

Thyroid tumors & nodules info

Epidem	<ul style="list-style-type: none"> -Rare -females -if present, is benign (e.g.: follicular adenoma)
Gross	<ul style="list-style-type: none"> -If found, solitary small nodule -if a nodule is found, its highly Likely to be a tumor
Diagnosis Method	<p>using Radioactive Uptake Study RUS</p> <p>great uptake = active nodule (graves)</p> <p>minor uptake = inhibited nodule (tumors - require fine needle aspiration for diagnosis of which type of tumors)</p>
Diagnoses (clues)	<p>Tumors</p> <ul style="list-style-type: none"> -if a nodule found in a young male -if it was solitary <p>Benign tumor</p> <p>Hot tumor (using RUS - high uptake)</p> <p>Malignant tumor</p> <p>Patient had a history of head and/or neck radiation</p>
Toxic tumors	<ul style="list-style-type: none"> -A tumor so active that it causes local toxicity -high RUS uptake (shows very dark) -almost all thyroid tumors are NOT toxic (non-fun, & cold)

/thyroid tumors (lymphoma + adenomas + carcinomas)

Thyroid Lymphoma	
Types	Primary “within the thyroid” Secondary “metastes”
Risk	Hashimoto

Adenoma	
Is	A benign thyroid tumor
Gross	One mass
Origin	Follicular cells
Hallmark	Presence the fibrous capsule around the tumor
Symptoms	Nill - painless
Diagnosis	Biopsy & L/M
types	<p>Macro -simple colloid tumor</p> <p>Micro -aka: fetal -seen in emryo thyroid</p> <p>Hurthle -large eosinophilic -very rich in mitochondria</p>
Invasion & metastasis	Nill
Treatment	Excision
Prognosis	Excellent

Carcinoma

Is	Thyroid malignant tumor
Epidem	Very, very rare
Etiology	Radiation
Types	(From most common to rarest) 1-Papillary 2-follicular 3-medullary 4-anaplastic “undifferentiated”
(1) Papillary	
Gene	RET, KTRK, BRAF
Epedim	Markedly inc in the past 30 years
RUS	Cold
Risk	Hashimoto & radiation
Hallmark	N features
Gross	Papillaries (finger-like projections)
L/M BOSS	-grooved N “coffee <u>B</u> ean-like” - <u>O</u> rphan annie N “very clear” - <u>P</u> sammoma bodies “calcification” - <u>P</u> seudo-inuclusions “cytoplasm invasion mimic IC inclusions”
Symptoms	-mostly asymptomatic thyroid nodule -may spread to cervical lymphs -doesn’t cause death “no mortality”
Prognosis depends on	-presence of extrathyroidal extensions -metastases -age
Prognosis in general	Excellent
(2) follicular	
Gene	RAS, PAX, PPAR-gamma
Hallmark	Capsule & vascular invasion (Can metastase hematogenously)
Risk	Iodine def.
Invasion	Minute -only invade the capsule and/or one vessel, but still remains encapsulated Wide -systemic invasion

(3) medullary

Gene	RET, MEN2(multiple endocrine neoplasms)
Hallmark	Neuroendocrine - calcitonin
Origin	Parafollicular cells
Hormones	-Significant inc in calcitonin secretion, causing hypocalcemia -calcitonin deposit within the tumor causing local amyloidosis
Epidim	-mostly sporadic -only familial are ass with the genes above -if familial and ass with MEN, then its called FMTC (familial medullary thyroid carcinoma) -if present, it affects all paraT glands
L/M	-neurogranular cytoplasm -multicentricity (having multiple centers) -amyloidosis -necrosis & hemorrhage
Stains	-ImmunoHistoChemistry IHC (for calcitonin) -congo red (for amyloid)

(4) anaplastic

Gene	P53 (tumor suppressor gene)
Epidim	commonest of all anaplastic tumors, but rarest of thyroid
Origin	-Follicular cells -can arise from another tumorous cells (papillary)
L/M types	-giant pleomorphic cells (osteoclast-like multiN giant cells) -spindle cells -giant & spindle cells -small cells
Prognosis	Worst! -highly malignant, rapidly growing, local&systemic metastasis (usually to trachea & eso "dysphagia")