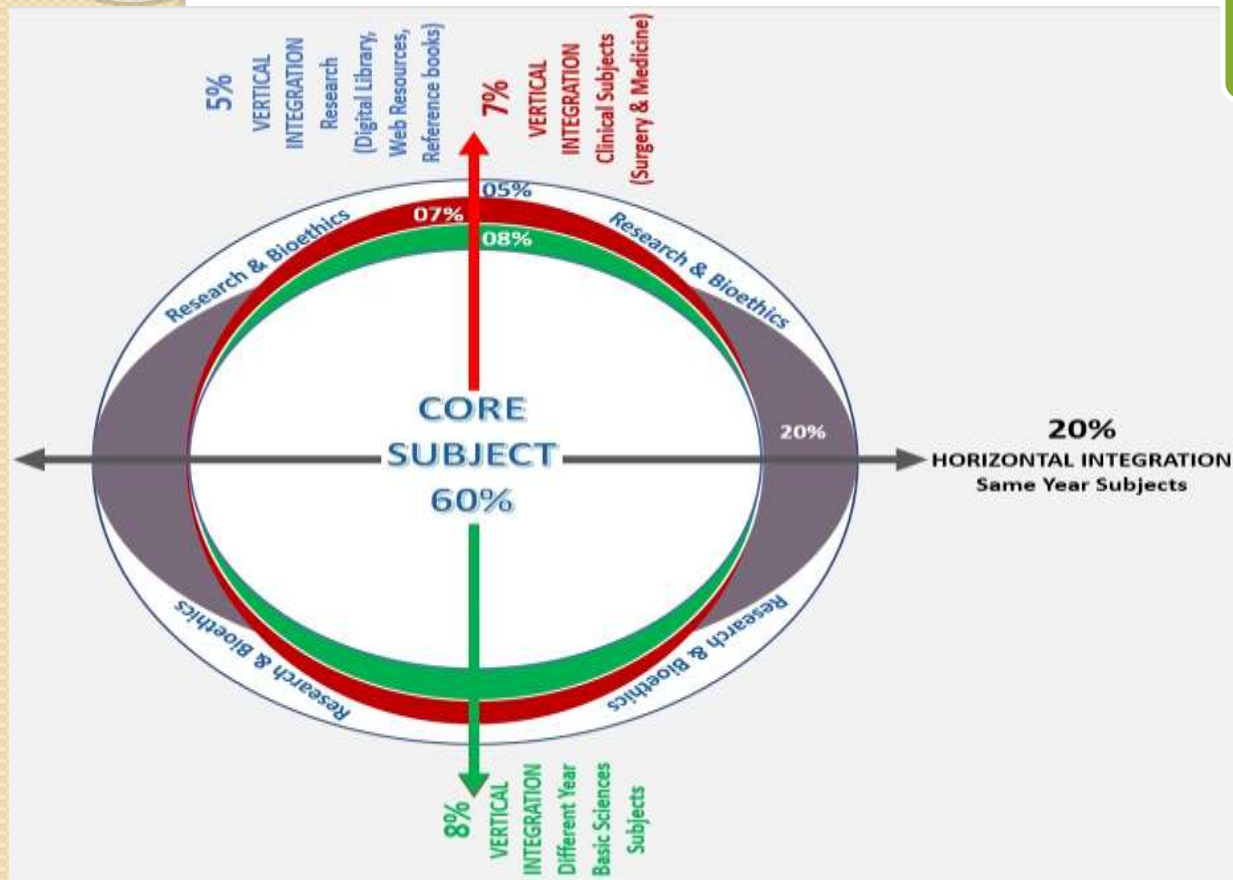


Mission Statement of RMU



- To impart evidence based research oriented medical education
- To provide best possible patient care
- To inculcate the values of mutual respect and ethical practice of medicine

Professor Umar Model of Integrated Lecture



Learning Outcomes

By the end the participants will be able to :

- Describe the applied anatomy and physiology of adrenal glands.
- Enumerate the different types of adrenal disorders.
- Describe adrenal disorders of surgical importance.

Screening

- Testing can be expensive and requires clinical suspicion and knowledge of limitations of different tests
- General principles:
 - New onset HTN if <30 or >50 years of age
 - HTN refractory to medical Rx ($>3-4$ meds)
 - Specific clinical/lab features typical for dz
 - i.e., hypokalemia, epigastric bruits, differential BP in arms, episodic HTN/flushing/palp, etc

