

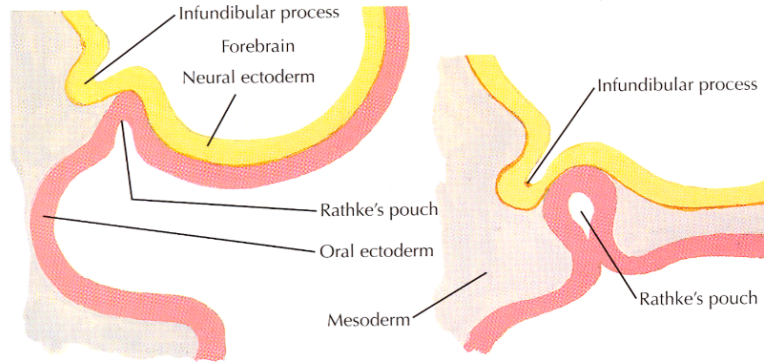
Development and teratology of the endocrine system and pharyngeal apparatus of the embryo

Anna Mac Gillavry Danylevska

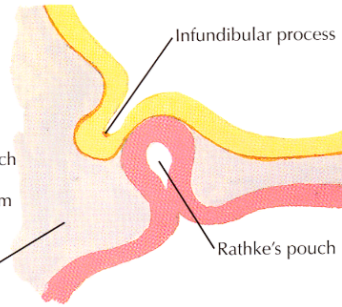
15.5.2023

Pituitary gland

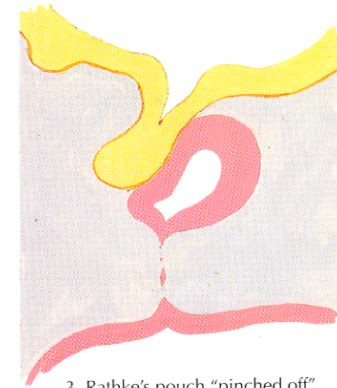
- Ectoderm (Rathke's pouch)
- Neuroectoderm of ventral wall of diencephalon



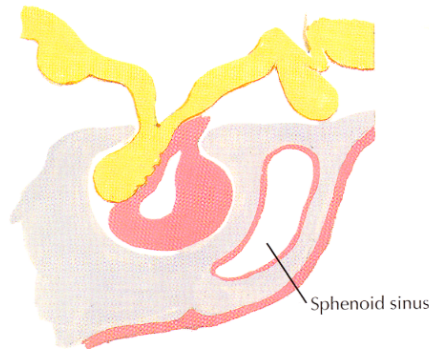
1. Beginning formation of Rathke's pouch and infundibular process



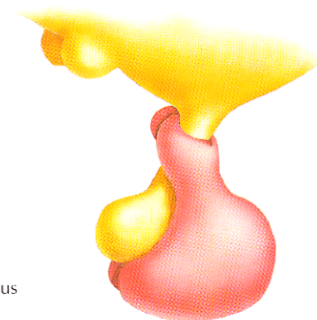
2. Neck of Rathke's pouch constricted by growth of mesoderm



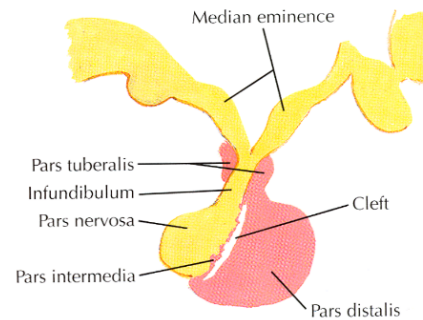
3. Rathke's pouch "pinched off"



4. "Pinched off" segment conforms to neural process, forming pars distalis, pars intermedia and pars tuberalis

