



CUSHING SYNDROME

KETSETZI LAURA EVGENIA 2ND YEAR ED; ENGLISH DIVISION SCIENCE CLUB

SUPERVISOR: DR. HAB. N. MED. DAVID AEBISHER, PROF. UR 2

STRUCTURE



General
Information



Cause



Physiology



Symptoms
& Signs



Diagnosis



Treatment

GENERAL INFORMATION

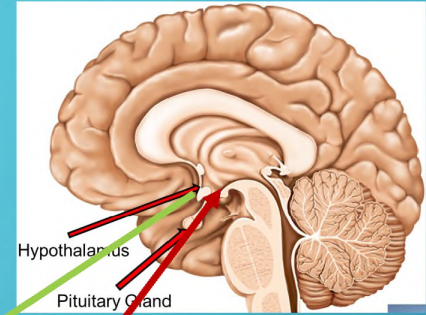
= Disorder, production of excessive cortisol (=stress hormone) over long period

- **Two types of Cushing syndrome:** Exo- & Endogenous
- Affects adults, aged 30 – 50
- Affects 3x more women than men
- 10-15 per million people affected p.a.

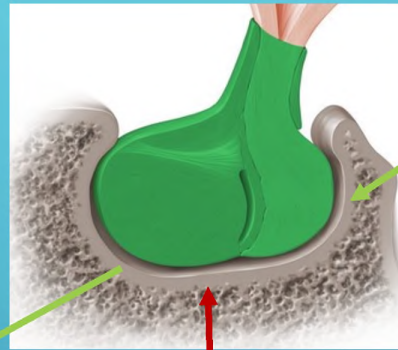
CAUSE

1. Iatrogenic (long-term treatment with steroids)
2. Over-secretion by adrenal glands (adrenal adenoma/carcinoma)
3. Over-stimulation of the adrenal glands (pituitary tumor)
4. Over-stimulation of the adrenal glands by an ectopic ACTH producing tumor

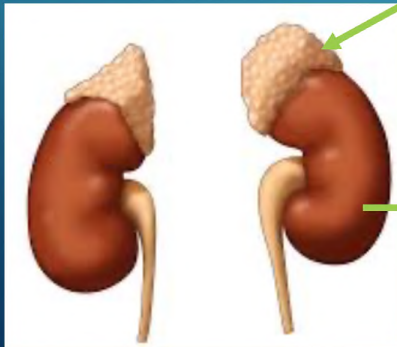
PHYSIOLOGY



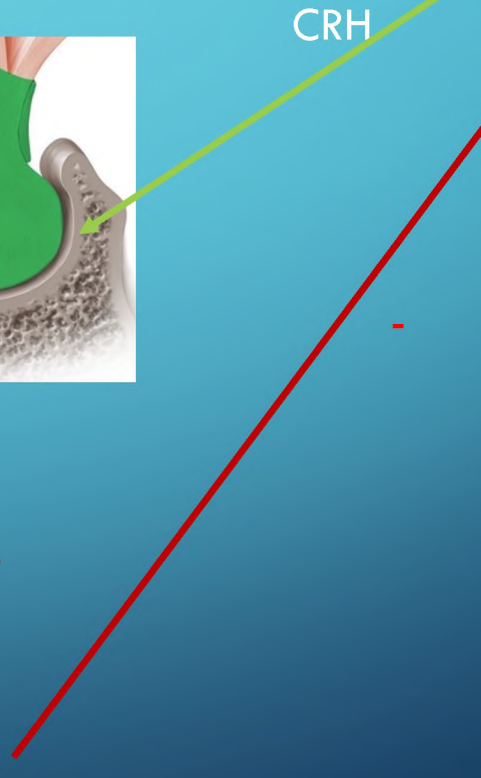
CRH



ACTH



Glucocorticoids: Cortisol



TWO TYPES



EXOGENOUS = OUTSIDE OF BODY

- **Most common cause:** Taking cortisol-like medications
- **Cortisol-like medications:** For treating inflammatory diseases; Suppressing immune system after organ transplantation
- Goes away after discontinuing the medication → **Temporary Disease**