Causes of Secondary HTN

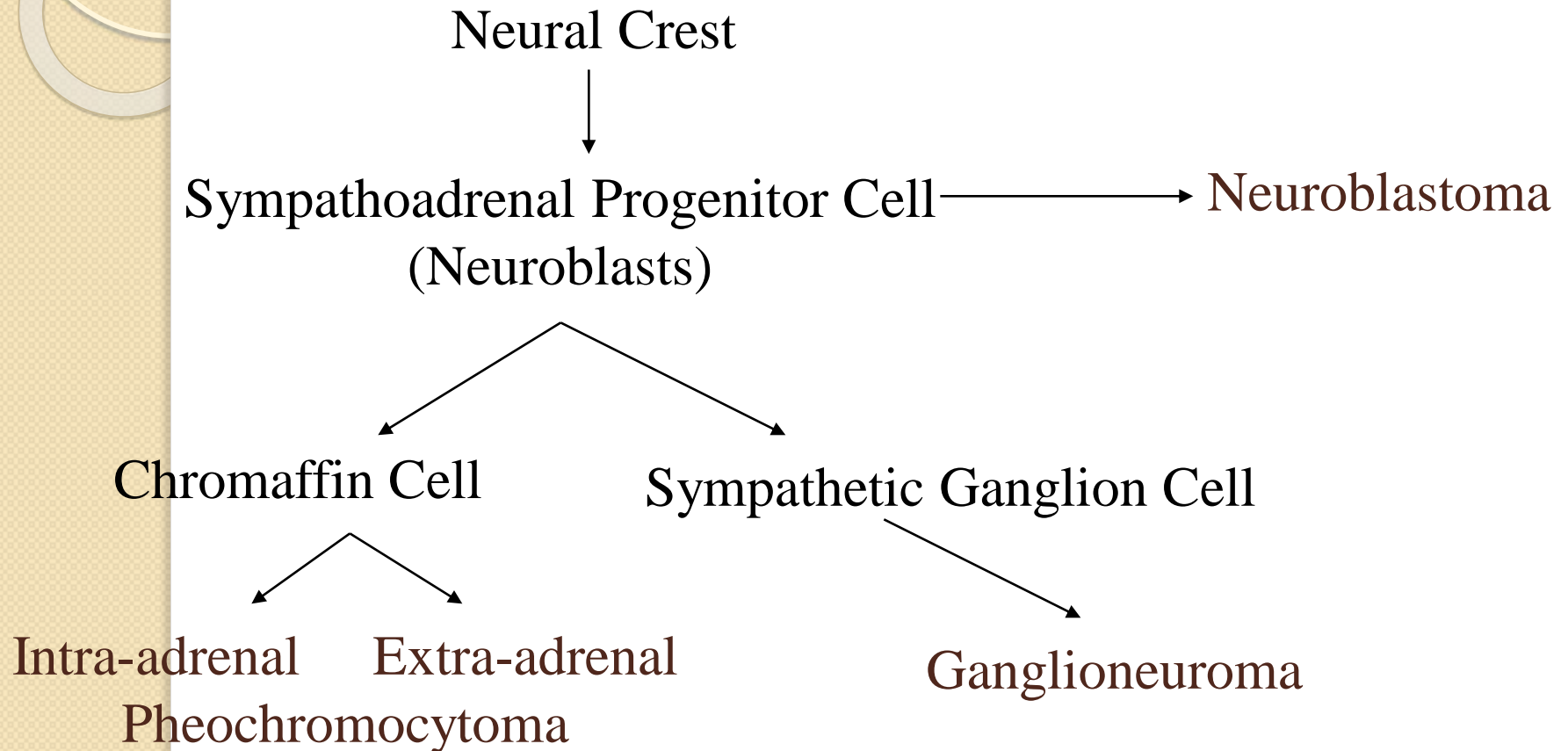
- Common

- Intrinsic Renal Disease
- Renovascular Dz
- Mineralocorticoid excess/ aldosteronism
- ? Sleep Breathing d/o

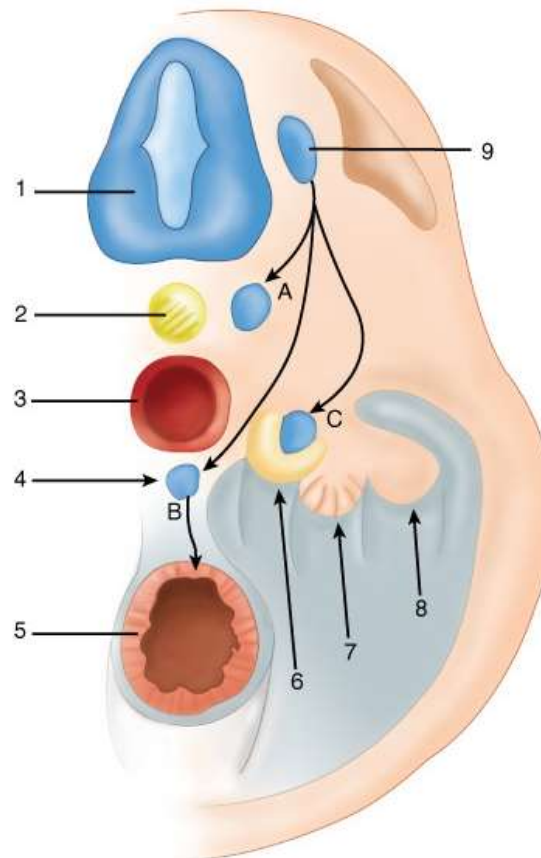
- Uncommon

- Pheochromocytoma
- Glucocorticoid excess/ Cushing's dz
- Coarctation of Aorta
- Hyper/hypothyroidism

Catecholamine Producing Tumors



1. Neural tube 2. chorda 3. aorta 4. base of mesentery 5. digestive tube 6. adrenal cortex 7. undifferentiated gonad 8. mesonephros 9. Neural crest



Source: Brunicaudi FC, Andersen DK, Billiar TR, Dunn DL, Hunter JG, Matthews JB, Pollock RE: *Schwartz's Principles of Surgery, 9th Edition*: <http://www.accessmedicine.com>
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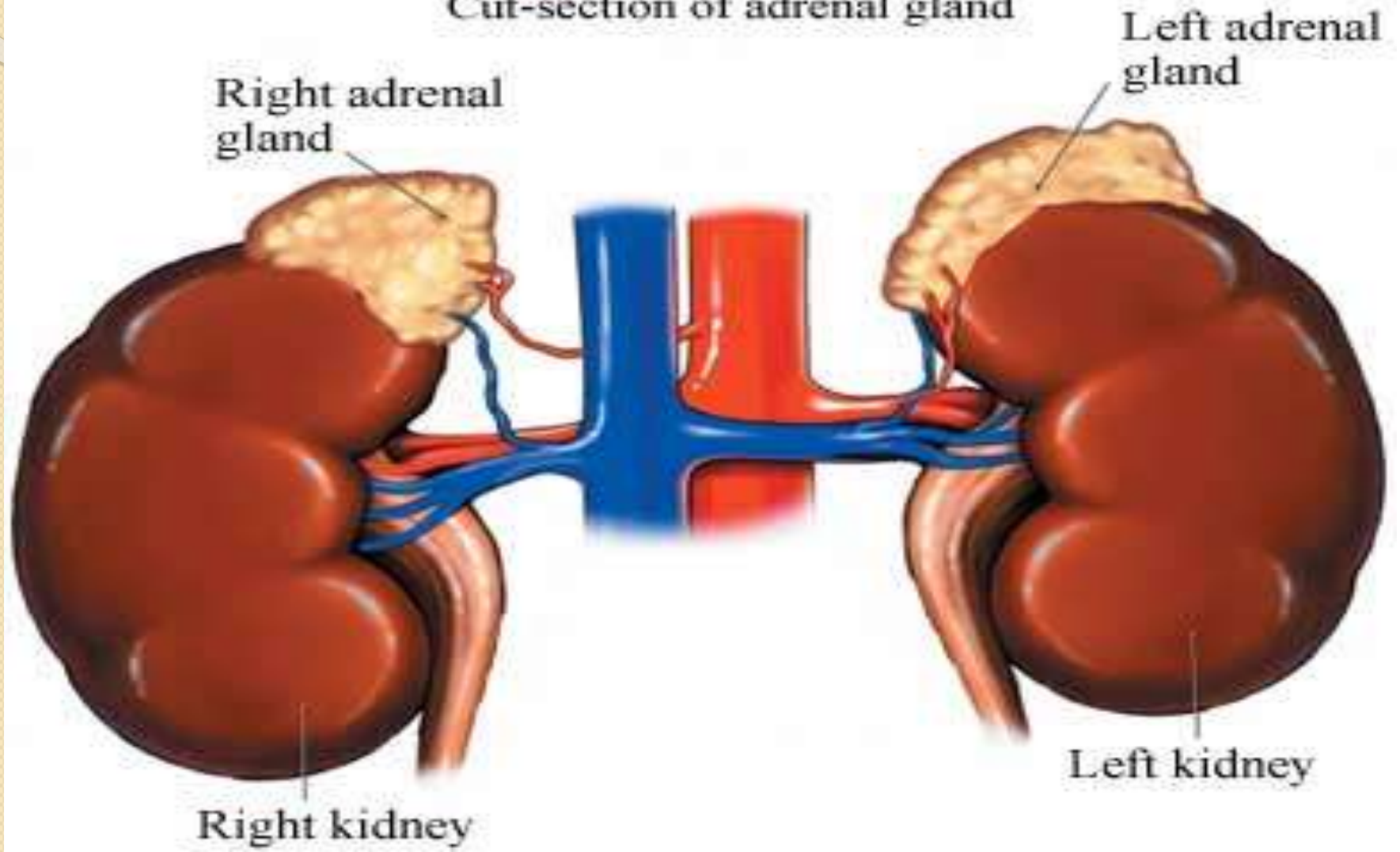
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Go to PC settings to activate Windows.