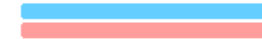


5. Pars tuberalis encircles infundibular stalk (lateral surface view)



6. Mature form

Development of the Hypophysis



F. Netter M.D.

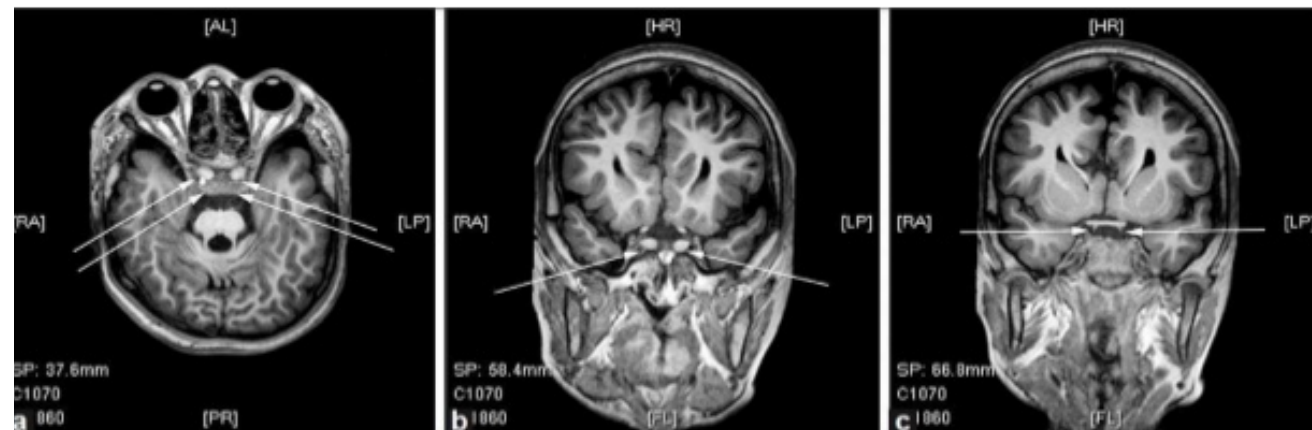
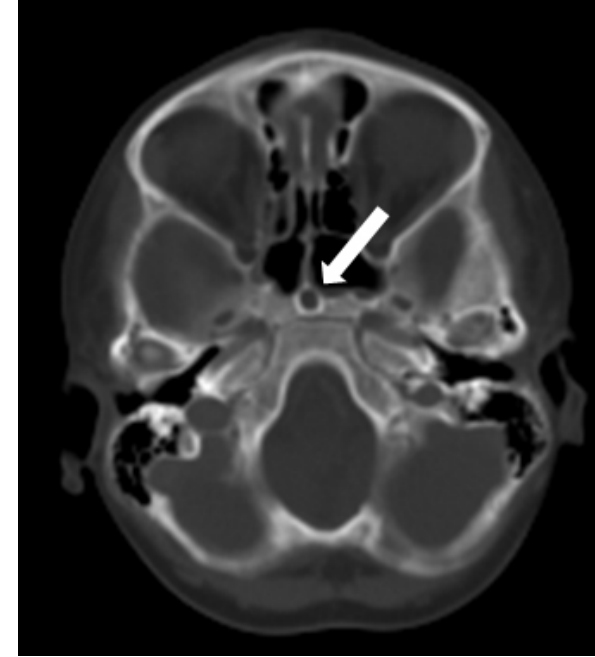
Craniopharyngeal canal

Pharyngeal hypophysis

Agensis/hypoplasia - agensis is incompatible with life; panhypopituitarism

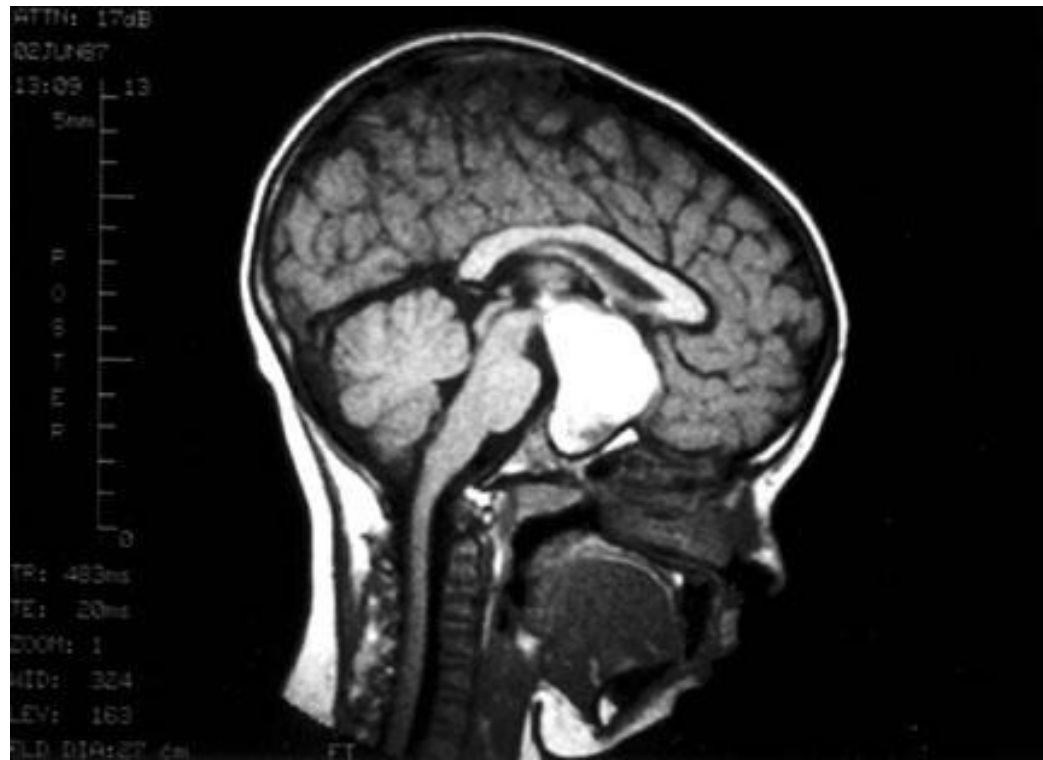
Duplication of the gland – very rare

Ectopic posterior pituitary – pituitary dwarfism



[Duplication of the pituitary gland associated with multiple blastogenesis defects: Duplication of the pituitary gland \(DPG\)-plus syndrome. Case report and review of literature - Surgical Neurology International](#)

