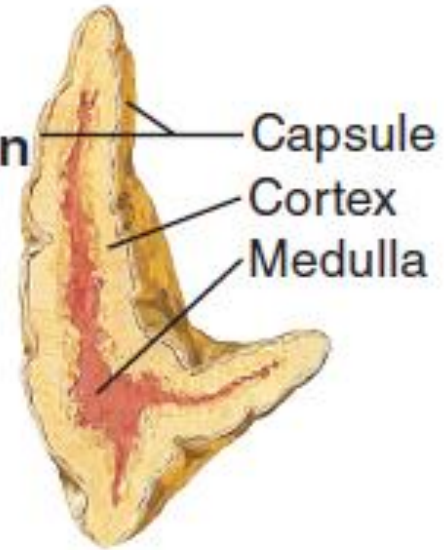
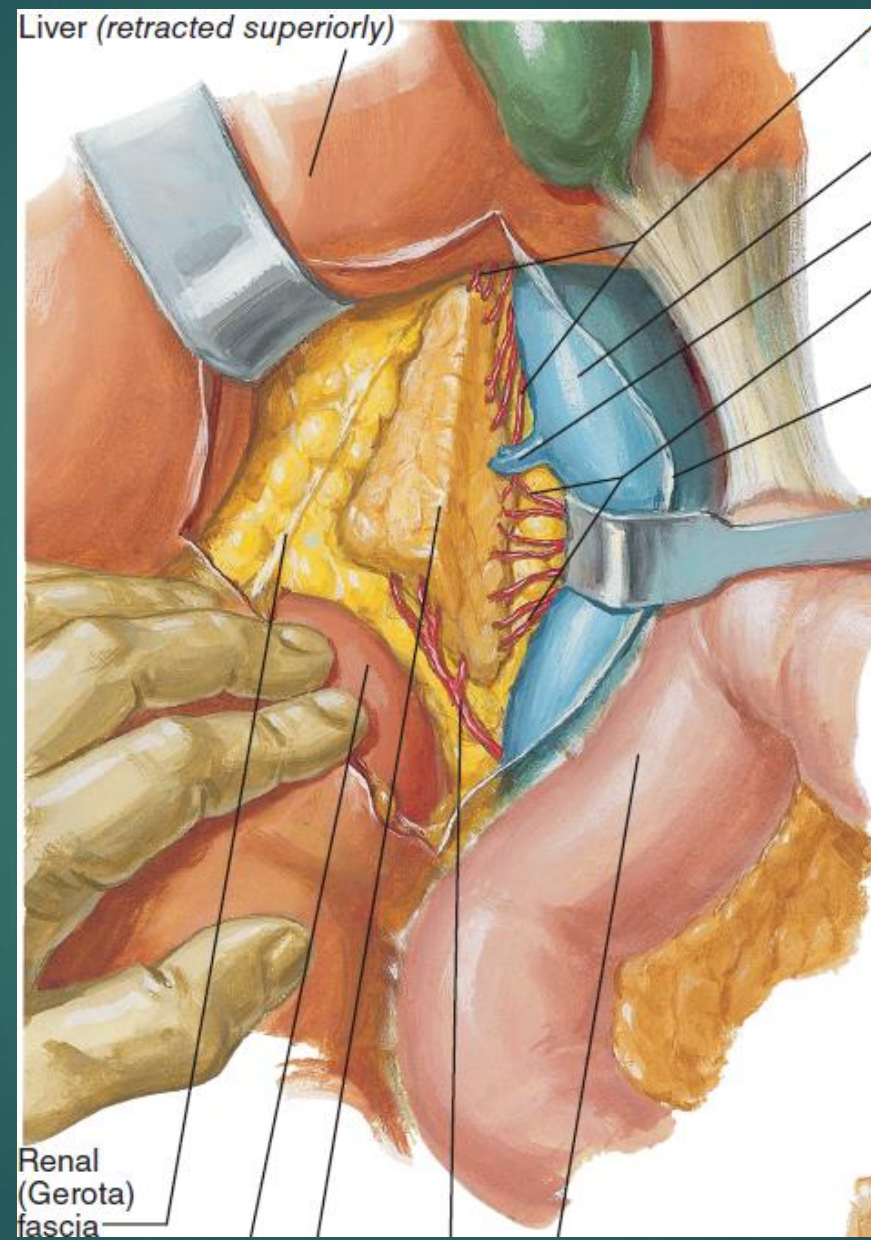