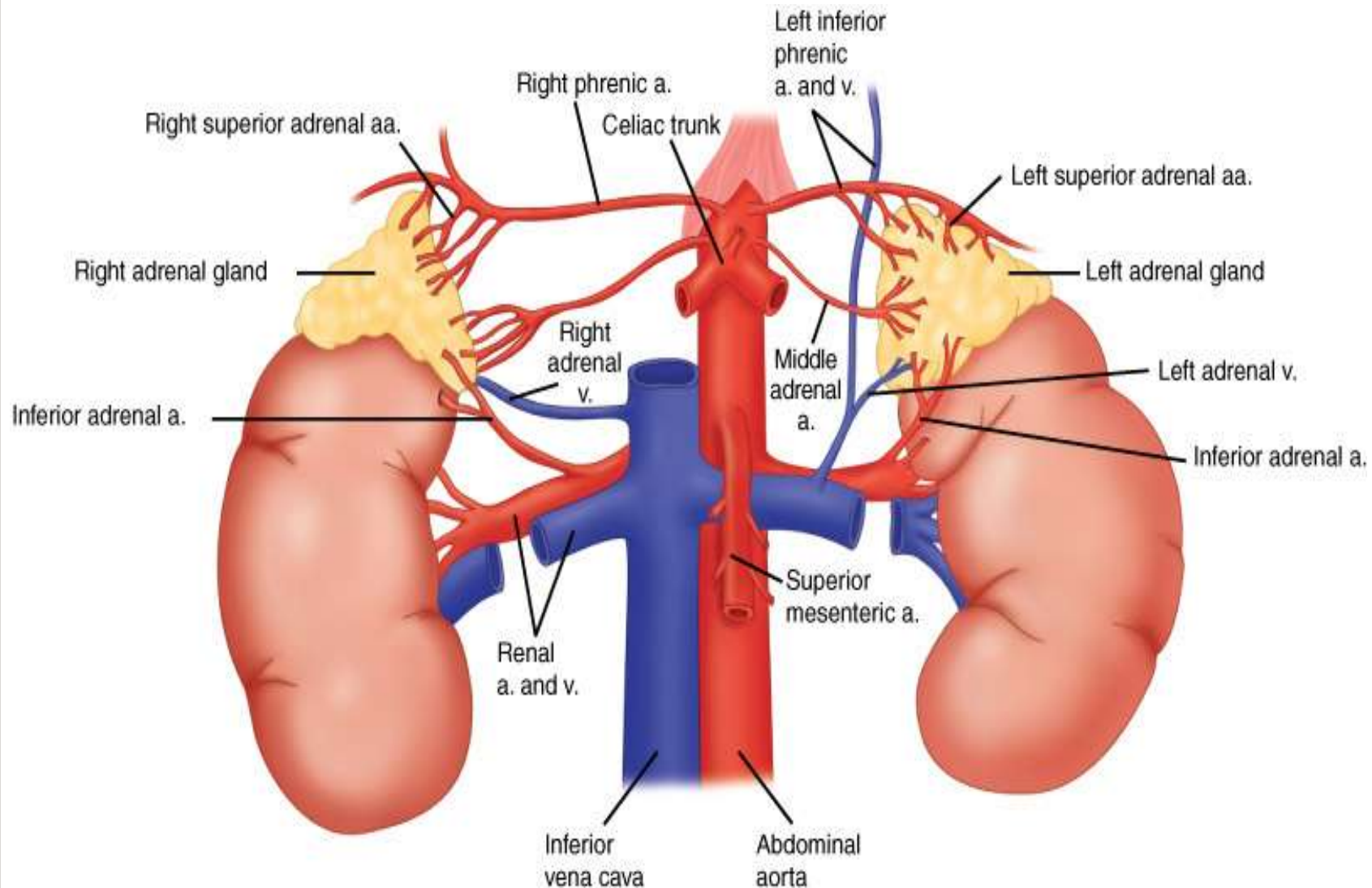
Anatomy

- Flattened yellowish
- Less than 10 gm
- Pyramidal and crescentic
- Rich blood supply

Historical background

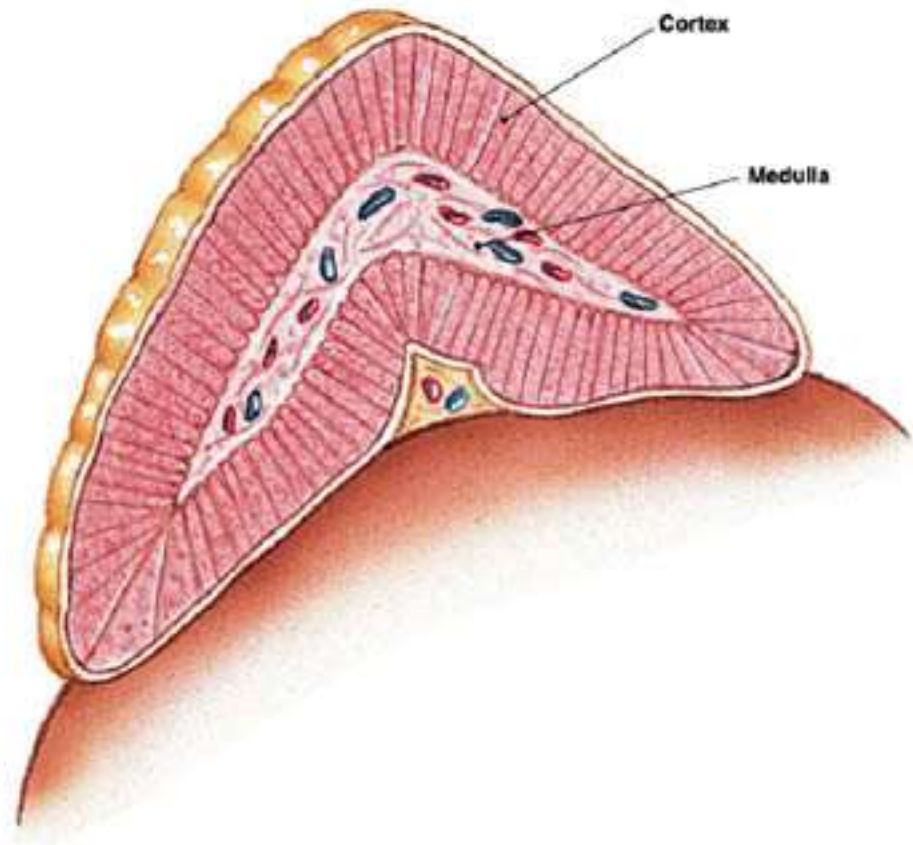
- Eustachius : anatomic account of the adrenals in 1563.
- Anatomic division of the adrenals into the cortex and medulla was described much later, by Cuvier in 1805.
- Thomas Addison in 1855 features of adrenal insufficiency, which still bear his name.
- .Pheochromocytomas were first identified by Frankel in 1885, but were not named as such until 1912 by Pick, who noted the characteristic chromaffin reaction of the tumor cells
- .Adrenaline was identified as an agent from the adrenal medulla that elevated blood pressure in dogs and was subsequently named epinephrine in 1897.
- The first successful adrenalectomies for pheochromocytoma were performed by Roux in Switzerland, and Charles Mayo in the United States.