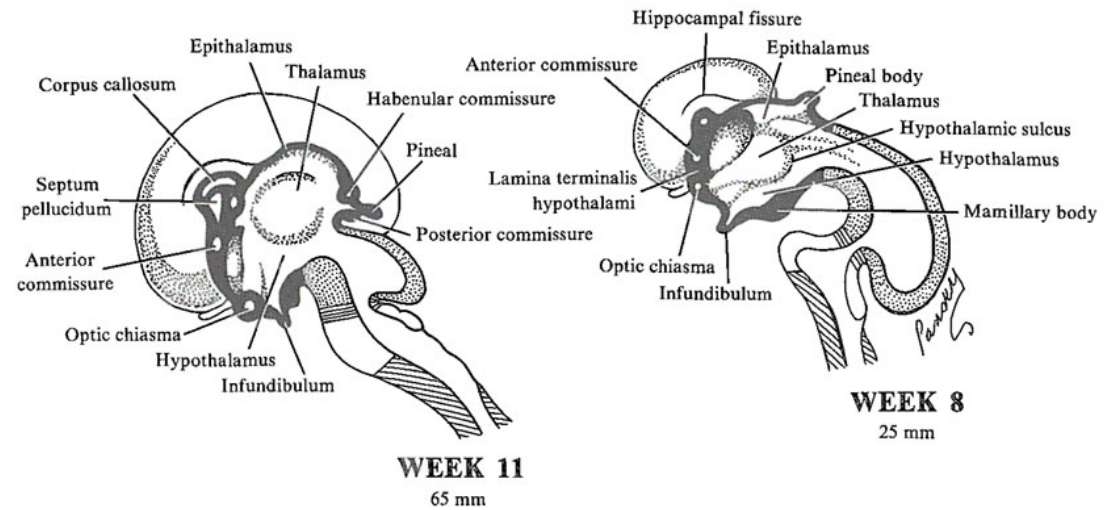
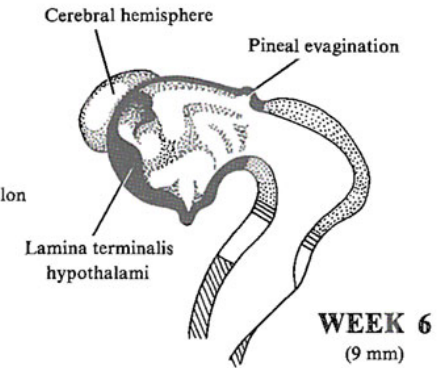
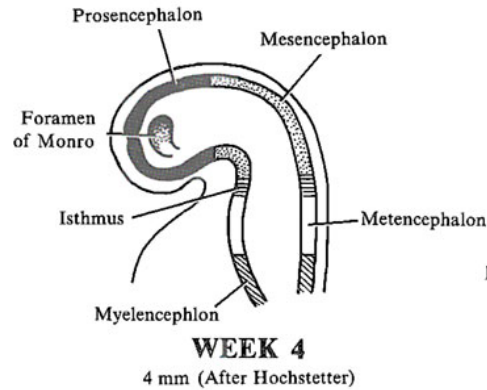
Craniopharyngiomas – usually lie above the sella; cause hydrocephalus, growth failure, diabetes insipidus, lose of peripheral vision



[Pediatric Craniopharyngioma: Background, Pathophysiology, Epidemiology \(medscape.com\)](https://www.medscape.com/lookup/topic/pediatric-craniopharyngioma)

Epiphysis

- thickening of caudal part of ependyma that does not contribute to development of choroid plexus at the roof of diencephalon
- neuroectoderm



Pineal gland agenesis – mutations PAX6 (paired box gene 6)

