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A must-buy book for FMGE Entrance Exam



Quick Review of **SURGERY** for NEET and FMGE

Including Latest Exam Pattern Questions, Important Annexures and Image-based Questions

Updated from Bailey and Love 27/E, Sabiston 20/E, Schwartz 11/E, Smith Urology 18/E, Campbell Urology 11/E and Harrison's 20/E

Updated with 8th AJCC (2017)



Pritesh Singh

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Most Important Very Important Important ****

THYROGLOSSAL CYST

THYROGLOSSAL CYST

Thyroid

- Cystic swelling developed in the remnant of the thyroglossal duct or tract
- Present in any part of the thyroglossal tract^Q (thyroglossal tract extends from foramen caecum to isthmus of thyroid)

Common Sites

- Subhyoid (MC)^Q
- Region of the thyroid cartilages
- Floor of mouth
- Region of the thyroid cartilaSuprahyoid
- Beneath the foramen caecum

- Clinical Features
- It is a midline swelling^Q, except in the region of thyroid cartilage, where thyroglossal tract is pushed to one side, usually to the left.
- Though it's a congenital swelling^Q MC age of presentation is between 15 and 30 years^Q.
- Cyst can be **moved sideways** but not vertically.
- · Peculiar characteristic which helps in distinguishing thyroglossal cyst from other neck swelling
- Moves up with protrusion of tongue^Q as the thyroglossal tract is attached to the tongue.
- Moves with deglutition^Q so do all thyroid swellings, subhyoid bursitis.
- Cyst is lined by pseudostratified columnar epithelium and squamous epithelium with heterotopic thyroid tissue present in 20% of cases.

Complications

• Recurrent infections^Q • Formation of thyroglossal fistula^Q • Carcinomatous change (papillary carcinoma^Q)

Treatment

• Sistrunk operation: En-bloc cystectomy and excision of central hyoid bone^Q to minimize recurrence.

RETROSTERNAL GOITER

RETROSTERNAL (SUBSTERNAL/MEDIASTINAL/INTRATHORACIC GOITER)

- A goiter is said to be retrosternal, substernal or mediastinal if > 50% of thyroid tissue is below the opening of thoracic cage^Q.
- Usually arises from lower pole of a nodular goiter^Q.

Clinical Features

- Often symptomless, discovered on a routine chest X-ray^Q.
 - Can lead to tracheal deviation and scabbard trachea^Q (flattening of trachea caused by compression)

Severe Symptoms due to Mass Effect on the Trachea, Esophagus, Great Vessels and Nerves

- 1. Dyspnea (MC symptom) particularly at night, cough and stridor^Q
- 2. Dysphagia
- 3. Enlargement of neck veins and superficial veins on the chest wall
- 4. Recurrent nerve palsy
- 5. **Pemberton's sign**^Q: Symptoms of faintness with evidence of facial congestion and external jugular venous obstruction when the arms are raised above the head.

Treatment

• Virtually all intrathoracic goiters can be removed via a cervical incision^Q.

■ INVESTIGATIONS IN THYROID DISORDERS

THYROID SCAN

- Whereas ultrasound allows anatomic evaluation, radionuclide scans allow assessment of thyroid function^Q.
- 123I and 131I iodine scintigraphy is also used to evaluate the functional status of the gland.
- Advantages of scanning with ¹²³I include a low dose of radiation (30 mrad) and short half-life^Q.
- ¹³¹I has a longer half-life (8 days) and emits higher levels of β-radiation^Q.
- ¹³¹I is optimal for **imaging thyroid carcinoma**. It is the screening modality of choice for the evaluation of distant metastasis.

Isotope	t _{1/2}
1 ¹³²	2.3 hours ^Q
1 ¹²³	13 hours ^o
I ¹³¹	8 days ^q

RADIOACTIVE IODINE (I¹³¹) THERAPY

- I¹³¹ is an effective agent for delivering high radiation doses to thyroid tissue^Q
- It emits mainly beta radiation (90%), which penetrates only 0.5 mm^Q of the tissue & allow therapeutic effects on thyroid without any damage to the surrounding structures, particularly parathyroids.

Mechanism of Action

- **I**¹³¹ emits **beta particles**^Q and *γ*-rays.
- Beta rays are utilized for their destructive effects on thyroid^Q cells.
- X-rays are useful for tracer studies.

1. 2. 3.

Indications in Carcinoma Thyroid		Contraindications of I ¹³¹ Therapy
Distant metastasis ^Q at diagnosis	1.	Childhood ^Q
Incomplete tumor resection ^Q	2.	Pregnancy ^Q
Patients at high risk for mortality or recurrence ^Q	3.	Lactation ^Q

SOLITARY THYROID NODULE

SOLITARY THYROID NODULE

- MC solitary thyroid nodule is benign colloid nodule^Q, it accounts for 60% cases of solitary thyroid nodule.
- 2nd MC cause of solitary thyroid nodule is follicular adenoma (30%)^Q.



ACUTE (SUPPURATIVE) THYROIDITIS

ACUTE (SUPPURATIVE) THYROIDITIS

- Acute thyroiditis is rare and due to **suppurative infection** of the **thyroid**^Q.
- More common in children and often is preceded by an upper respiratory tract infection or otitis media^Q.

Etiology

• Thyroid gland is inherently resistant to infection due to its extensive blood & lymphatic supply, high iodide content, and fibrous capsule.

Infectious Agents Can Seed Thyroid

- 1. Via hematogenous or lymphatic route^Q
- 2. Via direct spread from persistent pyriform sinus fistulae or thyroglossal duct cysts^Q
- 3. As a result of penetrating trauma
- 4. Due to immunosuppression^Q
- MC organism responsible: Staph. aureus >Streptococcus^Q
- In children & young adults, MC cause is presence of a pyriform sinus^Q (remnant of the fourth branchial pouch that connects the oropharynx with thyroid), such sinuses are predominantly left-sided^Q.
- Long-standing goiter and degeneration in thyroid malignancy are risk factors in elderly^Q

Clinical Features

- Thyroid pain, often referred to the throat or ears, and a small, tender goiter^Q
- Fever, dysphagia and erythema over the thyroid.
- Systemic symptoms of a febrile illness and lymphadenopathy^Q.