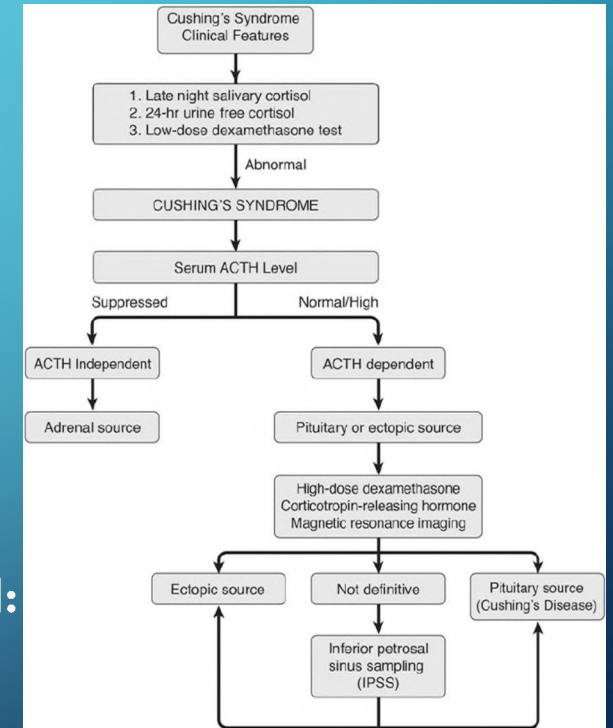
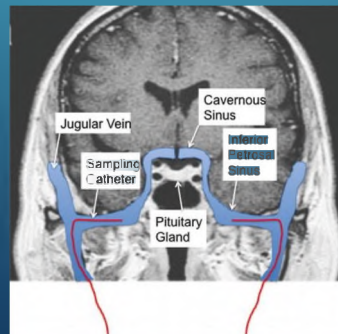
ENDOGENOUS = INSIDE THE BODY

- **Uncommon cause:** Adrenal glands overproduction of cortisol
- **Caused by hormone-secreted tumors:** In adrenal glands or pituitary tumors. Tumors produce excessive ACTH
- Pituitary adenomas responsible for >70% of cases
- Does not go away if not treated



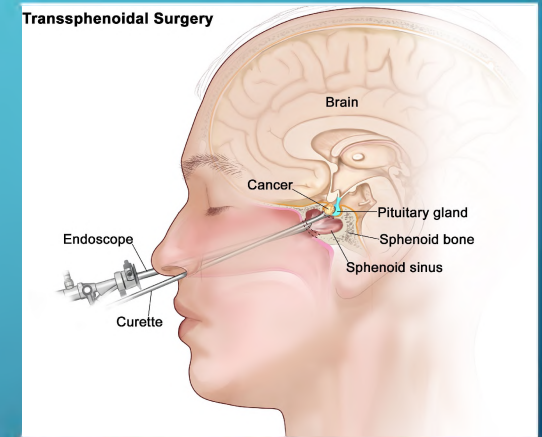
DIAGNOSIS

- Firstly, tests for **determination of excessive cortisol**:
 - 24-hours urinary cortisol test; Levels $>50-100\mu\text{g}$
 - Low dose of dexamethasone suppression test; High cortisol
 - Late night salivary cortisol test; High cortisol \rightarrow Positive
- Afterwards, tests for **determination of the source of excessive cortisol**:
 - Magnetic resonance imaging (MRI); tumor 5mm
 - Petrosal sinus sampling



TREATMENT

- Treatment depends on cause
- **Options:** surgery, radiation, chemotherapy, or cortisol-reducing medicines
- **Treatment for pituitary tumors:** Surgery; Success rate: 90%
- If surgery fails → surgery can be repeated
- After removal of tumor, pituitary doesn't produce ACTH for a while → Intake of cortisol medicine
- If surgery fails/ isn't possible: Radiation therapy
- **Treatment of adrenal tumors:** Surgery
- If both adrenal glands removed → intake of medicine to replace cortisol



SOURCES

- Book: Clinical pathophysiology made ridiculously simple by Aaron Berkowitz
- Chaudhry HS, Singh G. Cushing Syndrome. [Updated 2021 Jul 30]. In: StatPearls [Internet]. Treasure Island (FL): StatPearls Publishing; 2022 Jan-. Available from: <https://www.ncbi.nlm.nih.gov/books/NBK470218/>
- Miyachi Y. Pathophysiology and diagnosis of Cushing's syndrome. Biomed Pharmacother. 2000 Jun;54 Suppl 1:113s-117s. doi: 10.1016/s0753-3322(00)80026-8. PMID: 10915006.
- Lonser RR, Nieman L, Oldfield EH. Cushing's disease: pathobiology, diagnosis, and management. J Neurosurg. 2017 Feb;126(2):404-417. doi: 10.3171/2016.1.JNS152119. Epub 2016 Apr 22. PMID: 27104844.
- <https://www.niddk.nih.gov/health-information/endocrine-diseases/cushings-syndrome>
- <https://www.endocrine.org/patient-engagement/endocrine-library/cushings-syndrome-and-cushing-disease>