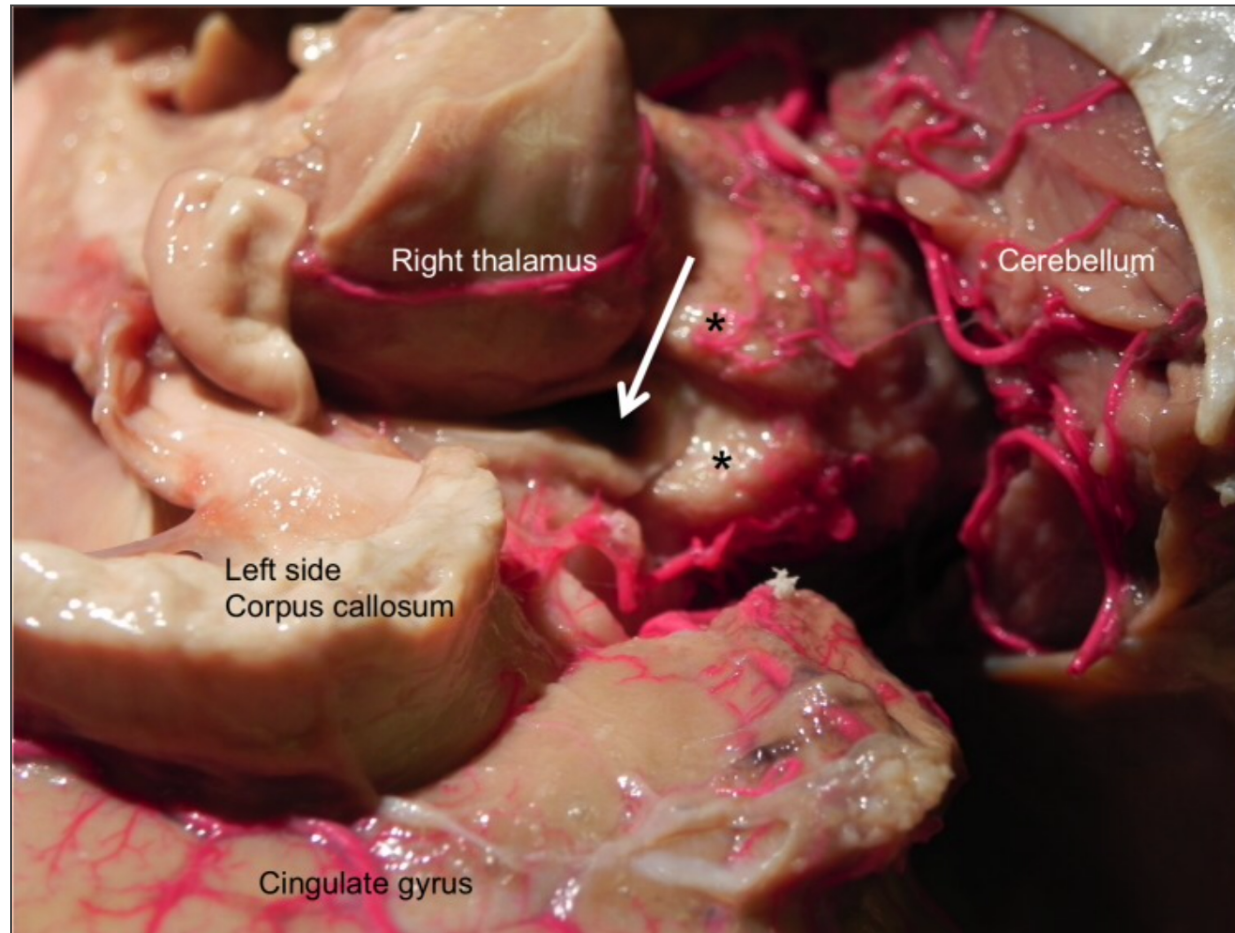
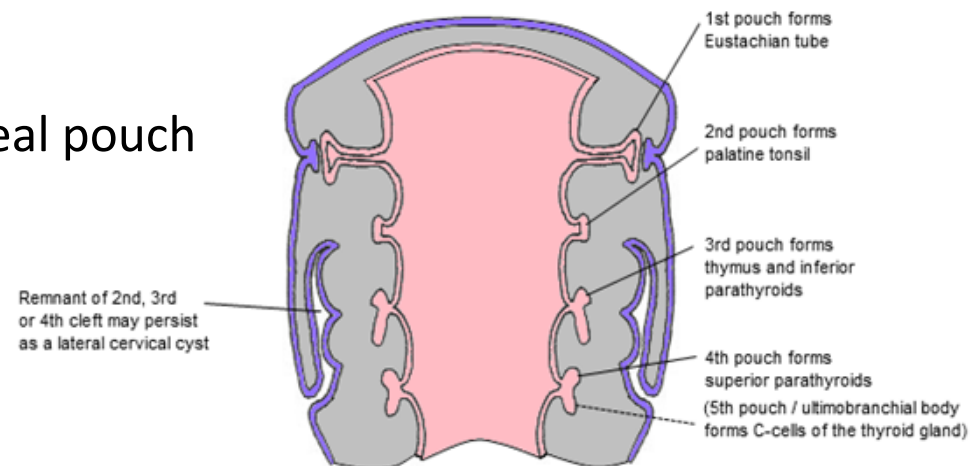


Figure 1

Absent Pineal Gland

Thyroid gland

- endodermal proliferation of pharyngeal floor between tuberculum impar and copula
- obliterating ductus thyreoglossus
- foramen caecum
- bilobed diverticulum
- lobus pyramidalis
- C-cells
 - neural crest origin
 - ultimobranchial body of 5th pharyngeal pouch



Pyramidal lobe – in 50 % of population

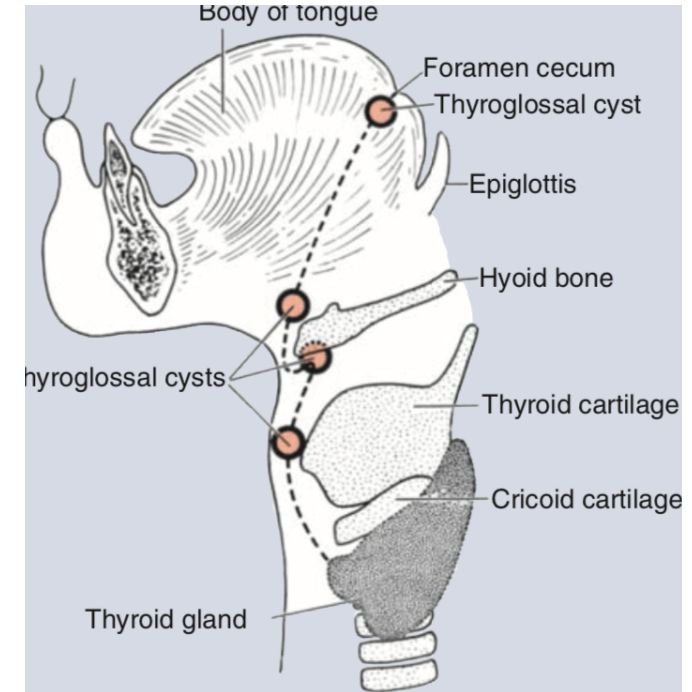
Congenital hypothyroidism (1/3000)

- ectopic thyroid
- hypoplasia, agenesis
- TSH deficiency

Ectopic thyroid gland – in 90 % cases it is lingual thyroid gland; sublingual thyroid gland