Diagnosis

- ESR and WBC count are usually increased, but thyroid function is normal.
- FNA biopsy shows infiltration by polymorphonuclear leukocytes.
- Culture of the sample can identify the organism.
- Persistent pyriform sinus fistula should be suspected in children with recurrent acute thyroiditis. A barium swallow demonstrates the anomalous tract with 80% sensitivity^Q.

Treatment

- Parenteral antibiotics and drainage of abscesses^Q.
- Patients with **pyriform sinus fistulae** require **complete resection**^Q of the sinus tract, including the area of the thyroid where the tract terminates, **to prevent recurrence**.

SUBACUTE THYROIDITIS

SUBACUTE /DE QUERVAIN'S/ GRANULOMATOUS/ VIRAL THYROIDITIS/GIANTCELL THYROIDITIS

- Also termed de Quervain's thyroiditis, granulomatous thyroiditis, or viral thyroiditis.
- Peak incidence: **30–50 years**; women are affected three times more frequently than men.
- Usually follows upper respiratory tract infection^Q
- A viral etiology has been proposed
- Strong association with the HLA-B35 haplotype^Q

	The Disorder Classically Progresses through Four Stages
1.	Initial hyperthyroid phase, due to release of thyroid hormone
2.	Euthyroid phase
3.	Hypothyroidism, occurs in about 20 to 30% of patients
4.	Resolution and return to the euthyroid state in 90% of patients.

- In the **early stages** of the disease, **TSH** is **decreased**, and **Tg**, **T4**, and **T**₃ levels are **elevated** due to the release of preformed thyroid hormone from destroyed follicles.
- ESR is typically >100 mm/h^Q.
- **RAIU** is decreased^Q

Clinical Features

- Painful and enlarged thyroid, sometimes accompanied by fever.
- Features of thyrotoxicosis or hypothyroidism, depending on the phase of the illness.

34 Quick Review of Surgery

Contd...

- Malaise and symptoms of an **upper respiratory tract infection** may **precede** the **thyroid-related features**^Q by several weeks.
- The patient typically complains of a sore throat and small exquisitely tender goiter^Q
- Pain is often referred to the jaw or ear.
- Complete resolution is the usual outcome^Q
- Permanent hypothyroidism can occur, particularly in those with coincidental thyroid autoimmunity.

Laboratory Findings

- ESR is markedly elevated^Q
- Antithyroid antibodies are low with T4, T3 and TSH levels depend on the stage of disease.
 - RAIU is decreased during the hyperthyroid stage (distinguishes from Grave's disease)
 In doubt: FNAC (shows characteristic giant multinucleated cells)^Q

Treatment

- Treatment is primarily symptomatic, as disease is self-limited^Q.
- Aspirin or other NSAIDs are sufficient to control symptoms in most cases.
- Severe cases with marked local or systemic symptoms may require glucocorticoids.
- Short-term thyroid replacement may be needed in the hypothyroid phase.
- Thyroidectomy is reserved for the rare patients who have a prolonged course not responsive to medical measures.

RIEDEL'S THYROIDITIS

RIEDEL'S THYROIDITIS

- A rare variant of thyroiditis also known as **Riedel's struma**^Q or **invasive fibrous thyroiditis**
- Characterized by the replacement of all or part of the thyroid parenchyma by fibrous tissue
- Also invades into adjacent tissues^Q.
- Etiology: Primary autoimmune etiology (probably)

Riedel's Thyroiditis is Associated with

- Mediastinal and retroperitoneal fibrosis
- Periorbital and retro-orbital fibrosis
- Sclerosing cholangitis

Clinical Features

- Occurs predominantly in **women**, **30–60 years**.
- Presents as a painless, hard anterior neck mass⁰, which progresses over weeks to years to produce symptoms of compression, including dysphagia, dyspnea, choking, and hoarseness.
- Patients may present with symptoms of hypothyroidism and hypoparathyroidism^Q as the gland is replaced by fibrous tissue.
- Physical examination: Hard, "woody" thyroid gland with fixation^Q to surrounding tissues.

Treatment

- Surgery^Q is the mainstay of the treatment (decompress the trachea by wedge excision of isthmus)
- Some patients show dramatic improvement with tamoxifen & corticosteroids.

HASHIMOTO'S THYROIDITIS

HASHIMOTO'S THYROIDITIS

- First described by Hashimoto as struma lymphomatosa⁰, i.e. a transformation of thyroid tissue to lymphoid tissue.
- MC inflammatory disorder of the thyroid and leading cause of hypothyroidism^Q.
 - Thyroid lymphoma^Q is a rare but well-recognized complication
 - Papillary thyroid carcinoma^Q may be occasionally associated
- Genetic association has been noted with HLA B8, DR3 and DR5^o.
- More common in **women** (Male:female, 1:10), near menopause (**30-50 years**).

Etiopathogenesis

- Autoimmune disease
- Thought to be initiated by activation of CD4+T (helper) lymphocytes which further recruit cytotoxic CD8+T cells.

• Thyroid tissue is destroyed by cytrotoxic T cells and autoantibodies^Q.

Autoantibodies are Directed against

- 1. Thyroglobulin (Tg): 60%
- 2. Thyroid peroxidase (TPO): 95%^Q
- 3. TSH-R: 60%
- It is also thought to be **associated with**:
 - Increased intake of **iodine**
 - Drugs such as interferon alpha, lithium, amiodarone

Pathology

• Gross examination: Mildly enlarged thyroid^Q with pale, gray-tan cut surface

• Microscopic examination:

- Gland is diffusely infiltrated by small lymphocytes and plasma cells^Q
- Follicles are lined by Hürthle or Askanazy cells^Q (characterized by abundant eosinophilic, granular cytoplasm).

Clinical Features

- MC presentation: Minimally or moderately enlarged firm gland^Q.
- On examination an **enlarged pyramidal lobe** is often palpable.
- Mild hyperthyroidism may be present initially (due to destruction of thyroid tissue).
- Hypothyroidism is inevitable and usually permanent^Q.

Laboratory Findings

- Elevated TSH and presence of thyroid autoantibodies confirm the diagnosis^Q.
- Elevated TSH, reduced T4 and T3 levels^Q.
- Presence to thyroid autoantibodies (particularly **TPO antibody**)^Q.
- In case of doubt, diagnosis is confirmed by FNA biopsy.