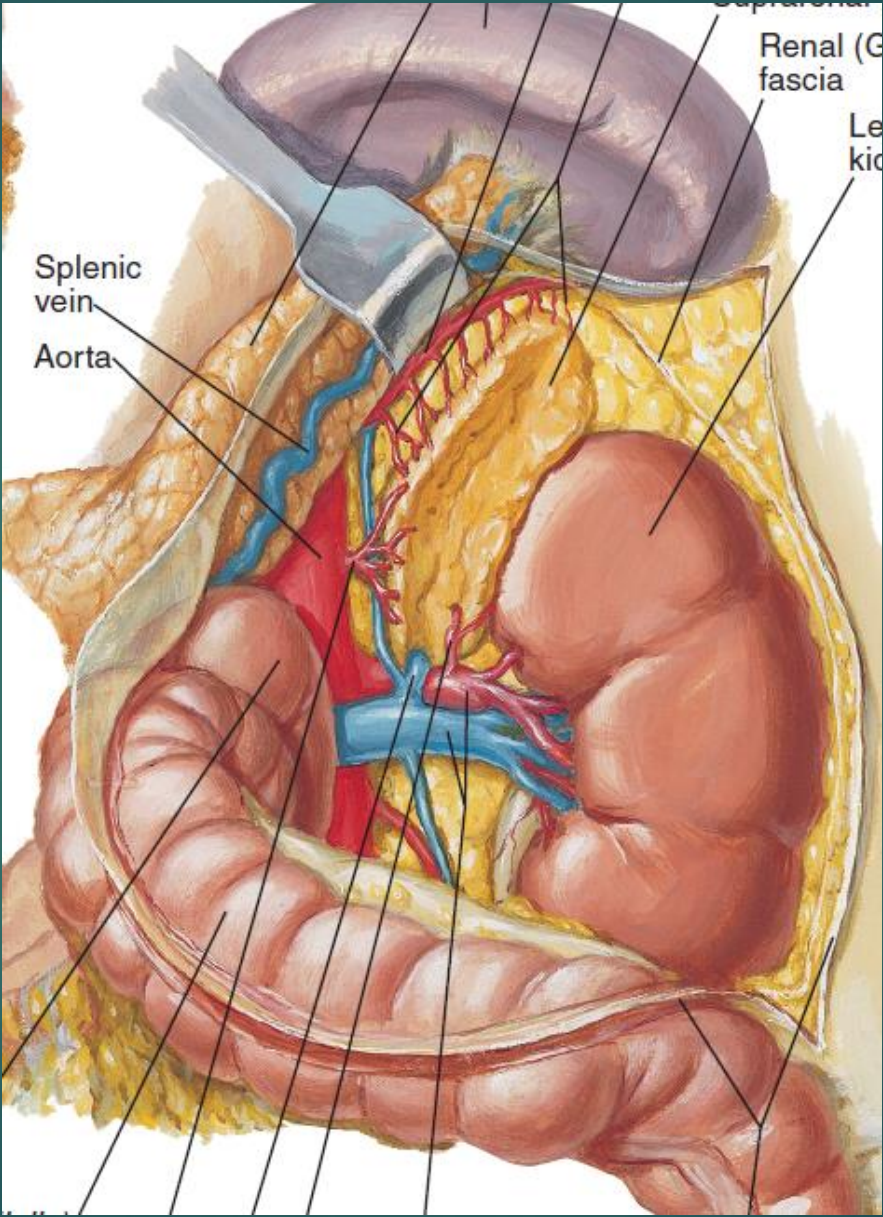


# Surgical Approaches to the Adrenal Gland

