ADRENAL TUMORS

Specific Objectives

At the end of the class students will be able to:-

- Define Adrenal Tumor .
- Enlist the etiological factors Adrenal Tumor.
- Explain about the pathophysiology of Adrenal Tumor
- Enlist the clinical manifestations of Adrenal Tumor
- Enumerate the diagnostic evaluation of Adrenal Tumor.
- Explain the management of Adrenal Tumor

DEFINITION

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Pheochomocytoma is a catecholamine secreting tumor of the chomaffin cells of the sympathetic nervous system, it is usually found in adrenal medulla.
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ETIOLOGY

Idiopathic

May be neuroectodermal disease with multiple endocrine neoplasia.

PATHOPHYSIOLOGY

- Due to causes
- It affect on adrenal medulla
- Develop on chromatin tissues found in sympathetic pone ganglia
- Excessive secretion of epinephrine & norepinephrine
- Signs & symptoms

SIGNS & SYMPTOMS

- Hypertension
- Headache
- Polyuria
- Hyper-metabolism
- Hyperglycemia
- Tremor
- Anxiety, flushing, nausea & vomiting

DIAGNOSTIC EVALUATION

- History collection
- Physical examination
- Urine & plasma catecholamine level
- MRI
- CT scan
- Scintigraphy

MEDICAL MANAGEMENT

- Preliminary preparation includes control of blood pressure & blood volume.
- Bed rest
- Hydration to be maintained before, during & after surgery.
- Phenoxybenzamine is a long acting alpha blocker is used after the blood pressure is stable.

SURGICAL MANAGEMENT

- Adrenalectomy but contraindicated for this surgery if there is-
- Hypertensive retinopathy, nephropathy.
- Increased platelet aggregation
- Stroke
- Heart, renal failure.

Nursing Management

- Obtain history of sign & symptoms
- Monitor ECG changes, arterial pressure, fluid
 & electrolyte balance
- Blood glucose levels

COMPLICATIONS

- Excessive discharge of hormone during induction of anesthesia
- Metastasis of tumor.