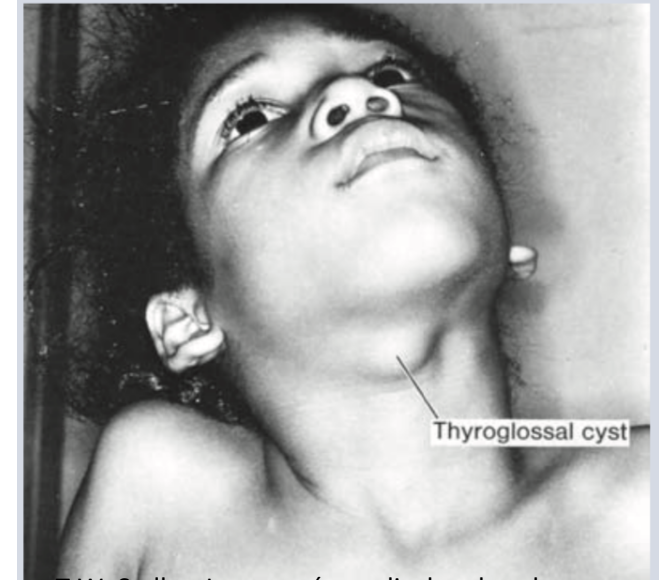
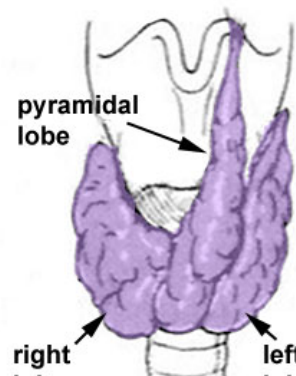
Thyroglossal duct cyst – clinically important to distinguish from ectopic thyroid gland!

Thyroglossal fistula



ysts. These cysts, most frequently found in the hyoid region, a

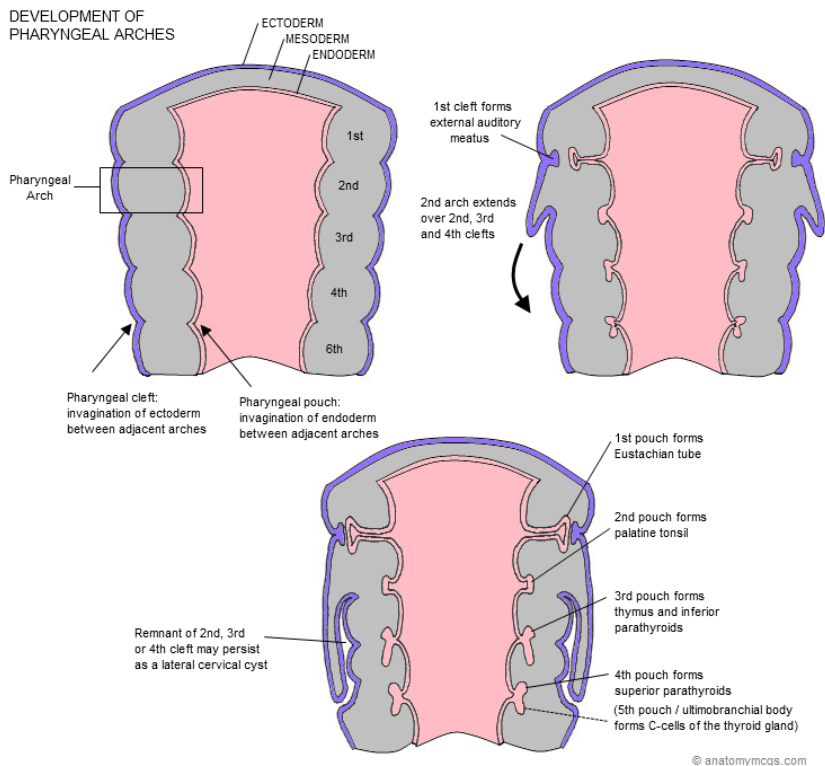
Thyroid Pyramidal Lobe (neck ventral view)



T.W. Sadler, Langman's medical embryology, 12th edition

Embryonic development of parathyroid gland

- glandulae parathyroideae superiores from endoderm of 4th pharyngeal pouch
- glandulae parathyroideae inferiores from dorsal process of 3rd pharyngeal pouch
- together with thymus descend to lower poles of thyroid



Ectopic parathyroid tissue –
the inferior parathyroids are more variable in their position
Supranumerary parathyroid glands

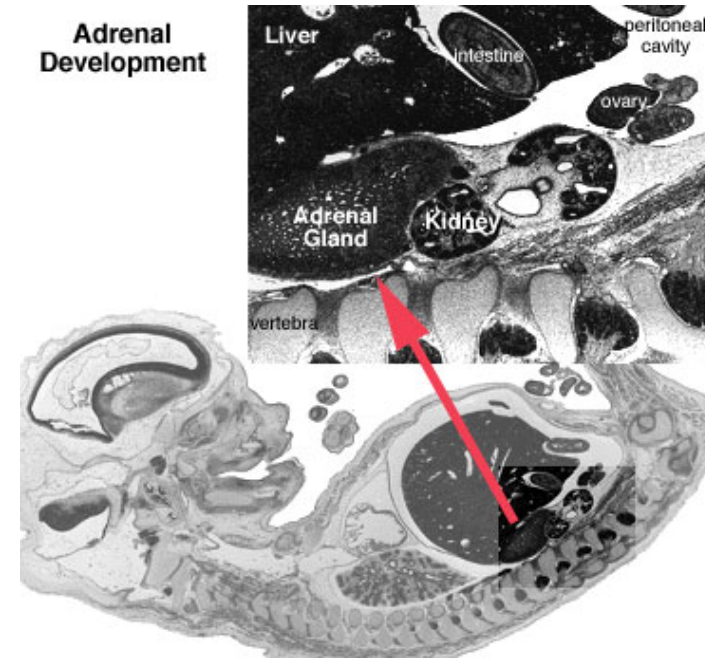
Suprarenal gland

Cortex

- Mesoderm ---> coelomic epithelium
- primitive fetal cortex 5-6th week
- definitive cortex
- zona reticularis fully differentiates within 3 years