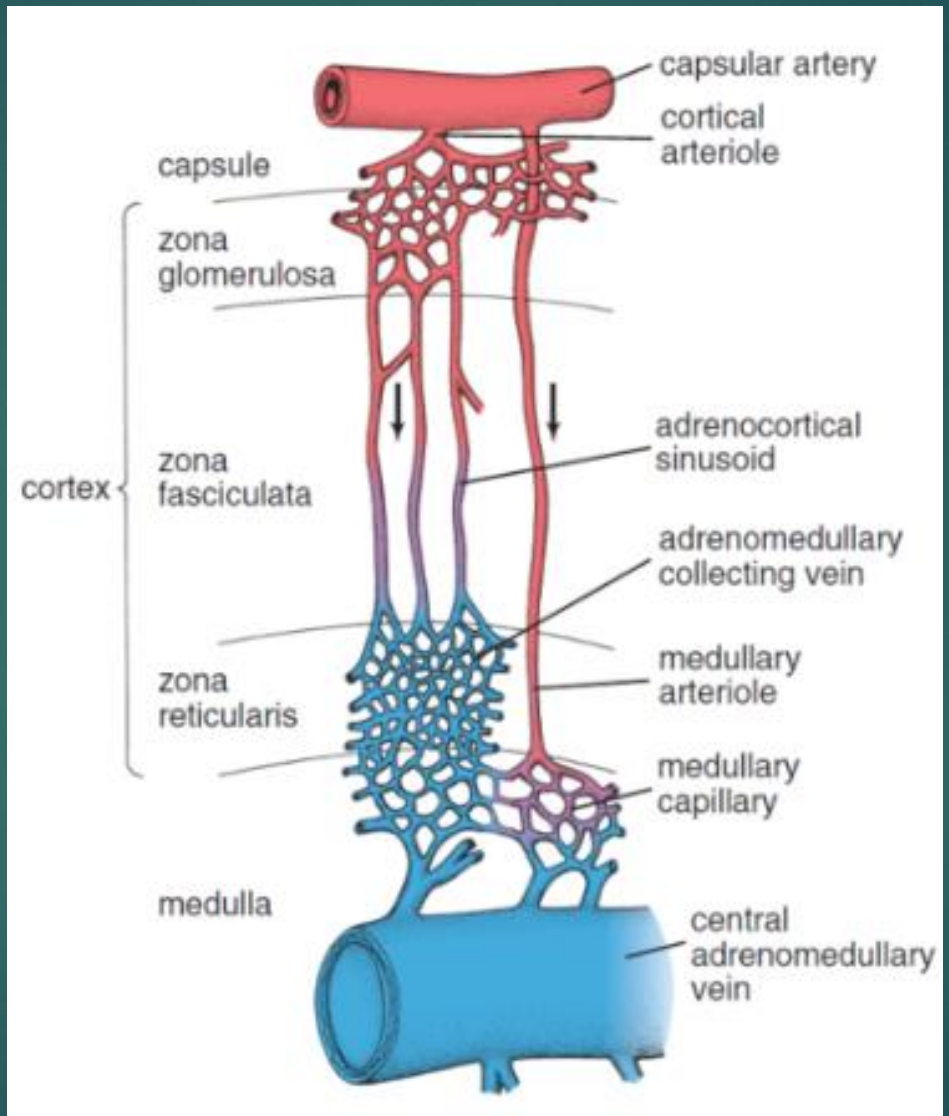
**Cross section  
through  
suprarenal  
gland**











