

1) Anterior neck lump moving with deglutition & tongue protrusion

	Suggested by	Confirmed by	Initial management
1- Ectopic thyroid tissue	Solid lump at any point of the course thyroglossal tract.	- US scan shows non-cystic lesion, (mainly) - radioisotope scan : nodule taking up iodine CT scan, histology of excised tissue.	It's the only source of thyroid tissue in body, so managed conservative : - Medical TTT by L-thyroxine → to decrease its size
2- Thyroglossal cyst	Cystic lump , fluctuant, in midline or just to the left, (commonly subhyoid in midline)	- US scan shows cystic lesion, (mainly) - radioisotope scan : cyst is cold CT scan, histology of excised tissue.	Surgical management : "sistrunk op." Excision of the cyst, track and central part of hyoid bone (due to its different relation with it, can't differentiate above, center or below it)

- Very important to differentiate between both types, because each has a different management

- Both are deep to the fascia, so their consistency is similar (firm) → diagnosed by **U/S**.

2) Neck lump moving with deglutition but not with tongue protrusion → Thyroid swelling

1- Ask about "**toxic manifestations**".

2- If no toxic manifestations → ask about "**Symptoms suggesting malignant invasion**".

3- If no → the patient is non-toxic, non-malignant → **Simple goiter**: ask about "**Pressure symptoms – Dysphagia & Dyspnea**":

- Huge goiter → occurs in *colloid Goiter & Simple nodular Goiter*.
- Retrosternal goiter.
- Certain malignant types: *Anaplastic tumors*.



goitre	Suggested by	Confirmed by	Initial management
1- Toxic goiter			
Graves's disease	- Young age , Clinical thyrotoxicosis → in 100% of patients Ophthalmopathy "Exophthalmos" → in 50% Pretibial myxoedema → in 1% No nodules.	- ↑FT4 or ↑FT3 & ↓TSH - TSH receptor antibody +ve. - U/S → Diffuse gland enlargement - Isotope scan → Diffusely increased uptake	Propranolol 40 to 80mg 8 hourly to control symptoms. Carbimazole 40mg od reduced to 5-10mg over 1-3mo with monthly TFT. FBC before starting. Written warning for agranulocytosis causing sore throat. Radioiodine or thyroidectomy offered if relapse after 6-18mo carbimazole.
Toxic multinodular goitre	- Old age , with a history of previous nodular goitre, multiple nodules and clinically thyrotoxic.	↑FT4 or ↑FT3 & ↓TSH and - U/S → Multiple Nodules. - Isotope scan → increase uptake from nodules themselves while other cold, or from paranodular tissue or combination of both.	carbimazole (± β-blocker for symptoms). Radioiodine very effective (not used if compression of adjacent structures in the neck and thoracic inlet—surgery offered instead).
Toxic Adenoma (Solitary Toxic nodule)	- At any age , - Solitary hyperactive autonomous nodule	↑FT4 or ↑FT3 & ↓TSH - U/S → Solitary nodule. - Isotope scan → single hot nodule.	
2- Malignant goiter	Symptoms suggesting malignant invasion : 1- Invasion of recurrent laryngeal nerve → hoarseness of voice. 2- Invasion of superior laryngeal nerve → choking & hoarseness of voice. 3- Invasion of the vagus nerve → painful ear. 4- Invasion of sympathetic chain → Horner's syndrome. - <i>No dysphagia or dyspnea because tracheal rings resist malignant invasion & underlying esophagus is protected by these rings.</i> - <i>No invasion of carotid sheath, it just push it backward.</i> 5- No dyspnea or dysphagia except in Anaplastic tumor .	- Normal thyroid function - U/S → cystic or solid. - Isotope scanning shows cold nodule - Biopsy	
3- Simple goiter			
<u>Simple diffuse goiter</u> 1- Endemic goiter 2- Physiological goiter 3- Dys-hormonogenesis 4- Sporadic goiter	Not nodular , clinically euthyroid Physiological goiter : - Most common goiter. - Age: Young female (15 – 20) at puberty or during pregnancy & lactation.	FT4 & FT3 normal, TSH normal thyroid antibodies -ve.	Reassurance, no treatment.
<u>Simple nodular goiter</u>	Multiple nodules , clinically euthyroid. The commonest disease of thyroid gland Age: 30 – 40 years. Sex: female > male.	FT4 & FT3 normal, TSH normal. Nodules on US scan or thyroid isotope scan .	Surgery only if indications: 1- Cosmetic 2- Pressure symptoms 3- Suspicion of malignancy.
4- Retrosternal goiter	Small swelling in closed space, causing pressure symptoms: (dyspnea & dysphagia)	CT scan CXR → shadow in sup. Mediastinum	Surgical excision

3) Neck lump doesn't move with deglutition nor tongue protrusion → Other neck swellings