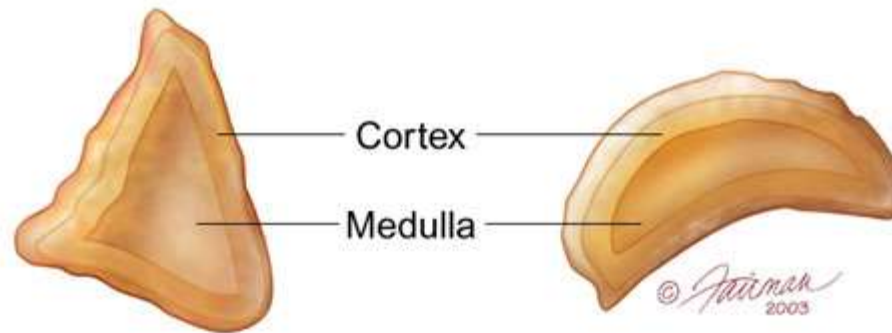
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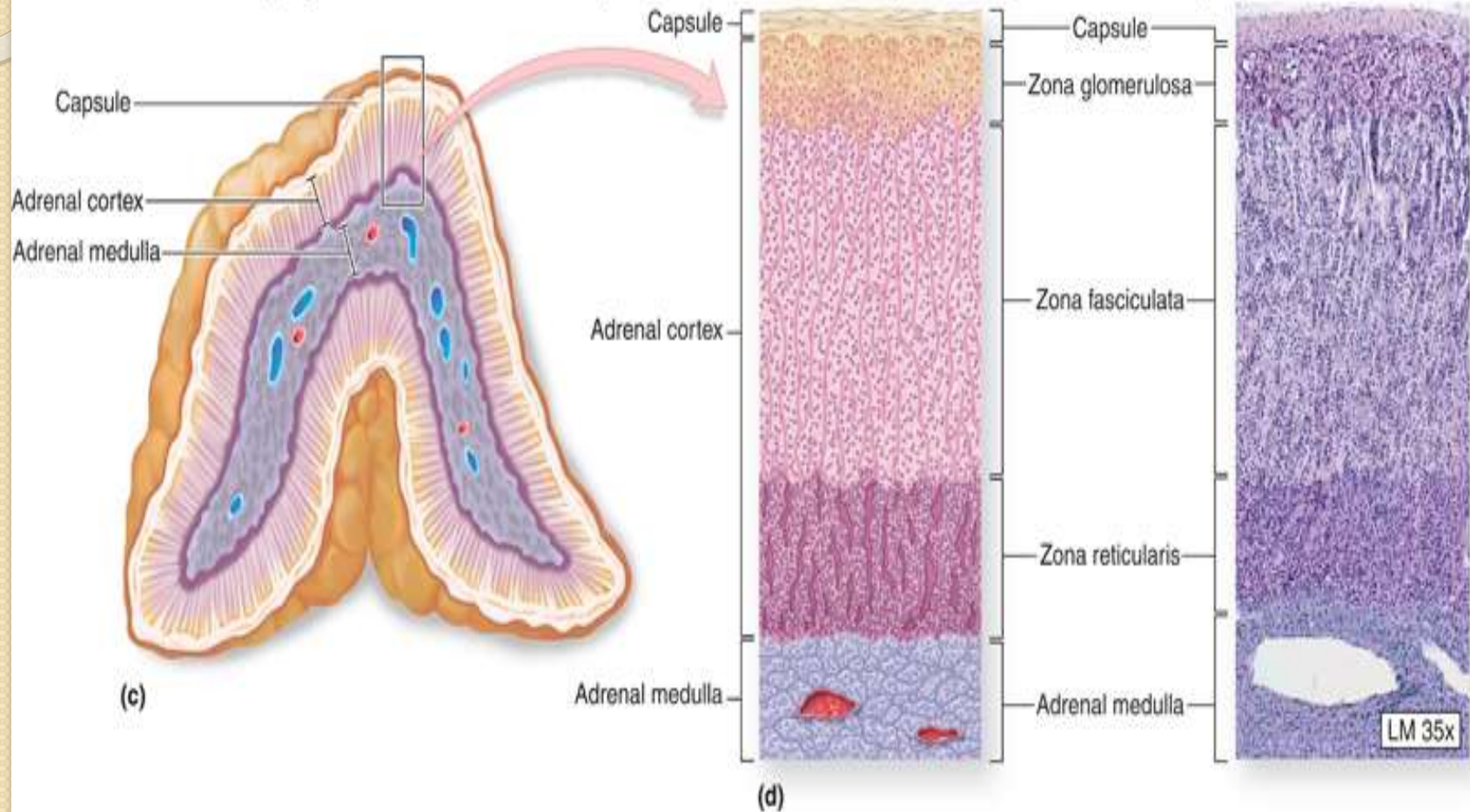