Management

- Thyroid hormone replacement therapy for overtly hypothyroid patients or in euthyroid patients to shrink large goiters^Q.
- Surgery may occasionally be indicated for suspicion of malignancy or for goiters causing compressive symptoms or cosmetic deformity.

PAINLESS OR SILENT THYROIDITIS

PAINLESS OR SILENT THYROIDITIS

- Painless thyroiditis, or "silent" thyroiditis, occurs in patients with underlying autoimmune thyroid disease and has a clinical course similar to that of subacute thyroiditis.
- Occurs in up to 5% of women 3-6 months after pregnancy termed as postpartum thyroiditis.
- Associated with presence of TPO antibodies antepartum, three times more common in women with type 1 DM.

Clinical Features

• Characterised by brief phase of thyrotoxicosis lasting 2-4 weeks, followed by hypothyroidism for 4-12 weeks, and then resolution.

Diagnosis

- Uptake of ^{99m}Tc pertechnetate or radioactive iodine is initially suppressed.
- In addition to the painless goiter, silent thyroiditis can be distinguished from subacute thyroiditis by a normal ESR & presence of TPO antibodies.

Treatment

- · Glucocorticoid treatment is not indicated for silent thyroiditis.
- Severe thyrotoxic symptoms: Propranolol
- Thyroxine replacement for hypothyroid phase but should be withdrawn after 6–9 months, as recovery is the rule.

GRAVE'S DISEASE (DIFFUSE TOXIC GOITER)

GRAVE'S DISEASE (DIFFUSE TOXIC GOITER)

- MC cause of hyperthyroidism, caused by stimulatory autoantibodies to TSH-R^Q.
- Autoimmune disease with strong familial predisposition^Q.
- More common in **females** with peak incidence between **40–60** years.
- Characterized by **thyrotoxicosis**, **diffuse goiter** & **extrathyroidal conditions**^Q (ophthalmopathy, dermopathy, thyroid acropachy and gynecomastia).

Etiopathogenesis

- Autoimmune process with possible triggers (post-partum state, iodine excess, lithium therapy and bacterial or viral infections)
- Associated with HLA-B8, HLA-DR3, HLA-DQA1*0501 and CTLA-4^Q
- Thyroid stimulating antibodies^Q are hallmark of Grave's disease

Clinical Features

- Hyperthyroid symptoms^Q (heat intolerance, increased sweating and thirst, weight loss despite adequate caloric intake)
- Symptoms of adrenergic stimulation^Q (palpitations, nervousness, fatigue, emotional lability, hyperkinesis and tremors)
- MC GI symptom is increased frequency of bowel movements and diarrhea^Q
 - Female patients often develop amenorrhea, decreased fertility and increased incidence of miscarriage^Q
 - Children experience rapid growth with early bone maturation^Q
 - Older patients present with CVS complications (AF and CHF)^Q
- Weight loss, facial flushing, warm and moist skin, tachycardia, cutaneous vasodilatation, collapsing pulse is seen on examination
- A fine tremor, muscle wasting and proximal muscle group weakness with hyperactive tendon reflexes often are present^Q
 - Overlying bruit or thrill at upper pole^Q due to increased vascularity
 - Loud venous hum^Q in supraclavicular space
 - Ophthalmopathy (orbital proptosis) occurs in 50%, dermopathy in 1–2%.^Q
 - Dermopathy is characterized by deposition of glycosaminoglycans leading to thickened skin in pretibial region and dorsum of foot^o (pretibial myxedema).
- Gynecomastia is common in young men^Q
- Rare bony involvement leads to **subperiosteal bone formation** and **swelling in metacarpals**^Q (thyroid acropachy).

Diagnosis

- Suppressed TSH with or without an elevated free T4 or T3 level. If eye signs are present, other tests are generally not needed^Q.
- In absence of eye signs, elevated RAIU with diffusely enlarged gland^Q confirms the diagnosis.
- Elevated TSH-R or thyroid-stimulating antibodies (TSAb) are diagnostic⁰ of Grave's disease and increased in about 90% patients.
- Anti-Tg and Anti-TPO antibodies are non-specific and elevated in up to 75% cases.
- MRI of orbits are useful in evaluating Grave's ophthalmopathy.

Treatment

• Treatment modalities: Antithyroid drugs, thyroid ablation with radioactive ¹³¹I and thyroidectomy^Q.

TREATMENT OF GRAVE'S DISEASE

TREATMENT OF GRAVE'S DISEASE

• Treatment modalities are antithyroid drugs, thyroid ablation with radioactive ¹³¹I, and thyroidectomy.

Antithyroid Drugs

- Administered in **preparation for RAI ablation** or **surgery**^Q.
- Drugs commonly used: Propylthiouracil and methimazole^Q.
- Propylthiouracil can cause liver failure in pregnancy.
- Methimazole is associated with aplasia cutis & choanal atresia.
 - Antithyroid drug of choice in Graves: Methimazole^Q
 - Antithyroid drug of choice **in pregnancy: Carbimazole**^Q
 - Antithyroid drug of choice in thyroid storm: Propylthiouracil^Q
- Most patients have **improved symptoms** in **2 weeks** and become **euthyroid** in about **6 weeks**^Q.
 - Catecholamine response of thyrotoxicosis can be alleviated by propranolol^Q.