# OPERATIVE APPROACHES

- ▶ Transabdominally
- ▶ Retroperitoneally

# Selection of operative approach

## Surgeon expertise

When the surgeon is experienced with more than one approach,

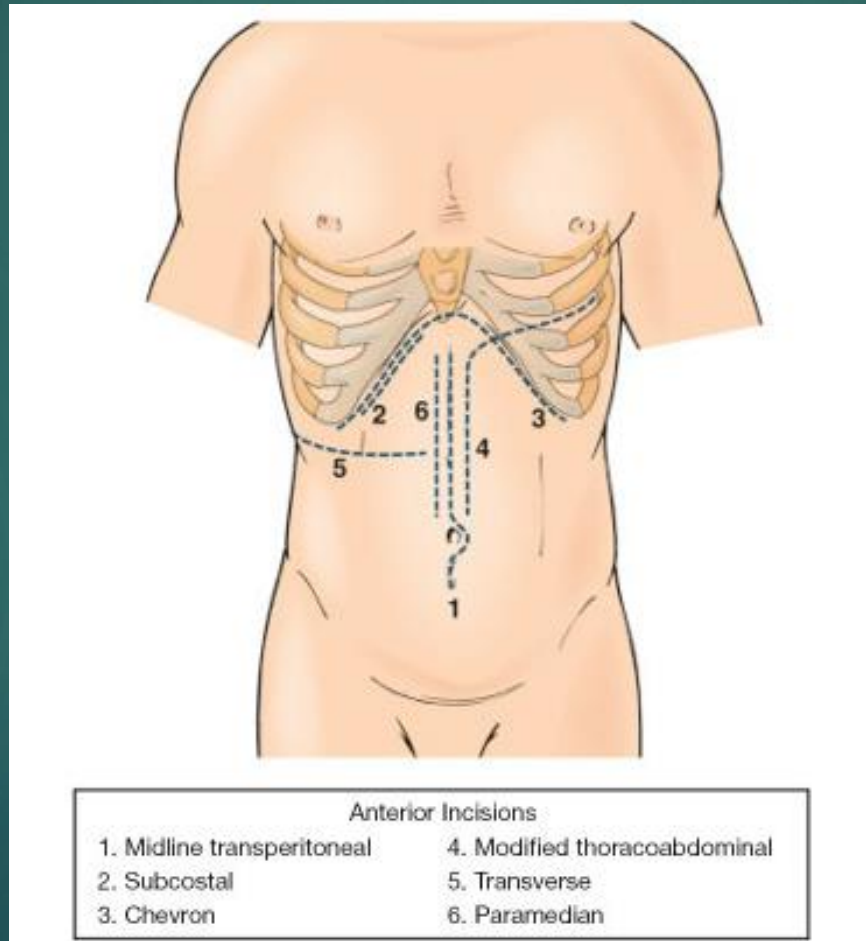
- ▶ large or malignant tumors
- ▶ bilateral tumors
- ▶ extensive prior abdominal surgeries
- ▶ patient body habitus