Right adrenal gland

Left adrenal gland

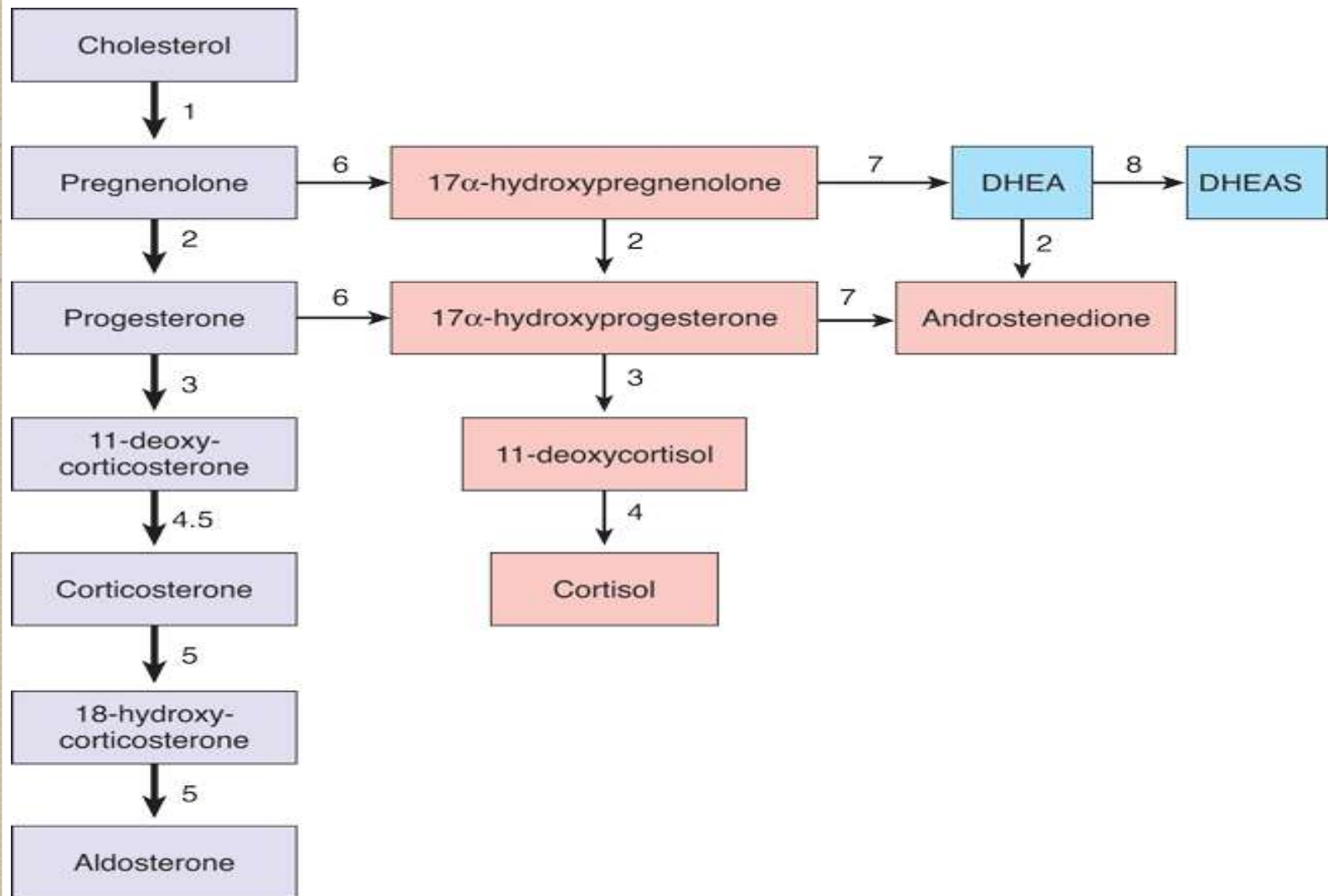


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Applied Physiology

- Essential for life
- Cortex secretes steroid hormones synthesized from cholesterol
- Zona glomerulosa produce mineralocorticoid aldosterone
- Zona fasciculata and reticularis synthesize glucocorticoids, androgens and oestrogens
- Adrenal medulla secretes catecholamines, adrenaline, noradrenaline and dopamine



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Adrenal Cortex

Hypocorticism

- i. Acute (Adrenal apoplexy of the newborn)
- ii. Chronic (Addisons Disease)

Hypercorticism

- i. Infantile
- ii. Prepubertal
- iii. Adult (Cushings Syndrome)
- iv. Postmenopausal
- v. Primary Aldosteronism

Adrenal Medulla

Tumours

- i. Ganglioneuroma
- ii. Neuroblastoma
- iii. Pheochromocytoma

Disorders of Surgical Importance

- Primary Aldosteronism
- Tumours of adrenal Medulla

Primary Aldosteronism

primary hyperaldosteronism PHA

- Conn's Syndrome
- Surgically Correctable type of Hypertension
- (1-2%) of all

Primary Aldosteronism

Pathology

- Hyperaldosteronism may be secondary to stimulation of the renin-angiotensin system from renal artery stenosis and to low-flow states such as congestive heart failure and cirrhosis
- Autonomous Excessive Aldosterone Secretion.
- 80% of patients with PHA have unilateral adrenocortical adenoma(Conn's syndrome).Remainder have bilateral hyperplasia

Primary Aldosteronism

- $[\text{Na}^+]$
- $[\text{K}^{++}]$



- $[\text{Na}^+]$

- $[\text{K}^{++}]$

Primary Aldosteronism

Clinical Features:

Characterized by hypertension and hypokalemia

1) Muscular Weakness

2) Polyuria

3) Polydypsia

4) Hypertension

5) Edema

Primary Aldosteronism

Investigations:

- S. potassium is low
- Elevated urinary potassium
- CT
- MRI



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Primary Aldosteronism

Treatment

Medical

Surgical

Unilateral Adrenalectomy

Neuroblastoma

- Neural Crest Cells
- Location

$\frac{3}{4}$ abdomen

$\frac{1}{2}$ of them in adrenals

Neuroblastoma

Clinical Features:

- Age:

90% < 8yrs

- Abdominal Mass

- Weight loss
- Abdominal Pain
- Distension

Neuroblastoma

- Metastasis:

50% of infants



Neuroblastoma

Investigations:

- Urine
- Abdominal Ultrasound
- CT Scan

Neuroblastoma

Treatment:

- Surgical Excision
- Neoadjuvant Chemo/radiotherapy




Pheochromocytomas

Pheochromocytomas

Definition

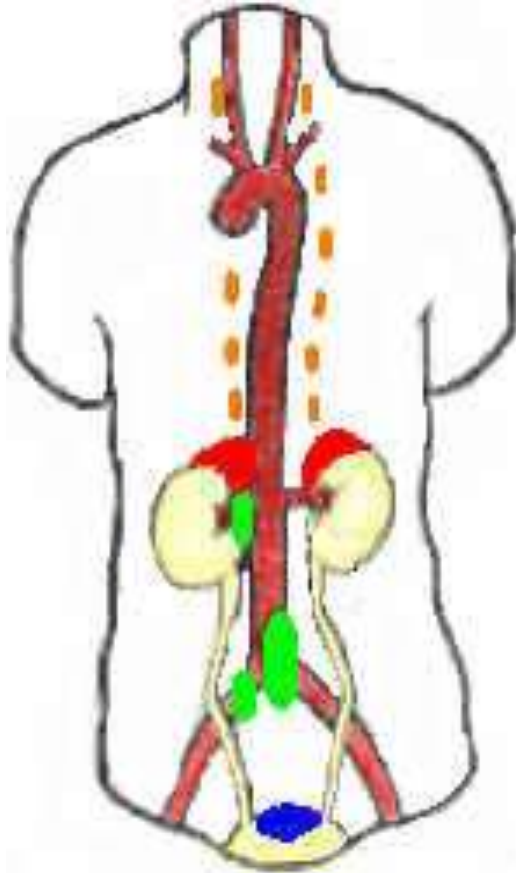
Pheochromocytomas are neoplasms derived from the chromaffin cells of