Radioactive lodine Therapy (131)

¹³¹I emits beta (90%) and gamma rays^Q

Indications of RAI Therapy

- 1. Older patients with small or moderate-sized goiters^Q
- 2. Patients relapsed after medical or surgical therapy^Q
- 3. Antithyroid drugs or surgery are contraindicated^Q

	Contraindications of RAI		
	Absolute Contraindications	Relative Contraindications	
•	Pregnancy ^Q	• Young patients (children and adolescents) ^Q	
•	Lactation ^Q	 Thyroid nodules^Q 	
		Ophthalmopathy ^q	

•

Surgical Treatment

- Surgery is recommended when RAI is contraindicated^Q
- Treatment of choice: Total Thyroidectomy

Indications of Surgery	
When RAI is Contraindicated	Relative Indications
 Confirmed cancer or suspicious thyroid nodules^q Young patients^q 	Smokers, with moderate to severe Grave's ophthalmopathy ^q Detinate desiring residence of
 Pregnancy and Lactation^a Severe reactions to antithyroid medications 	 Patients desiring rapid control of hyperthyroidism with a chance of being euthyroid Poor compliance to antithyroid medications
 Large goiters causing compressive symptoms Reluctant to undergo RAI therapy 	· · · · · · · · · · · · · · · · · · ·

TOXIC ADENOMA

TOXIC ADENOMA (PLUMMER'S DISEASE)

- Hyperthyroidism from a single hyperfunctioning nodule^Q typically occurs in younger patients
- Increased thyroid hormone production occurs independent of TSH control^Q.
- Characterized by **somatic mutations** in the **TSH-R gene**^Q

Clinical Features

- Recent growth of a long-standing nodule along with the symptoms of hyperthyroidism^Q.
- Hyperthyroidism from a single hyperfunctioning nodule typically occurs in younger patients
- Physical examination: Solitary thyroid nodule without palpable thyroid tissue on the contralateral side.
- Eye signs are not common, mainly **CVS dysfunction**
- Rarely malignant^Q.

Diagnosis

• RAI scanning shows a "hot" nodule^Q with suppression the rest of the thyroid gland.

Treatment

- Smaller nodules may be managed with antithyroid medications and RAI^Q.
- Most patients are euthyroid after radioiodine therapy^Q (radioiodine preferentially accumulates in hyperfunctioning nodules)
- Surgery (Hemithyroidectomy) is preferred in young patients with larger nodules^Q.

THYROTOXICOSIS

CVS FINDINGS IN THYROTOXICOSIS

- MC cardiovascular manifestation is sinus tachycardia^Q, often associated with palpitations, occasionally caused by supraventricular tachycardia^Q.
- Exertional dyspnea^Q
- Hyperactive precordium with **loud first heart sound**, an accentuated pulmonic component of the second heart sound, and a **thirds heart sound**^Q.
- Systolic ejection click^Q
- The high cardiac output produces a **bounding pulse**, **widened pulse pressure**^Q, and an **aortic systolic murmur** and can lead to worsening of angina or heart failure in the elderly or those with preexisting heart disease.
- Atrial fibrillation is more common in patients >50 years of age^Q.
- A systolic scratch, also known as Means-Lerman scratch^Q, is occasionally heard in 2nd left intercoastal space during expiration.
- Systolic hypertension

Cardiovascular Manifestations of Thyrotoxicosis			
Increased Atrial Irritability	High Cardiac Output		
 Sinus tachycardia (MC)^Q 	Bounding pulse		
 Palpitations^Q 	 Wide pulse pressure^Q 		
 Supraventricular tachycardia^Q 	 Hyperdynamic precordium^Q 		
Atrial fibrillations ^q	 Loud first heart sound^Q, an accentuated pulmonic component of the second heart sound, and a thirds heart sound^Q. 		
	 Aortic systolic murmur^Q 		
	Means-Lerman scratch ^Q		

THYROID STORM (THYROTOXIC CRISIS)

• It is an emergency due to decompensated hyperthyroidism^Q.

Treatment

_

- Non-selective beta blocker (Propranolol):
 - Most valuable measure in thyroid storm^Q.
 - In thyroid storm most of the symptoms are because of adrenergic over activity due to increased tissue sensitivity to catecholamines in hyperthyroidism.
 - This increased sensitivity is due to increased number of beta receptors^Q.
- Quick relief is obtained by blocking beta receptors.
- **Propylthiouracil:**
 - Antithyroid drug of choice for thyroid storm^Q
 - **Reduces hormone synthesis** as well as peripheral conversion of T_4 to T_3^Q _
- **Corticosteroids (Hydrocortisone):**
 - Inhibits both release of thyroid hormone from the gland and peripheral conversion of T, to T, o
- Iodides (Potassium iodide or ipanoic acid):
- Used to **inhibit** further **hormone release**^Q from the gland.
- **Other Measures:**
 - Diltiazem, if tachycardia is not controlled by propranolol alone.
 - Rehydration, anxiolytics, external cooling and appropriate antibiotics

CARCINOMA THYROID

Type of Thyroid Carcinoma	Prevalence
Papillary (MC)	80–90%
Follicular	5–10%
Medullary	10%
Anaplastic	Rare
Lymphoma	1%

Well Differentiated Thyroid Cancer

1. Papillary carcinoma of thyroid^Q 2. Follicular carcinoma of thyroid^Q

3. Follicular variant of papillary carcinoma thyroid^a 4. Hurthle cell carcinoma (variant of follicular carcinoma thyroid)^a

Carcinoma Thyroid		
Туре	Mode of spread	MC site of metastasis
Papillary carcinoma	Lymphatic ^q spread	Lungs
Follicular carcinoma	Hematogenous ^q spread	Bones
Medullary carcinoma	Both lymphatic and hematogenous ^Q spread	Liver
Anaplastic carcinoma	Direct invasion ^q Lungs	
Pulsating Secondaries		

Follicular carcinoma thyroid^Q 2. RCC^Q

Bone Metastasis in Carcinoma Thyroid		Bone Metastasis in Carcinoma Thyroid	
	Follicular carcinoma	Osteolytic metastasis (Pulsating secondaries in flat bones) ^Q	
	Medullary carcinoma	Osteoblastic metastasis ^q	

PAPILLARY CARCINOMA OF THYROID

PAPILLARY CARCINOMA OF THYROID

- Accounts for 80% of all thyroid malignancies in iodine-sufficient areas^Q
- MC thyroid cancer in children & individuals exposed to external radiation^Q. •
- More often in women, 30-40 years.

Pathology

- Grossly: Hard & whitish remain flat on sectioning with a blade with macroscopic calcification, necrosis, or cystic changes
 - Multifocality^Q is common (up to 85% of cases) on microscopic examination.
 - Multifocality is associated with an increased risk of cervical nodal metastases^Q, rarely
 - invade adjacent structures such as the trachea, esophagus & RLNs.
- Rarely encapsulated^Q (PCT are seldom encapsulated)
- Other variants: Tall cell⁰, insular⁰, columnar, diffuse sclerosing, clear cell, trabecular, and poorly differentiated types; account for about 1%; associated with a worse prognosis.