Medulla

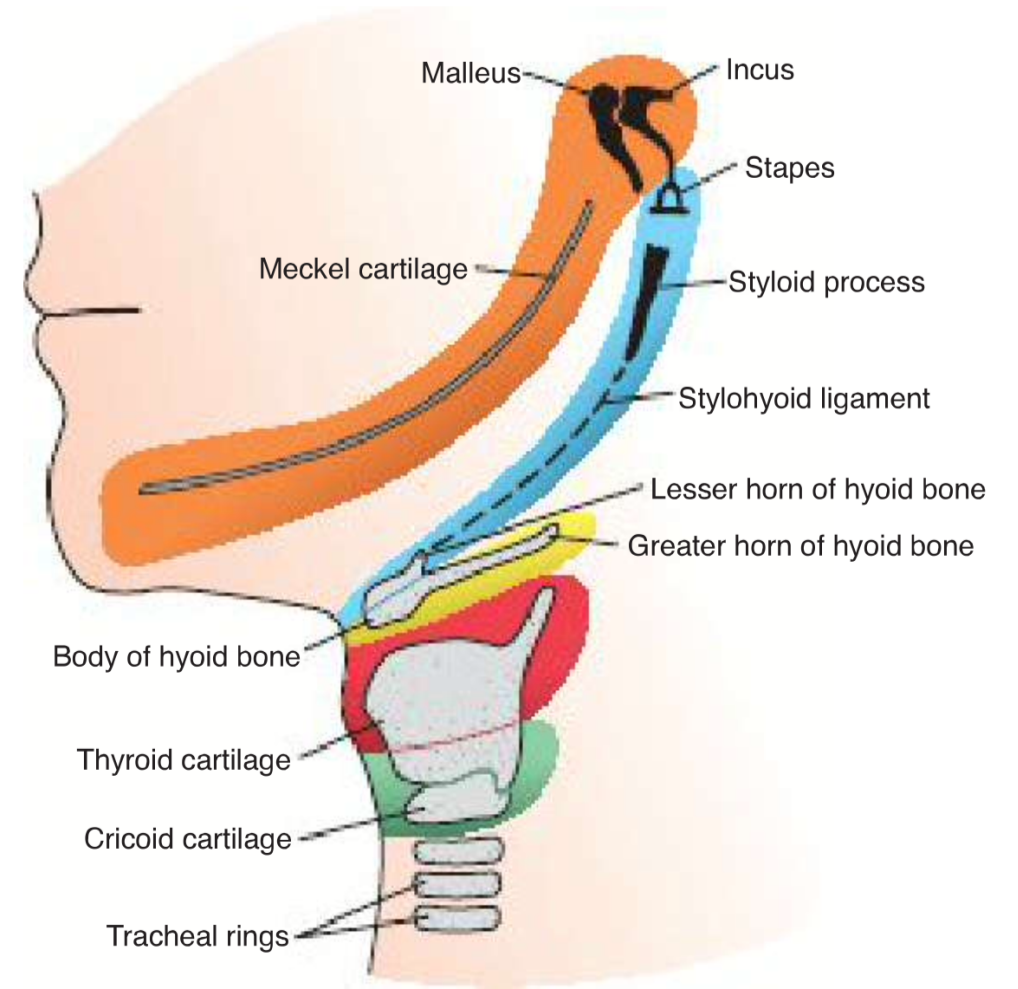
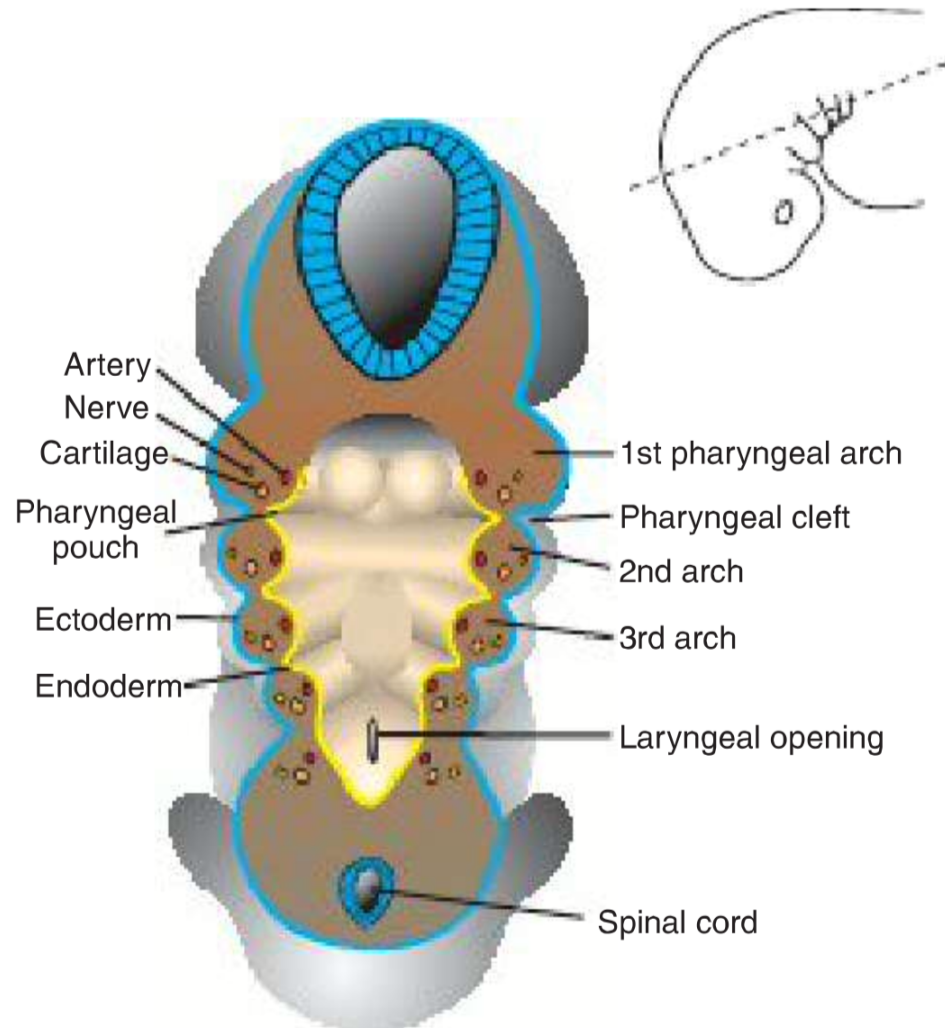
- neural crest