Parotid region swellings

A- Localized :

<u>From skin & subcutaneous tissue:</u>		- Sebaceous cyst - Subcutaneous abscess - Lypoma - hemangioma - lymphangioma	
	<u>From margins :</u>	- Masseter hypertrophy - Zygomatic tumor - Mastoiditis	
<u>LN's :</u>	Multiple	- Pre-auricular LNs - Parotid LNs - Buccinator LNs	<i>LN is diagnosed by two items :</i> 1- Anatomical site 2- Multiplicity (but can be single) . - Both are deep to parotid fascia (which is strong deep fascia) . - To differentiate between both by either : 1- U/S , is there line of cleavage ? If yes → LNs If no → Parotid neoplasm. 2- CT , is there line of cleavage ? We ask for images because the least biopsy in parotid gland swelling is superficial parotidectomy because of branches of facial nerve ! (it might be just LN).
	Single	- Pre-auricular LNs or - Parotid gland neoplasm	

If Parotid gland neoplasm

Pathological differentiation of parotid neoplasm : (can't be assessed by clinician).

A) Benign :

- 1- Pleomorphic adenoma (most common 85%) → It's pleomorphic adenoma till proven otherwise.
- 2- Monomorphic adenoma "Adenolymphoma" , "Warthin's Tumor" , " papillary cystadenoma lymphomatosum"

B) Malignant :

- 1- Adenocarcinoma on top of pleomorphic adenoma (most common malignancy) .
- 2- Adenocarcinoma from the start
- 3- Adenoid cystic carcinoma
- 4- Acinic cell tumor.
- 5- Epidermoid carcinoma
- 6- Mucoepidermoid carcinoma



Clinical estimation of type of tumor :

- If old male , with history of remission & exacerbation → think of "adenolymphoma"



Ask for technetium scan → Hot spot → adenolymphoma (only hot spot tumor)
Cold spot → other benign & malignant tumors

TTT of adenolymphoma: The only tumor will be treated by evacuation (not by superficial parotidectomy), because it's very localized tumor.

- If history of **pain** before swelling because tumor spread along sheaths of facial nerve branches → think about "adenoid cystic carcinoma" → Ask for CT or MRI (not felt because parotid is covered by very dense fascia).

The pathognomonic signs are **early** in adenoid cystic carcinoma because the tumor spread along myelin sheaths of facial nerve branches → so if you neglected the pain of the patient and didn't diagnose the neoplasm , the patient might come early with facial nerve palsy.

The pathognomonic signs of parotid malignancy are late signs:

- 1- Facial nerve palsy
- 2- Fixity of mandible

B- Diffuse :

Diffuse Parotid swellings

Acute

Obstructive :

- 1- Stone
- 2- Stricture

- C/P : colicky facial pain
- Can't be differentiated clinically so ask for X-ray :
 • If radiopaque → stone
 • If no → Stricture.

TTT:

- **Proximal** (near gland) → superficial Parotidectomy
- **Distal** (near duct) → meotomy
- **Intermediate** → expectant ttt : by dilating duct every month by dilator till we found the stone or relieve stricture.

Non-Obstructive :

- Acute inflammation

• Viral :

- 1- Mumps
- Usually bilateral (may start unilateral)
- Occurs in children .
- we scared of 3 complications
2ry encephalitis, Pancreatitis. Orchitis
- Require isolation , bed rest ,antibiotics & vitamins

2- Coxsackie virus

• Bacterial (usually unilateral)

TTT:

- 1- TTT of predisposing factors
- 2- Analgesia & Massive antibiotics
- 3- Hilton incision (pre-auricular longitudinal incision & open the fascia transversely to avoid injury of facial nerve) & evacuation
- Don't wait for fluctuation because of dense parotid fascia.

Chronic (all bilateral)

1- Endemic parotitis

- Bilateral.
- ttt: reassurance & conservative.

2- Sialosis (Sialadenosis): conservative.

- Better seen (inspection) than felt
- Associated with :
 • Acromegaly
 • Diabetes (controlled or not)

3- Lipomatous pseudohypertrophy

- Exaggerated form of sialosis
- Sagging of the enlargement.

4- Sialectasis (ectatic duct) conservative.

→ by x-ray: Sand ground appearance.

5- Sarcoidosis :

- Generalized lymphadenopathy with hilar shadow except submental LNs
- Renal calcinosis & Renal stones

6- Sjogren's syndrome

- Lympho-epithelial disease complex
- Rheumatoid arthritis
- Dry eye
- Dry mouth (due to chronic diffuse parotitis)

Predisposing factors of acute bacterial parotitis :

- 1- Immunosuppressive
- 2- Local irradiation
- 3- Chemotherapy
- 4- Diabetic, neglected, poor control
- 5- Bad oral hygiene.