# Open Vs MIS?

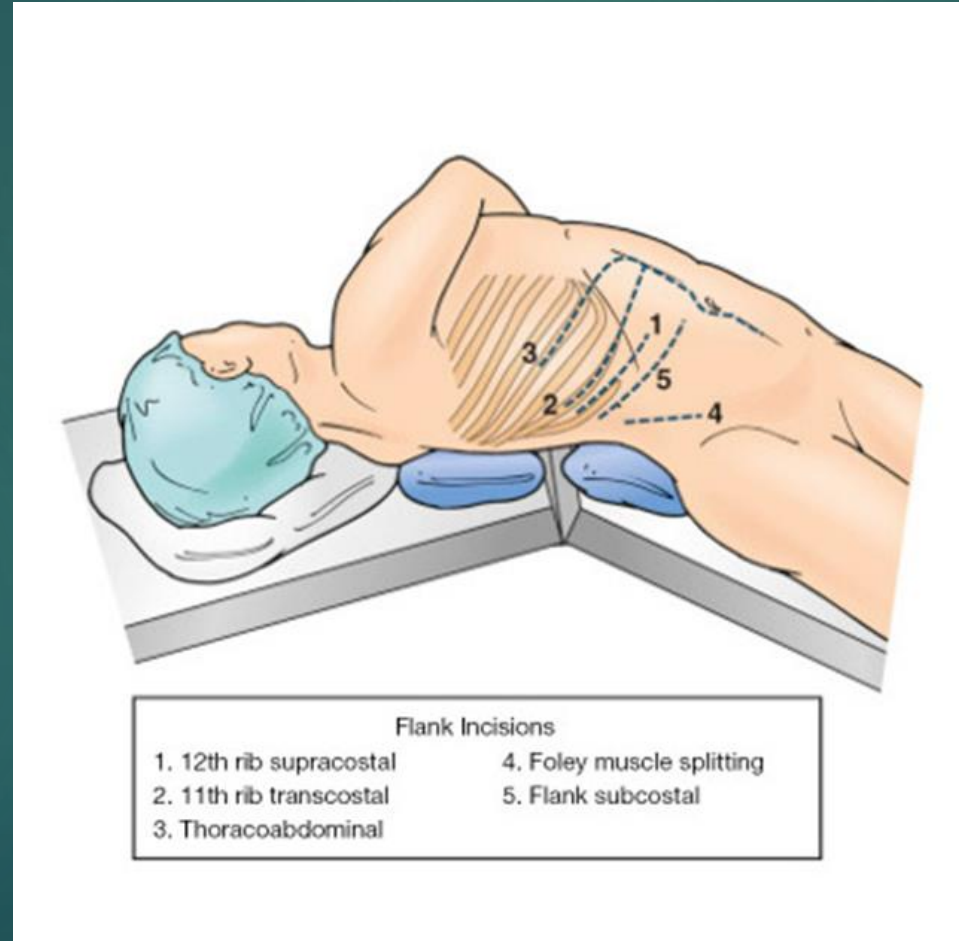
- ▶ Primary adrenal **malignancy** of any size (eg, adrenal cortical carcinoma, malignant pheochromocytoma).
- ▶ Adrenal mass **invading** surrounding structures (eg, liver, kidney, inferior vena cava).
- ▶ Adrenal mass **suspicious** but not confirmed as a primary adrenal malignancy (eg, >6 cm, irregular margins, hemorrhage or central necrosis, hypervascular).
- ▶ Extensive prior upper abdominal or retroperitoneal procedures that preclude a minimally invasive surgical (MIS) approach.
- ▶ Concomitant procedures (eg, hepatic resection) not amenable to laparoscopic approach.



# Open Transabdominal Adrenalectomy



# Open Retroperitoneal Adrenalectomy



# Adrenocortical carcinoma

In the United States, open adrenalectomy with removal of adjacent lymph nodes is recommended regardless of size.

- ▶ National Comprehensive Cancer Network (NCCN) American Association of Clinical Endocrinologists (AAACE), and American Association of Endocrine Surgeons (AAES)

In Europe, laparoscopic adrenalectomy is performed for stage I or II ACC with a diameter < 10 cm.

- ▶ European Society of Endocrine Surgeons



Data on the safety and efficacy of MIS for suspected ACC are inconsistent

no significant differences in recurrence rates or complications between laparoscopic and open adrenalectomy among 517 patients undergoing surgery for adrenal malignancy

Sautter AE, et al. Am Surg. 2016

# Pheochromocytoma

laparoscopically or robotically when safe and feasible

- ▶ convert to open surgery in case of intraoperative **bleeding** (pheochromocytomas are hypervascular lesions) or **capsular violation** (risk of hypertensive crisis)

## Open adrenalectomy

- ▶ large (>6 cm)
- ▶ invasive pheochromocytomas
- ▶ paragangliomas



# Intraoperative Complications

## ACCESS RELATED

Abdominal wall hemorrhage  
Cutaneous nerve injury  
Visceral injury by Veress needle or trocar

## HEMORRHAGE

Inferior vena cava or aorta  
Adrenal vein  
Lumbar vein  
Hepatic vein  
Remnant adrenal gland after partial adrenalectomy

## ISCHEMIA

Ligation of renal artery or vein  
Ligation of superior mesenteric artery and vein

## INJURY TO NEIGHBORING ORGANS AS A RESULT OF THERMAL ENERGY OR INCORRECT PLANE OF DISSECTION

Lung—pneumothorax  
Pancreas  
Liver  
Spleen  
Stomach and bowel, especially duodenum  
Kidney