- adrenal medulla (90%)
 - extra-adrenal paraganglia (10%)
-
- result in unregulated, episodic oversecretion of catecholamines

- 
- Phaeochromocytoma, unilateral or bilateral occurs in association with following hereditary syndromes
 - Multiple endocrine neoplasia type 2 (Sipple's syndrome)
 - Von Hippel-Lindau (VHL)
 - Renal cell carcinoma
 - CNS and retinal haemangioblastoma
 - Neurofibromatosis type I
 - Familial paraganglioma syndrome

Pheochromocytoma : location

- Virtually all (99%) arise within the abdomen



	Adrenal medulla	:90%
	Extra-adrenal	:10%




Pheochromocytomas

- Uncommon neoplasms
- 0.1% to 0.3% of all hypertensive patients
- Surgically correctable form of hypertension

Phaeochromocytoma

THE 10% TUMOUR

- 
- extra-adrenal
 - children
 - familial
 - bilateral
 - malignant

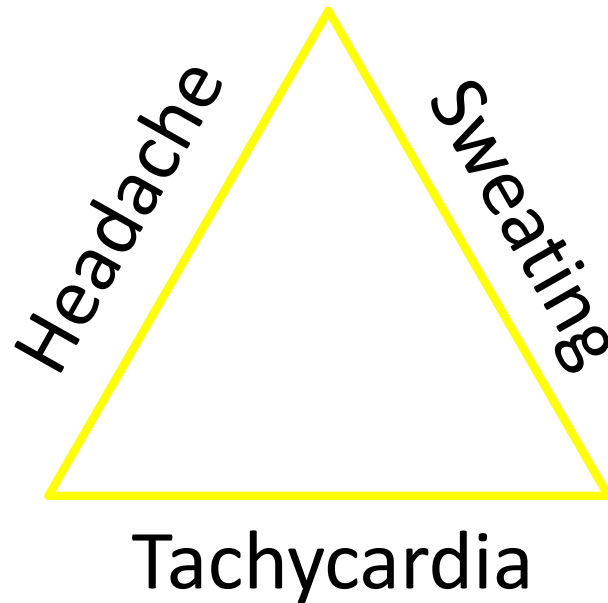
10 %

Clinical Features

- Usually present before the age of 50 years
Commonly 3rd to 5th decades
- Male = Female
- Manifests by effects of adrenaline and noradrenaline excess

Classical Presentation

- Paroxysms manifesting as a triad of:



Classical Presentation

- **Additional symptoms during an attack :**
 - Blood pressure may rise to 200/100 mmHg
 - Paroxysmal hypertension in 50% patients
 - Palpitations
 - Extreme anxiety
 - Tremors
 - Paresthesias
 - Chest or abdominal pain
 - Nausea, vomiting
 - Dyspnea

Other Presentations

- **Persistent Hypertension**
 - All young hypertensive patients should be screened for a catecholamine-secreting tumor
 - Surgically correctable

Catecholamine crises

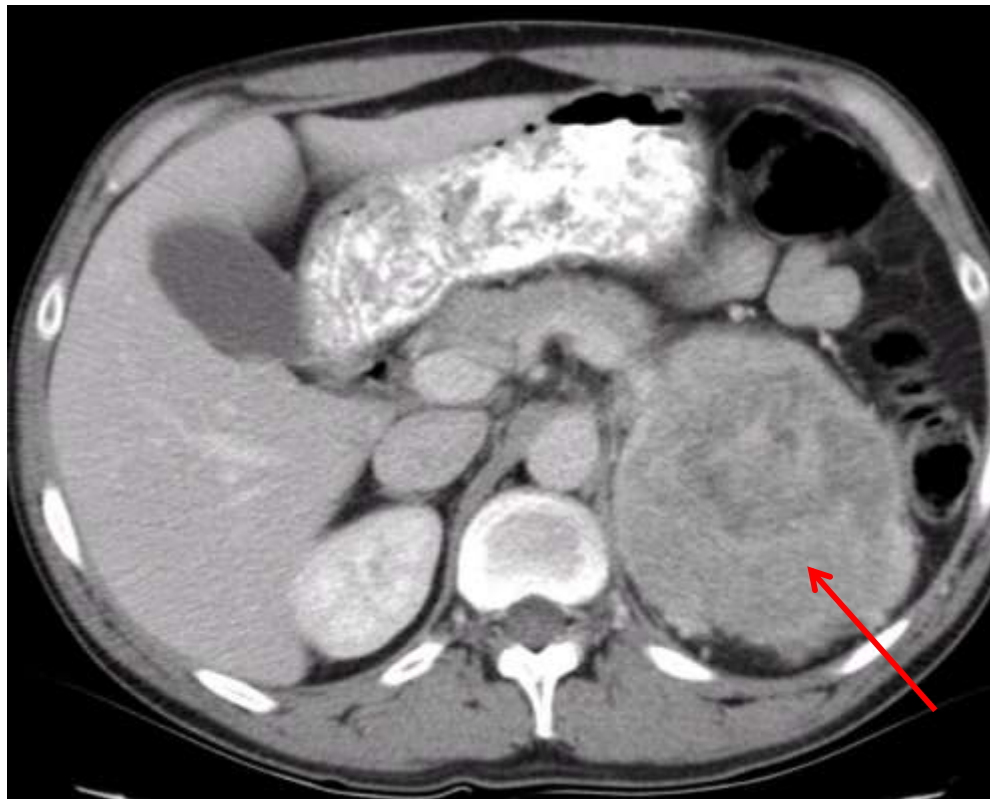
- Sudden and severe increase in blood pressure
- Can lead to
 - Heart failure
 - Pulmonary edema
 - Arrhythmias
 - Intracranial hemorrhage

24 Hours Urinary Catecholamines

- 24 hours urine collection
- Analyzed for catecholamines and their metabolites:
 - Adrenaline and Noradrenaline
 - Metanephrines
 - Vanillylmandelic acid (VMA)
- Diagnostic values:
 - > 2-3 fold elevation