Histological Characteristics of Papillary Carcinoma Thyroid

- Papillary projections^q: PTC contains branching papillae of cuboidal epithelial cells
- Orphan Annie eye nuclei:
 - The nuclei contain finely dispersed chromatin, which imparts an optically clear or empty appearance, giving rise to term ground glass or Orphan Annie eye nuclei^Q.
 - Invaginations of cytoplasm in cross-sections: Intranuclear inclusions^Q (pseudo-inclusion) or intranuclear grooves^Q.
 - Diagnosis of PTC is based on these nuclear characteristics^q even in the absence of papillary structures.
 - Psammoma bodieso: Microscopic, calcified deposits representing clumps of sloughed cells

Clinical Features

- Most patients are euthyroid & present with a slow-growing painless mass^Q in the neck.
- Dysphagia, dyspnea dysphonia are associated with locally advanced invasive disease.
- Lymph node metastases are common^Q, especially in children young adults, and may be the presenting complaint.

• "Lateral aberrant thyroid" denotes a cervical lymph node that has been invaded by metastatic cancer^Q.

- Distant metastases are uncommon at initial presentation, but may ultimately develop in up to 20% of patients.
- MC sites of metastasis: Lungs^Q >bone >liver >brain.

Diagnosis

- Diagnosis is established by FNAC of the thyroid mass or lymph node^Q.
- Once thyroid cancer is diagnosed on FNAC, a **complete neck ultrasound** to evaluate the **contralateral lobe** and for **LN metastases** in the central & lateral neck compartments.

Treatment

- Total or near-total thyroidectomy^Q
- During thyroidectomy, enlarged central neck nodes should be removed^Q.
- Biopsy-proven lymph node metastases detected clinically or by imaging in the lateral neck in patients with papillary carcinoma are managed with modified radical neck dissection.

Prognosis

• PTC have an excellent prognosis with a >95% 10-year survival rate^Q.

FOLLICULAR CARCINOMA THYROID

FOLLICULAR CARCINOMA OF THYROID

- FTC account for 10% of thyroid cancers
- Occurs more commonly in **iodine-deficient areas**^Q.
- More common in women with mean age of 50 years
- Genes implicated in FCT: p53^Q, PTEN^Q, Ras^Q, PAX8/PPAR1

Pathology

- Usually **solitary lesion** surrounded by **capsule**^Q.
- Malignancy is defined by the presence of capsular and vascular invasion^Q.
- Tumor infiltration and invasion, as well as tumor thrombus within the middle thyroid or jugular veins, may be apparent at operation.

Clinical Features

- Usually present as solitary thyroid nodules, occasionally with a history of rapid size increase, and long-standing goiter^Q.
- Preoperative clinical diagnosis of cancer is difficult unless distant metastases are present.
- Large follicular tumors (>4 cm) in older men are more likely to be malignant^Q.

• MC site of metastasis is bone (Osteolytic metastasis with pulsating secondaries in flat bones)^Q

• MC site of metastasis: Vertebra^Q >Ribs >Pelvis Bones >Skull

Diagnosis

- FNAC is unable to distinguish benign follicular lesions from follicular carcinomas^Q.
- Intraoperative frozen-section examination usually is not helpful, but should be performed when there is evidence of capsular or vascular invasion, or when adjacent lymphadenopathy is present.

Treatment

- Follicular lesion: Hemithyroidectomy^Q (80% of these patients will have benign adenomas)
- Thyroid cancer: Total thyroidectomy^Q
- Prophylactic nodal dissection is unwarranted^Q because nodal involvement is infrequent.

Prognosis

• Most important prognostic factor: Age and distant metastasis.

	Poor Long-term Prognosis		
•	Age > 50 years ⁰ Tumor size > 4 cm ⁰ Higher tumor grade⁰	 Marked vascular invasion^a Extrathyroidal invasion^a Distant metastases^a 	

MEDULLARY CARCINOMA THYROID

MEDULLARY CARCINOMA THYROID

- Neuroendocrine carcinoma **arising from parafollicular 'C' cells**^Q of thyroid
- Parafollicular 'C' cells are derived from the ultimobranchial bodies^Q & secrete calcitonin^Q
- 'C' Cells are concentrated superolaterally in thyroid lobes, from where MTC usually develops
- Most MTCs (75–80%) arise sporadically^Q
- Spread is both lymphatic & hematogenous^Q
- MC site of metastasis: Liver^Q

Medullary Carcinoma Thyroid		
Sporadic: 80% ^o	Familial: 20% ^Q	
	(Non-MEN setting/ MEN-2A/MEN-2B)	
 Originate in one lobe^Q 	Multicentric and bilateral ^Q	
Seen in 6th decade	 Occur in younger age^Q 	
• RET protoncogene ^Q mutation	 Associted with C-cell hyperplasia^Q 	
	 RET protoncogene^a mutation 	

Clinical Features

Medullary Carcinoma Should Be Suspected

- High level of serum Calcitonin^Q & CEA^Q
- · Cervical lymph nodes at time of presentation (LN involvement, thyroid and blood borne metastases occurs early)^q
- Diarrhea^Q at the time of presentation.
- Amyloid ^Q in stroma histologically.
- · MEN setting: Evidence of Pheochromocytoma/Hyperparathyroidism/Thyroid cancer in family.
- (Discovery of medullary carcinoma thyroid makes family surveillance advisable)^Q

Diagnosis

- Diagnosed by FNAC^Q
- I¹³¹ scan is of no use as MTC is TSH independent^Q.
- Tumor marker: Calcitonin is raised in almost all cases of MTC
- Calcitonin excess in MTC is not associated with hypocalcemia

Treatment

- Total thyroidectomy + Central LN dissection ± Ipsilateral MRND if tumor >1 cm^Q
- If nodes are positive on ipsilateral side: Bilateral MRND
- Vandetanib (EGFR inhibitor) is the only drug approved by US FDA for treatment of advanced & progressive MTC

Follow-up

• Level of Calcitonin falls after resection and is raises again in cases of recurrence, used for follow up^Q.