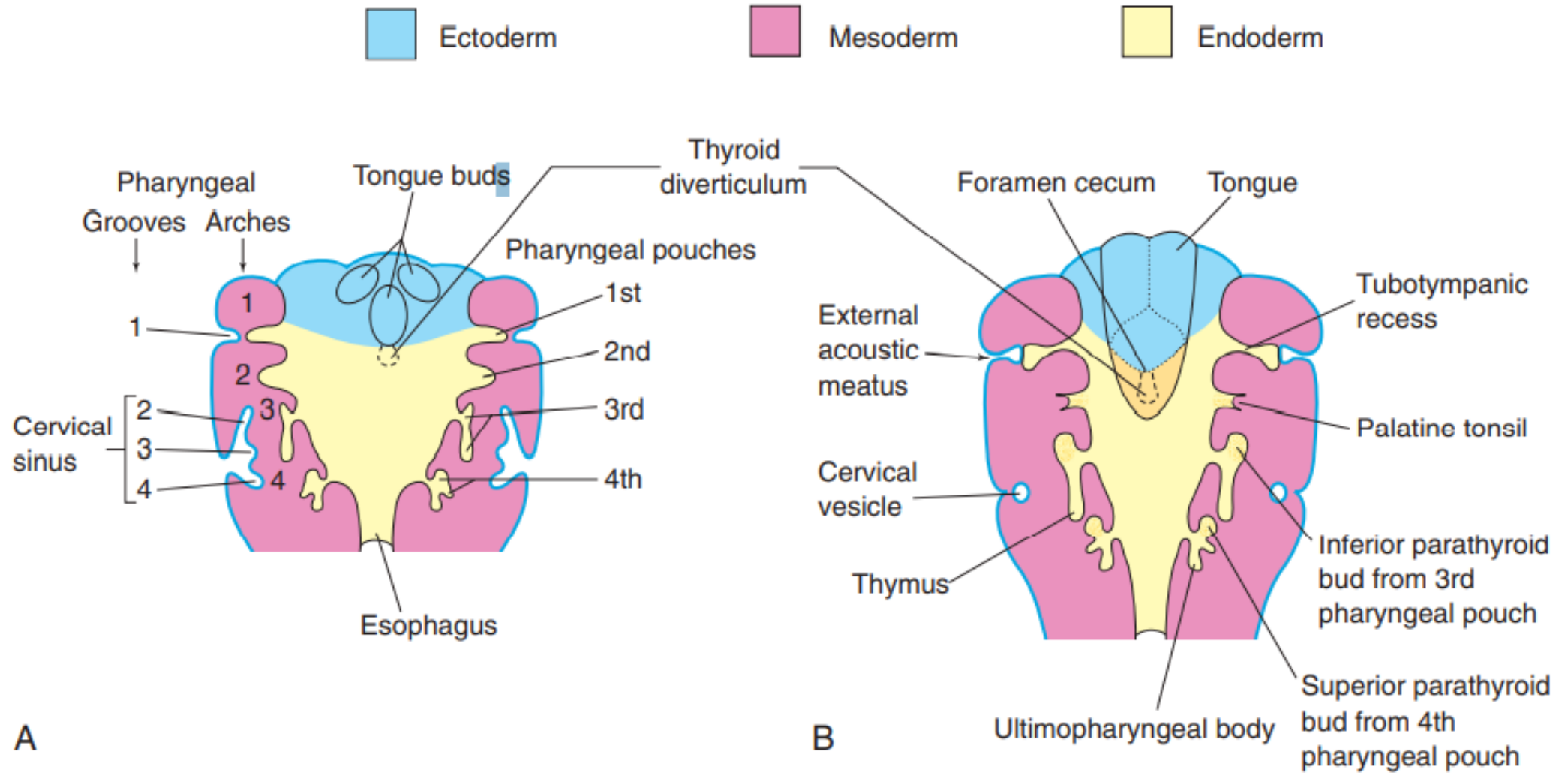
[Week10 adrenal - Endocrine - Adrenal Development - Embryology \(unsw.edu.au\)](http://www.unsw.edu.au)