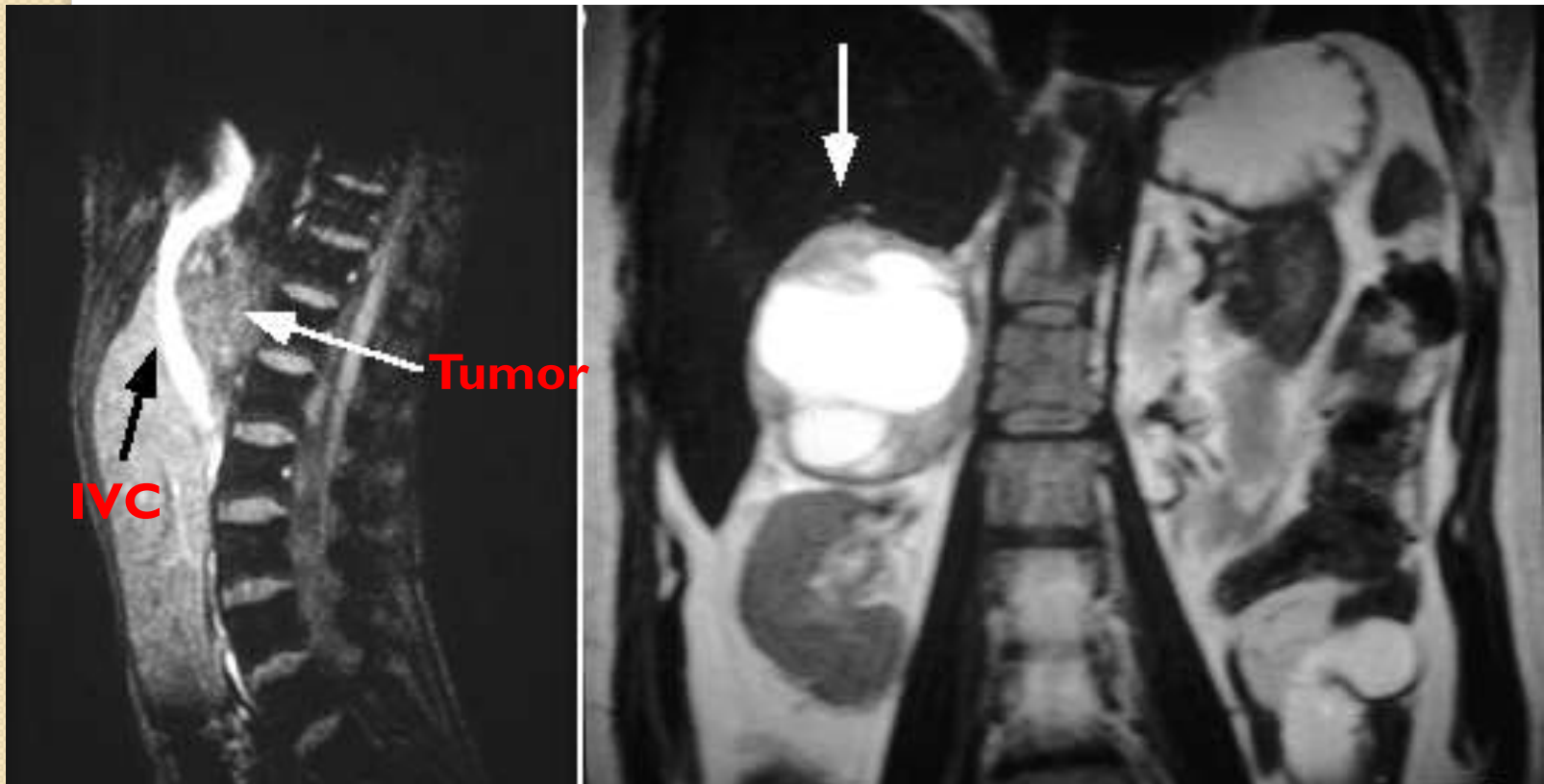
Localization: Imaging

- CT abdomen



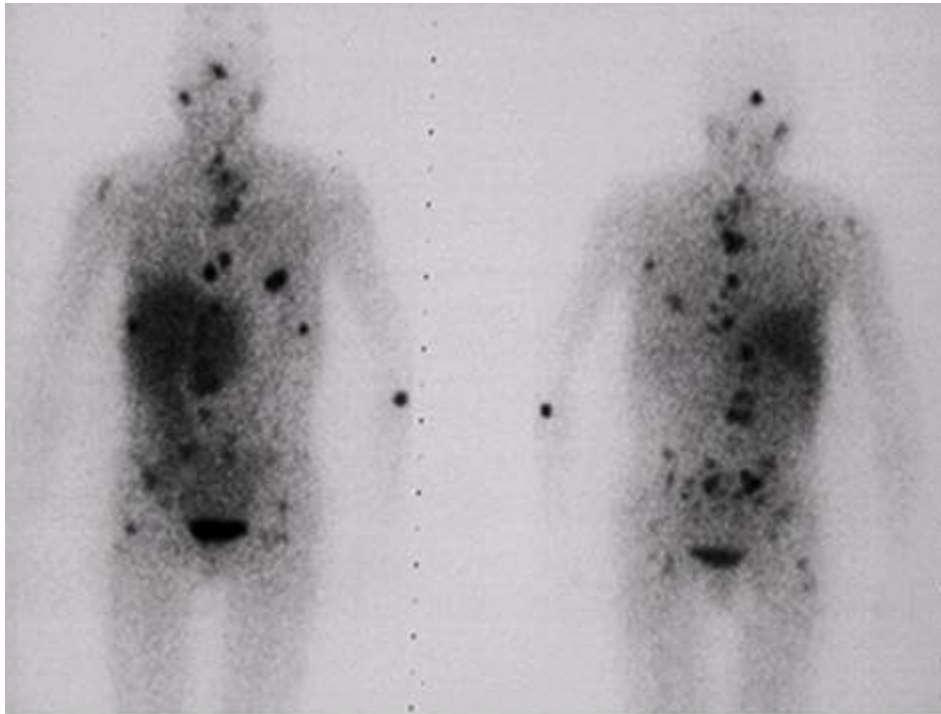
Localization: Imaging

- MRI
 - More sensitive for extra-adrenal pheochromocytoma



Localization: Imaging

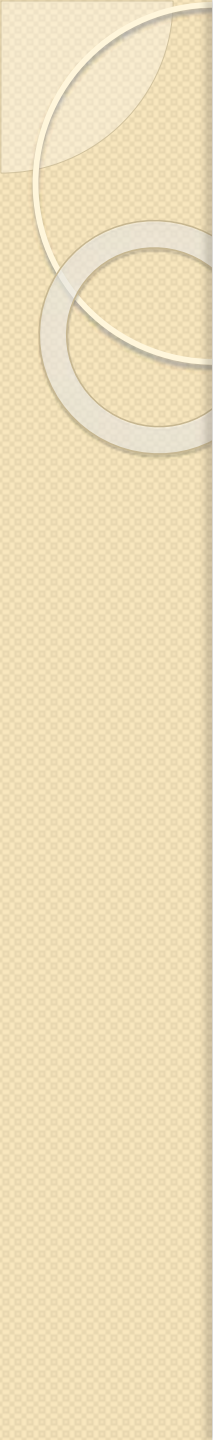
- MIBG (^{131}I -meta-iodobenzylguanidine)
Scintigraphy