Prognosis

• MTC is associated with poor prognosis^Q.

ANAPLASTIC CARCINOMA

ANAPLASTIC CARCINOMA

- Accounts for **1%** of all thyroid malignancies
- Mainly affect women in 7th and 8th decade^Q
- The typical patient has a long-standing neck mass, which rapidly enlarges and may be painful^Q.
- Most aggressive form of thyroid cancer^Q

Pathology

- Grossly: Firm & whitish in appearance.
- Microscopically, sheets of cells with marked heterogeneity & characteristic giant & multinucleated cells^Q.

Clinical Features

- Typical manifestation: An older patient with dysphagia, cervical tenderness & a painful, rapidly enlarging neck mass^Q
- Superior vena cava syndrome can also be part of the findings.
- The clinical situation deteriorates rapidly into tracheal obstruction & rapid local invasion^Q of surrounding structures.
- Associated symptoms: Dysphonia, dysphagia & dyspnea
 - Lymph nodes usually are palpable at presentation.
 - Evidence of metastatic spread also may be present.
 - MC site of metastasis: Lungs^Q

Diagnosis

- Confirmed by FNAC revealing characteristic giant & multinucleated cells^Q.
- Incisional biopsy occasionally is needed to confirm the diagnosis

Treatment

- Thyroidectomy for resectable mass^Q (may lead to a small improvement in survival, especially in younger individuals)
- Combined radiation & chemotherapy in an adjuvant setting in patients with resectable disease has been associated with prolonged survival^Q.

Prognosis

• Most aggressive thyroid malignancies^Q, with <6 months survival

METASTATIC TUMORS OF THYROID

METASTATIC TUMORS OF THYROID

- Rare, most cases are found in autopsy
- MC site of primary: CA Breast^Q > CA Lung
- If thyroid metastases is detected pre-mortem, MC site of primary: RCC^Q > CA Breast > CA Lung

THYROID LYMPHOMA

THYROID LYMPHOMA

- MC type is NHL B cell^Q type, of intermediate grade.
- Majority of patients have thyroid disease plus cervical or mediastinal lymph nodes^Q.
- More common in **females**.
- Most thyroid lymphomas develop in patients with Chronic Lymphocytic Thyroiditis^Q

Clinical Features

- Lymphomas are rapidly growing tumours, present with rapidly enlarging neck mass which is often painless.
- Patients may present with acute respiratory distress & dysphagia
- About 10–30% present with symptoms relating to local invasion, including hoarseness, dyspnoea with stridor, or dysphagia.
 - Painless^Q and associated with fever^Q
 - Patients with thyroid lymphoma virtually never have hyperthyroidism but frequently have hypothyroidism^Q.
 - Hypothyroid patients have evidence of autoimmune thyroiditis or Hashimoto's thyroiditis^Q.

Diagnosis

• Diagnosis is confirmed by **core-needle biopsy**^Q.

Treatment: External Beam Radiotherapy + Chemotherapy^Q

- Patients with **thyroid lymphoma** respond rapidly to chemotherapy (**CHOP**—cyclophosphamide, doxorubicin, vincristine, and prednisone) and associated with **improved survival**.
- Combined treatment with radiotherapy & chemotherapy often is recommended.
- To alleviate pressure symptoms, surgical resection (Thyroidectomy and nodal dissection) is recommended.

THYROIDECTOMY

STEPS OF THYROIDECTOMY

- A Kocher transverse collar incision, typically 4 to 5 cm in length, is placed in or parallel to a natural skin crease 1 cm below cricoid cartilage.
- Subcutaneous tissues & platysma are incised sharply and subplatysmal **flaps are raised superiorly** to the level of **thyroid cartilage** and **inferiorly** to the **suprasternal notch**^Q
- Strap muscles are divided in the midline along the entire length of mobilized flaps, & thyroid gland is exposed.
- Middle thyroid veins are ligated and divided^Q.
 - Dissection plane is kept as close to the thyroid as possible & superior pole vessels are individually identified, skeletonized, ligated, & divided low on the thyroid gland to avoid injury to external branch of superior laryngeal nerve^Q.
- RLNs can be most consistently identified at the level of cricoid cartilage.
- Parathyroids usually can be identified within 1 cm of the crossing of the inferior thyroid artery and the RLN^Q
 - Inferior thyroid vessels are dissected, skeletonized, ligated, and divided as close to the surface of thyroid gland as possible to minimize devascularization of the parathyroids (extracapsular^Q dissection) or injury to the RLN^Q.
- RLN is most vulnerable to injury in the vicinity of the ligament of Berry. Any bleeding in this area should be controlled with gentle pressure before carefully identifying the vessel & ligating it. Use of the electrocautery should be avoided in proximity to the RLN^Q.
- Once the ligament is divided, the thyroid can be separated from the underlying trachea by sharp dissection.

• **Parathyroid glands** that have been **inadvertently removed** during the thyroidectomy should be resected, confirmed as parathyroid tissue by frozen section, **divided into 1-mm fragments**, and **reimplanted into** individual pockets in the **sternocleidomastoid**^Q **muscle**. The sites should be **marked with silk sutures** and a **clip**^Q.

COMPLICATIONS OF THYROIDECTOMY

COMPLICATIONS OF THYROIDECTOMY

• Hemorrhage

- Due to slipping of ligature on the superior thyroid artery^Q, bleeding from muscular artery
- Hematomas may cause airway compromise and must be evacuated immediately^Q.
- Hematomas may occur immediately or later on.
- An immediate bleed occurs after or shortly before extubation when the patient lightens from anaesthesia and may begin to cough, causing a vessel to open.
- **Respiratory obstruction: Causes include**
- Tension hematoma^Q
- Laryngeal edema (by anesthetic intubation): MC cause of respiratory obstruction^Q
- Bilateral recurrent laryngeal nerve paralysis^Q
- Recurrent laryngeal nerve paralysis
 - May be unilateral or bilateral, transient or permanent.
 - Bilateral paralysis causes respiratory obstruction Dyspnea, stridor.
- Injury to other nerves
 - External branches of superior laryngeal nerve^Q (MC injured nerve during thyroid surgery: External laryngeal nerve^Q)
 - Cervical sympathetic trunk may cause Horner's syndrome.
- Parathyroid insufficiency
 - Due to removal of the parathyroid glands or infarction due to vascular injury^Q.
 - **Vascular injury**^Q is more important.
 - Cases usually present 2–5 days after operation^Q with symptoms of hypocalcemia (circumoral and fingertip numbness and tingling tetany, carpopedal spasm and laryngeal stridor)^Q
 - Treatment with oral calcium & vitamin D supplements^Q
 - **IV calcium gluconate**^Q may be required in severe cases.
- Thyroid insufficiency
- Thyrotoxic crisis
 - Occurs if the thyrotoxic patient has been inadequately prepared for thyroidectomy^Q.