It's difficult for parotid to get inflamed because it's highly vascular, so there should be predisposing factors ,, TTT of these predisposing factors

Anterior Triangle swellings

1- Submandibular Triangle	Cystic	Ranula	<p><i>Suggested by:</i> translucent cyst lateral to midline, with domed, bluish discoloration in floor of mouth lateral to frenulum → presents itself as swelling in submandibular or submental triangle.</p> <p><i>management:</i> 1- Marsupialization (deroofting) & suture cyst wall to oral mucous m. 2- Excision(difficult) in recurrent cases.</p>
	Solid	Multiple LNs	<p>To differentiate, roll the swelling:</p> <p>- If rolled → LNs</p> <p>- If not rolled → - Fixed LNs (long stand neglected inflamed LNs) - Salivary (due to floor muscle: mylohyoid muscle)</p> <p style="text-align: center;">↓</p> <p>Bimanual examination :</p> <p>- if 2 lobes are palpable → Salivary gland</p> <p>- if Not→ Fixed LNs <u>or</u> superficial parotid tumor</p> <p style="text-align: center;">↓</p> <p>Excision & pathology to differentiate.</p>
2- Submental Triangle	Cystic	Ranula	
	Solid Single or Multiple	LNs only (no salivary gland)	
3- Carotid Triangle	Cystic	Branchial cyst "Congenital"	<p><i>Suggested by:</i> fluctuant swelling at anterior border of sternomastoid muscle, <i>Confirmed by:</i> US scan, CT scan</p> <p><i>Initial management:</i> excision it till lateral wall of pharynx with segment of lat. Wall</p>
		Tuberculosis ('cold') abscess	<p><i>Suggested by:</i> fluctuant(cystic) swelling with low grade or no fever. <i>Confirmed by:</i> acid-fast bacilli (AFB), on microscopy or culture and sensitivity of aspirate.</p>
		Carotid aneurysm	<p>Pulsatile swelling coincides with carotid pulsation. – Carotid angiography</p>
		Pharyngeal diverticulum	<p><i>Suggested by:</i> intermittent, fluctuant, compressible, swelling (usually on left) under sternomastoid muscle , and dysphagia. <i>confirmed by:</i> barium swallow fills pouch. <i>Initial management:</i> surgical referral for excision.</p>
	Laryngocele	<p><i>Suggested by:</i> fluctuant swelling in neck which becomes prominent on straining. Swelling is resonant & compressible Occurs in musicians . <i>Confirmed by:</i> MRI & laryngoscope.</p>	
Solid	Multiple LNs	Carotid body tumor (potato Tumor)	<p><i>Suggested by:</i> Very slowly growing mass with history of mass for years</p> <p>- mobile : from side to side but not vertical</p> <p>- arising from chemoreceptors at carotid bifurcation (upper third of sternomastoid),</p> <p>- pulsatile , bruit maybe heard by stethoscope.</p> <p>- My extend to parapharyngeal space present in oropharynx.</p> <p><i>Confirmed by:</i> 1- Angiography: widening of carotid artery bifurcation (characteristic sign)</p> <p><i>management:</i> Surgical excision with preservation of ICA.</p>
	Single		

Posterior Triangle swellings

Cystic		Cystic hygroma (Cavernous Lymphangioma) "Congenital"	<p><i>Suggested by:</i> Single , large , irregular, ill-defined, swelling that transilluminates well (only translucent neck swelling) , appears at birth or <20y of age.</p> <p><i>Confirmed by:</i> US scan, CT scan, <i>Complications:</i> Recurrent infection , respiratory distress due to compression of trachea, increases in size on coughing or crying. <i>management:</i> Surgical excision at about the age of 3 years.</p>
		Tuberculosis ('cold') abscess (most common site)	<p><i>Suggested by:</i> fluctuant(cystic) swelling with low grade or no fever. <i>Confirmed by:</i> acid-fast bacilli (AFB), on microscopy or C & S of aspirate.</p>
		Esophageal diverticulum	<p><i>Confirmed by:</i> barium swallow fills pouch.</p>
		Pneumatocele	<p><i>Suggested by :</i></p> <p>- Cystic swelling in the supraclavicular region -Become prominent on straining</p> <p>- Resonant & compressible <i>management :</i> 1- Correct straining factors 2- plication of Sibson's fascia.</p>
Solid	Multiple LNs	LNs	<p>The presence of a cervical rib can cause a form of thoracic outlet syndrome due to compression of the lower trunk of the brachial plexus or subclavian artery → Compression of the brachial plexus may be identified by weakness of the muscles around the muscles in the hand, near the base of the thumb "neurovascular deficient". (diagnosed by x-ray or CT)</p> <p>- Mostly seen in newborns due to birth trauma - Rare condition in which ischemia of the muscle → fibrosis & mass</p>
	Single	Cervical rib	
		Sternomastoid tumor	