## HEMODYNAMIC INSTABILITY

Pheochromocytoma

# Postoperative Complications

## PRIMARY ALDOSTERONISM

Hypokalemia: secondary to continued potassium loss immediately postoperative

Hyperkalemia: secondary to failure of contralateral adrenal to secrete aldosterone

## CUSHING SYNDROME

Inadequate steroid replacement leading to hypocorticism

Fracture secondary to osteoporosis

Hyperglycemia

Poor wound healing

Increased risk of infections

## PHEOCHROMOCYTOMA

Hypotension secondary to  $\alpha$ -adrenergic blockade after tumor removal

## GENERIC COMPLICATIONS

Hemorrhage

Pneumothorax

Pancreatitis

Pneumonia

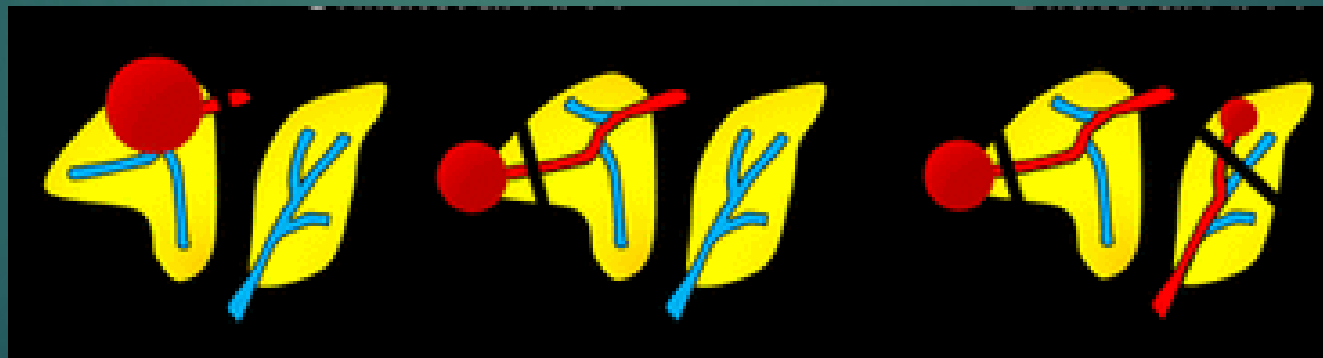
Prolonged ileus

Intra-abdominal collections

# Partial Adrenalectomy

no established consensus or guidelines

- ▶ lesions <3 cm that are located **anteriorly** or **laterally** in the adrenal gland (to preserve the adrenal vein)





The amount of adrenal tissue that must be left behind after partial adrenalectomy to avoid insufficiency **is not known**.

- ▶ at least a third of the adrenal gland should be preserved
- ▶ a 3- to 5-mm margin is recommended.

The overall recurrence rate was 8% (95% CI: 0.05–0.12)

- ▶ the least in the retroperitoneoscopic group 1% (95% CI: 0–0.04) and Conn's syndrome group 2% (95% CI: 0.01–0.05)
- ▶ highest in open group 15% (95% CI: 0.07–0.28) and Pheochromocytoma group 10% (95% CI: 0.07–0.16)

Steroid free rate were 85% (95% CI: 0.78–0.9)

- ▶ best in the Conn's syndrome group 97% (95% CI: 0.85–0.99) and laparoscopic group 88% (95% CI: 0.75–0.95)



Bilateral hereditary pheochromocytoma

Endocrine syndromes MEN I and II

von Hippel-Lindau syndrome

Neurofibromatosis type 1

Hereditary pheochromocytoma-paraganglioma syndrome  
(PGL)

Unilateral pheochromocytoma in documented genetic cases

Primary aldosteronism resulting from unilateral adrenal  
adenoma with the opposite normal adrenal gland

Unilateral adrenal adenoma in Cushing syndrome

Pseudocyst or adrenal endothelial cyst



Thanks For Your  
Attention