Multiple Choice Questions

PAPILLARY CARCINOMA

- 1. Psammoma bodies may be seen in all of the following, (Recent Question 2016, All India 2011) except:
 - a. Follicular carcinoma of thyroid
 - b. Papillary carcinoma of thyroid
 - Meningioma C.
 - d. Serous cystadenocarcinoma of ovary
- 2. Most common thyroid malignancy is:

(DNB 2012, MHPGMCET 2002)

(Orissa 2011)

- b. Follicular carcinoma Anaplastic carcinoma a.
- d. Papillary carcinoma c. Medullary carcinoma
- 3. Which thyroid malignancy is common after radiation exposure? (Recent Question 2016, MHSSMCET 2005)
 - a. Follicular b. Papillary
 - c. Medullary d. Anaplastic
- 4. Orphan Annie-eye nuclei seen in:
 - a. Papillary carcinoma of thyroid
 - b. Medullary carcinoma of thyroid
 - Anaplastic carcinoma of thyroid c.
 - Follicular carcinoma of thyroid d.
- 5. Which of the following would be the best treatment for a 2 cm thyroid nodule in a 50 years old man with FNAC revealing it to be a papillary carcinoma?
 - (All India 2009, Recent Question 2015)
 - Hemithyroidectomy a.
 - b. Subtotal thyroidectomy with modified neck dissection
 - Near total thyroidectomy with modified neck dissection c.
 - Hemithyroidectomy with modified neck dissection d.
- 6. True regarding papillary carcinoma of thyroid:
 - Undifferentiated carcinoma (MCI March 2006) a. Blood-borne metastasis is commoner b.
 - Excellent prognosis c.
 - Capsulated d.

a.

c.

- 7. Which type of thyroid carcinoma has the best prognosis?
 - a. Papillary carcinoma
- b. Anaplastic carcinoma

(DNB 2010, All India 96)

- Follicular carcinoma d. Medullary carcinoma C. 8. Occult thyroid malignancy with nodal metastasis is:
- (DNB 2005, 2001, AIIMS Sept 96)
 - Medullary carcinoma b. Follicular carcinoma
 - Papillary carcinoma
 - d. Anaplastic carcinoma
- 9. A 21 years old woman has 3 cm node in the lower deep cervical chain on the left. The biopsy is interpreted as revealing normal thyroid tissue in a lymph node. The most likely diagnosis is: (DNB 2012, DPG 2009 Feb) Subacute thyroiditis a.

 - b. Metastatic carcinoma thyroid
- C. Hashimoto's disease d. Lateral aberrant thyroid 10. Most common type of carcinoma thyroid having least
 - chances of hematogenous spread: (Recent Question 2015) b. Papillary a.
 - Follicular Anaplastic
 - d. Medullary
- 11. Orphan-Annie eve nuclei is seen in: (Recent Question 2017) a. Papillary carcinoma thyroid
 - b. Follicular carcinoma thyroid
 - c. Medullary carcinoma thyroid
 - d. Anaplastic carcinoma thyroid

- 12. Psammoma bodies are seen in: a. Papillary carcinoma thyroid
 - b. Follicular carcinoma thyroid
 - C. Medullary carcinoma thyroid
 - d. Thyroid lymphoma

C.

- 13. A 27 years old woman presents with 26 weeks of gestation with a lesion, which is found to be papillary carcinoma of thyroid. Which is the best treatment for this patient?
 - Thyroid ablation using radioactive iodine a. b.
 - (MCI June 2019) Total thyroidectomy Hemi-thyroidectomy
 - Observation

FOLLICULAR CARCINOMA

- 14. All of the following are true for follicular carcinoma of thyroid except: (COMEDK 2006) Lymph node involvement rare a.
 - Vascular involvement common b.
 - Younger patients have good prognosis c.
 - d. Diagnosis by FNAC
- 15. Thyroid carcinoma with pulsating vascular skeletal (COMEDK 2007, All India 95) metastasis is: Follicular b. Anaplastic а.
 - c. Medullary d. Papillary
- FNAC is useful in all the following types of thyroid 16. carcinoma except: (UPPG 2010, MCI March 2005, All India 95)
 - b. Follicular a. Papillary Anaplastic c.
 - d. Medullary
- 17. Most probable malignancy that develops in a case of longstanding goiter is: (MCI June 2018, Recent Question 2015, Kerala PG 2015, AIIMS Feb 97, Nov 2001)
 - Follicular carcinoma b. Anaplastic carcinoma a.
 - c. Papillary carcinoma d. Medullary carcinoma
- Bone metastasis is common in which thyroid tumor: 18. Follicular b. Papillary (AIIMS Nov 99) a. Hurthle cell tumour d. Anaplastic c.
- 19. Metastasis from follicular carcinoma should be treated by:
 - Radioiodine b. Surgery (MCI Sept 2006) a. d. Observation Thyroxine c.
- 20. True regarding follicular carcinoma of thyroid: a. Hematogemous spread (JIPMER 2014, 2013)
 - b. Commonly multifocal
 - Readily diagnosed by face c.
 - d. Most commonly carcinoma of thyroid
- 21. FNAC cannot detect which of the following?
 - (AIIMS Nov 2014) b. Papillary carcinoma a. Follicular carcinoma Colloid goiter Hashimoto's thyroiditis c. d.
- MEDULLARY CARCINOMA
 - 22. Screening method of medullary carcinoma thyroid is:
 - Serum calcitonin b. Serum calcium а
 - Serum alkaline phosphate (All India 97, AIIMS Nov 95) c.
 - d. Serum acid phosphatase
 - 23. Treatment of medullary carcinoma thyroid:
 - a. Surgery and Radiotherapy (AIIMS May 2011)
 - Radiotherapy and Chemotherapy b.
 - Surgery only c.
 - Radioiodine ablation d.