Congenital adrenal hyperplasia – group of autosomal recessive disorders – excessive production of androgens: causes rapid growth and accelerated skeletal maturation in both sexes

Neck region



Neck region – pharyngeal apparatus

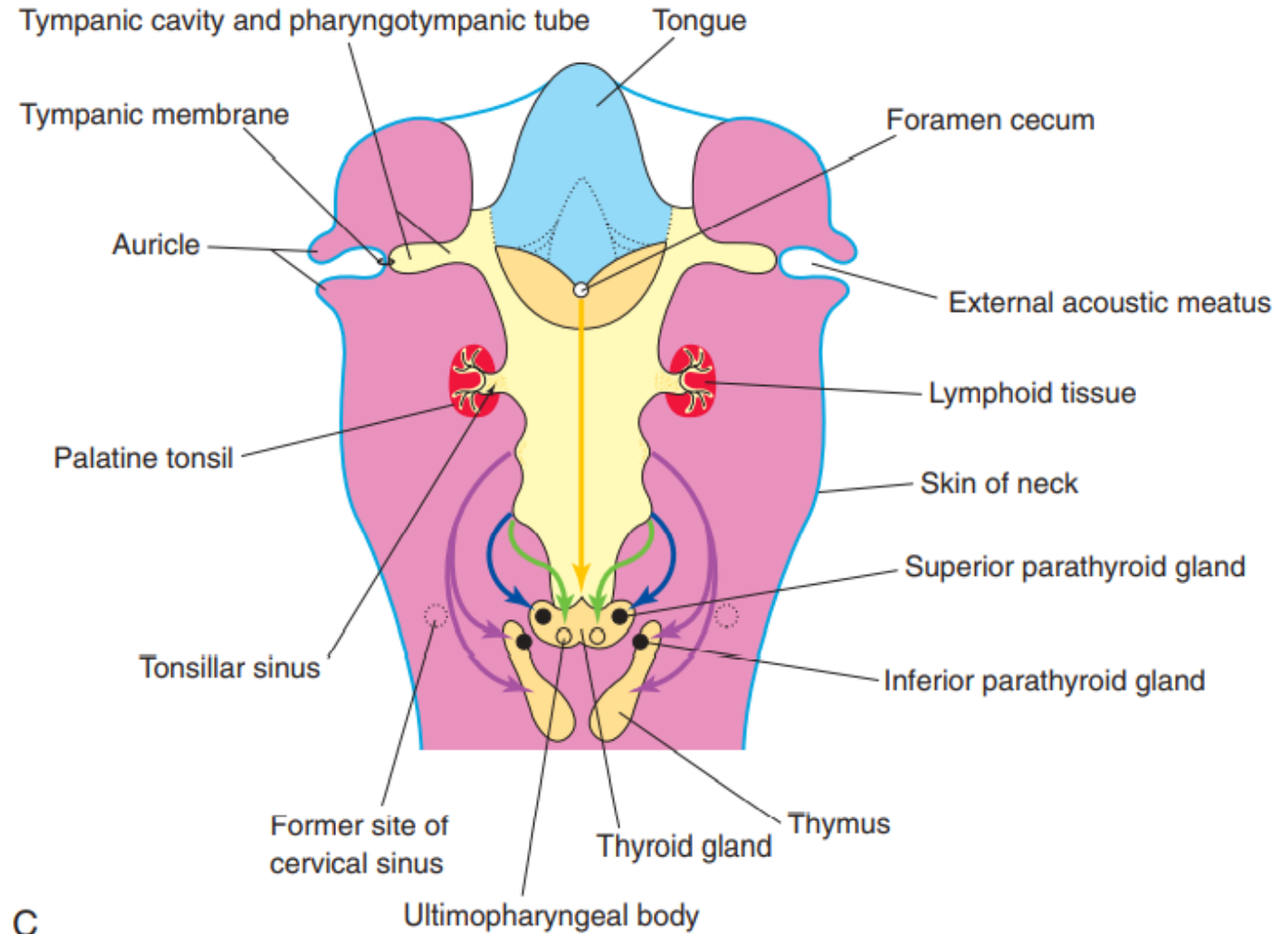


Ectopic thymic tissue

Ectopic parathyroid tissue

Branchial fistulas: external and internal

Cervical cysts



Craniofacial defects associated with neural crest cells

- Mandibulofacial dysostosis – Treacher Collins syndrome: autosomal dominant, 1/50000
- Robin sequence: first-arch structures, 1/8500 – micrognathia, cleft palate and glossoptosis
- 22q11.2 deletion syndromes: DiGeorge syndrome, DiGeorge anomaly, velocardiofacial syndrome etc.
 - 1/4000
- Hemifacial microsomia (oculoauriculovertebral spectrum – Goldenhar syndrome) – 1/5600, assymetry in 65% cases; involves maxillary, temporal, zygomatic bones, ears, eyes, vertebrae. Cardiac deffects in 50% cases.



<https://economictimes.indiatimes.com/magazines/panache/man-born-with-treacher-collins-syndrome-was-rejected-by-biological-parents-36-hrs-after-birth-for-his-face-but-his-adoptive-mom-didnt-care/articleshow/93541374.cms?from=mdr>