Diffuse metastatic pheochromocytoma

Preoperative Preparation

- Surgery without preoperative preparation is **dangerous**
 - Risks:
 - Perioperative hypertensive crisis
 - Post excisional hypotension

- 
- The patient should come to operation with
 - Blood pressure and pulse rate controlled
 - Adequate hydration


- 
- Preoperative control of blood pressure
 - α -adrenergic blocker
 - B-blockers

Preoperative Preparation

- Volume expansion to avoid post-excision hypotension
 - Liberal oral salt and water
 - Intravenous hydration night before surgery

Measures to avoid cardiovascular instability during surgery

- Continuous monitoring
 - ECG
 - Arterial line
 - CVP line / Swan-Ganz catheter
- Avoid excessive tumor manipulation, which can result in catecholamine release

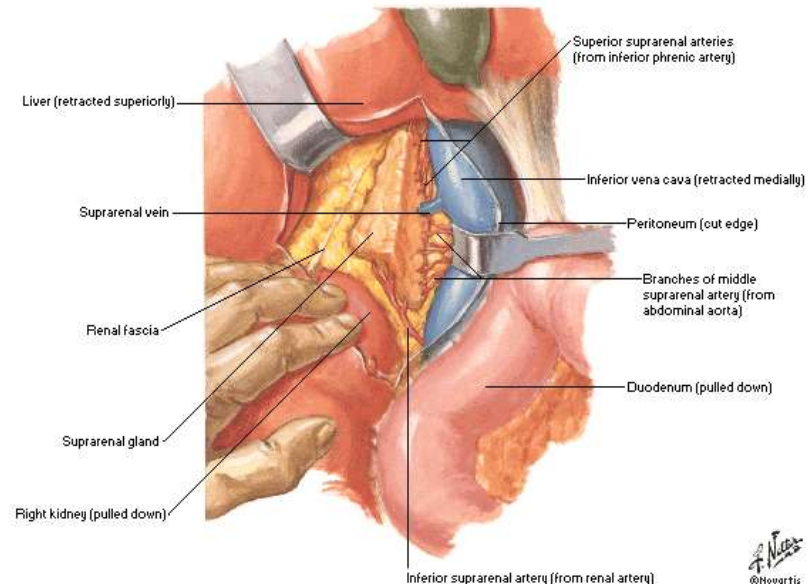
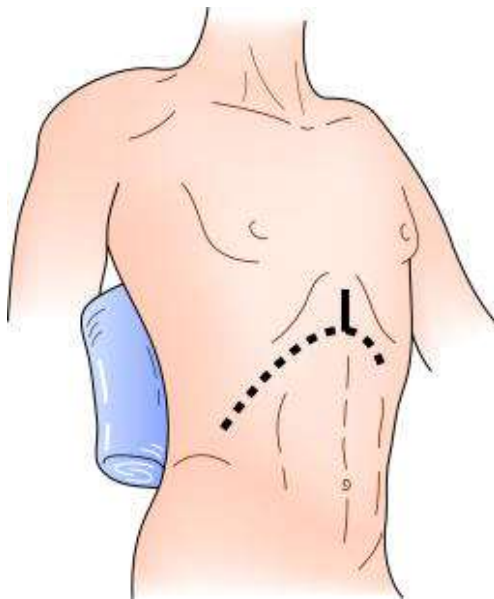
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- Operative mortality prior to 1950 :
24 - 50 %
 - Current operative mortality :
0 - 2.7 %

Operative Procedures

- Adrenalectomy
 - Open
 - Laparoscopic
- Excision of extra-adrenal paraganglioma

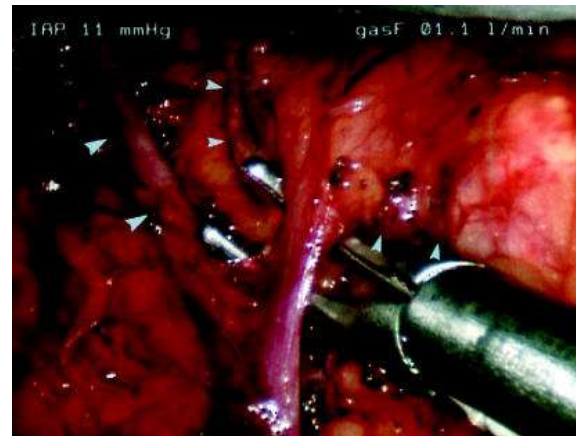
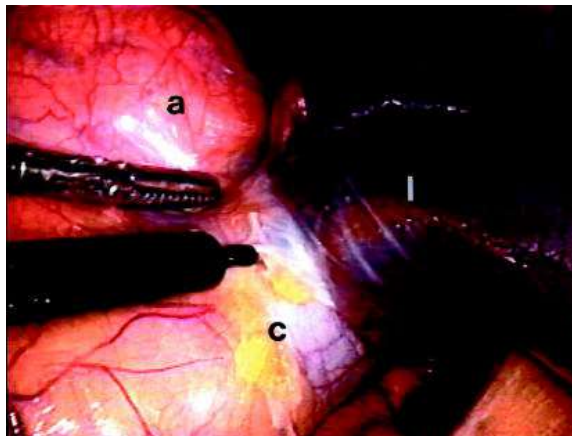
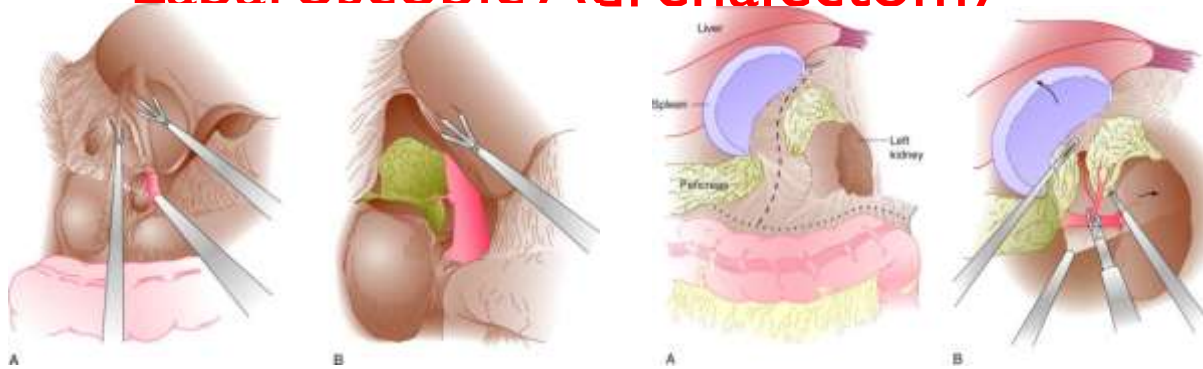
Operative Procedures

■ Open Adrenalectomy



Operative Procedures

■ Laparoscopic Adrenalectomy





Any Questions



Thanks