(Recent Question 2017)

Explanations

PAPILLARY CARCINOMA

1. Ans. a. Follicular carcinoma of thyroid (Ref: Schwartz 11/e p1647, 10/e p1542; Sabiston 20/e p902; Bailey 27/e p818)

	Psammoma Bodies (PSM)		
	1. Papillary carcinoma thyroid ^Q	2. Papillary carcinoma (RCC) ^Q	
	3. Serous cystadenoma ^Q	4. Meningioma ^Q	
y (carcinoma	3. Ans. b. Papillary	

- 2. Ans. d. Papillary carcinoma
- 4. Ans. a. Papillary carcinoma of thyroid
- 6. Ans. c. Excellent prognosis
- 8. Ans. c. Papillary carcinoma
- 9. Ans. d. Lateral aberrant thyroid
- 10. Ans. b. Papillary
- 11. Ans. a. Papillary carcinoma thyroid (Ref: Schwartz 11/e p1647, 10/e p1542; Sabiston 20/e p903)
- 12. Ans. a. Papillary carcinoma thyroid (Ref: Schwartz 11/e p1647, 10/e p1542; Sabiston 20/e p903).
- 13. Ans. b. Total thyroidectomy (Ref: Schwartz 11/e p1647; Sabiston 20/e p903)

FOLLICULAR CARCINOMA

14. Ans. d. Diagnosis by FNAC (Fef: Schwartz 11/e p1650, 10/e p1544, 1357; Sabiston 20/e p906; Bailey 27/e p818)

LIMITATIONS OF FNAC IN THYROID DISEASES

- 1. Not able to distinguish follicular adenoma from follicular carcinoma^Q
- 2. Not able to distinguish Hurthle cell adenoma from Hurthle cell carcinoma^Q
- 3. Useless in Reidel's thyroiditis^Q (Biopsy is preferred)^Q
- 4. FNAC is less reliable in patients who have history of head and neck irradiation or family history of thyroid cancer due to higher likelihood of multifocal lesions and occult cancer^Q
- 15. Ans. a. Follicular
- 17. Ans. a. Follicular carcinoma
- 19. Ans. a. Radioiodine
- 21. Ans. a. Follicular carcinoma

MEDULLARY CARCINOMA

- 22. Ans. a. Serum calcitonin (Ref: Schwartz 11/e p1655-1656, 10/e p1549-1550; Sabiston 20/e p909; Bailey 27/e p820; Harrison 20/e p2716)
- 23. Ans. c. Surgery only (Ref: Schwartz 11/e p1656, 10/e p1550; Sabiston 20/e p909; Bailey 27/e p820)
- 24. Ans. a. 5 years
- 25. Ans. a. Calcitonin (Ref: Schwartz 11/e p1656, 10/e p1549; Sabiston 20/e p909; Bailey 27/e p820)
- 26. Ans. c. CEA
- 28. Ans. b. Medullary carcinoma
- 29. Ans. c. Medullary carcinoma
- 30. Ans. c. It is dependent on TSH
- 31. Ans. a. Medullary carcinoma thyroid (Ref: Schwartz 11/e p1656, 10/e p1550; Sabiston 20/e p909; Bailey 27/e p820)
- 32. Ans. c. Medullary carcinoma thyroid (Ref: Schwartz 11/e p1656, 10/e p1550; Sabiston 20/e p909; Bailey 27/e p820)
- 33. Ans. a. Medullary carcinoma thyroid (Ref: Schwartz 11/e p1656, 10/e p1541; Sabiston 20/e p909)

ANAPLASTIC CARCINOMA

- 34. Ans. d. Anaplastic carcinoma
- 35. Ans. d. Anaplastic

- 16. Ans. b. Follicular
- 18. Ans. a. Follicular
- 20. Ans. a. Hematogenous spread

7. Ans. a. Papillary carcinoma

5. Ans. c. Near total thyroidectomy with modified neck dissection

27. Ans. c. Ret proto-oncogene

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Quick Review of SURGERY

Salient Features

- Synopsis added before questions to build concepts and to save precious time
- · New NBE based pattern (wider coverage, concept development, one-liner approach)
- · Image-based Questions with Answers (According to Recent Examinations) have been added
- Solved MCQs (PGMEEs 2019–1985) including All Recent Questions (2019–2013)
- · Thoroughly verified answers from MCh aspirants and residents
- Explanations in tabulated form
- · Explanations incorporating only relevant and high yielding facts
- · Highlighted important and golden facts
- Explanations from most authentic surgery books
- · Repeated questions grouped together to save your precious time
- · Line diagrams to minimize tedious efforts
- · Mnemonics for faster learning
- Controversial questions have been handled with special care.

This book is an attempt to provide you with the thorough knowledge of the subject but with a viewpoint that the most important thing is to clear the exam as the competition is intense. There is a difference between Just preparing for an exam, and preparing to achieve the topmost score. So, the book has covered all the topics but with special emphasis on the important and high yielding facts to minimize your tedious efforts and to save your precious time. Remember that without proper revision it is of no use to cover huge amount of syllabus. Last but not least, the surest way not to fail is to determine to succeed. All the best...

Dr Pritesh Singh Director Br Pritesh Institute MBBS (MAMC), MS (Surgery), FMAS, FIAGES

Dr Pritesh Singh, graduate from Maulana Azad Medical College and postgraduate from Lady Hardinge Medical College, New Delhi, India, is an excellent teacher and has been taking awe inspiring classes in various countries since 2009. He is amongst the best faculty and is very popular with students because of his spellbinding classes. He is a renowned educationist and author of Surgery Essence, which needs no introduction and AlIMS Essence and DPG entrance examination books. The students all over the country admire the way he teaches. He is not just a source of inspiration for his pupils rather he is their role model, as he is young and dynamic. He sets a positive example with his style of teaching, courtesy, cooperation and professionalism. Some students say he is a magician who keeps his students spellbound throughout his class. His performance speaks volumes about his knowledge and precision.

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