

THE 7TH ASIAN NEURO-OPTHALMOLOGY SOCIETY CONGRESS 27-29 SEPTEMBER 2013



CERTIFICATE
OF

ATTENDANCE

THIS IS TO CERTIFY THAT

M. HIDAYAT, MD

HAS ATTENDED THE 7TH ASIAN NEURO-OPTHALMOLOGY SOCIETY CONGRESS AS

SPEAKER

Muhammad Sidik, MD

Chairman

Masato Wakakura

President of Asian Neuro-Ophthalmology Society



Young Man with Pituitary Adenoma

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Introduction

The pituitary gland, or hypophysis, is an endocrine gland which produce number of hormones which control the secretions of many other endocrine glands

Its anatomical position is important in ophthalmological view

Development

The anterior pituitary (adenohypophysis) arises from Rathke's pouch, an upward growth from the ectodermal roof of the stomodeum

The posterior pituitary (neurohypophysis) arises from a downward growth from the floor of the diencephalon

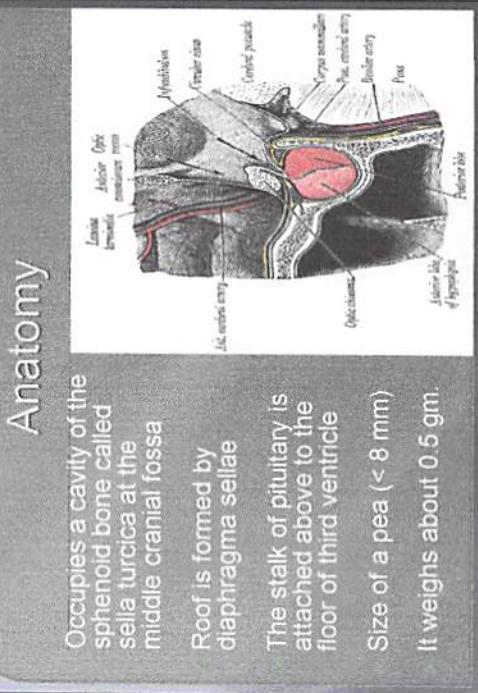
Anatomy

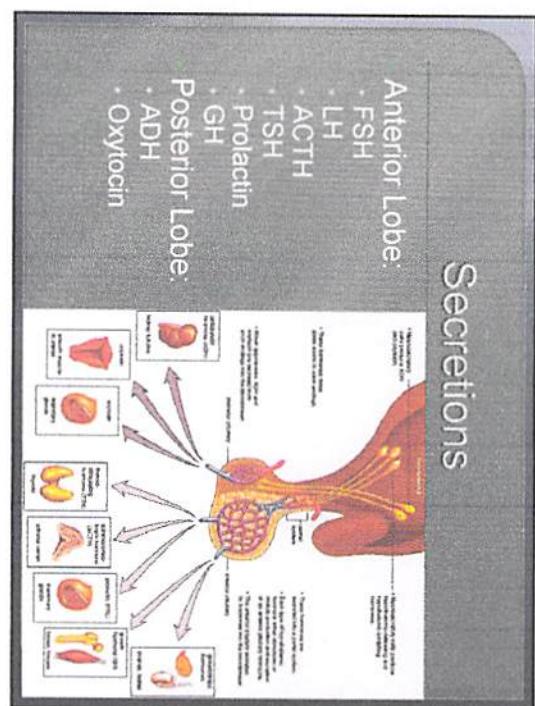
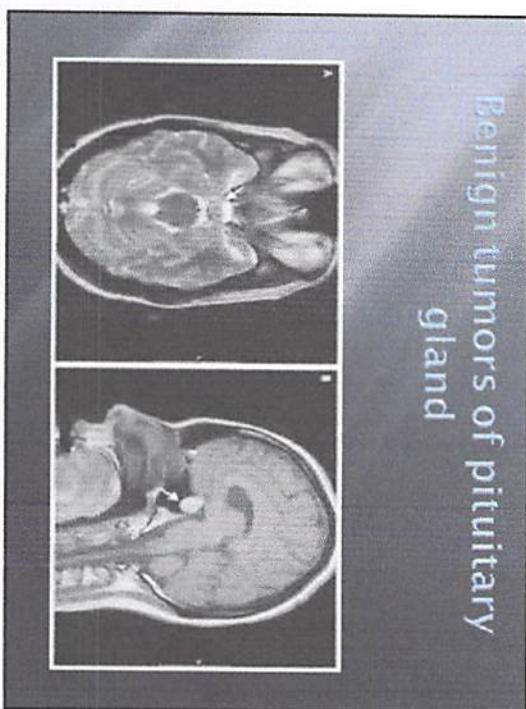
Occupies a cavity of the sphenoid bone called sella turcica at the middle cranial fossa

Roof is formed by diaphragma sellae

The stalk of pituitary is attached above to the floor of third ventricle

Size of a pea (< 8 mm)
 It weighs about 0.5 gm.





Endocrine gland	Hormone	Function	Secretion control is made by
Pituitary			
Posterior			
ADH	Stimulates retention of water, decreasing its permeability at distal convoluted tubules and collecting ducts of the nephrons	Stimulates the excretion of fluid and blood glucose	Orbit (Hypothalamus)
Oxytocin	Stimulates uterine contractions during parturition and stimulates milk ejection (letdown) in the mammary glands	Suckling by the neonate initiates a reflex arc	Orbit (Hypothalamus)
Anterior			
GH	Stimulates liver to produce growth factors that stimulate bone and cartilage growth	Growth (Hypothalamus)	
Prolactin	Stimulates mammary gland growth and milk production		
ACTH	Stimulates adrenal cortex to produce hormones [Glucocorticoids and mineralocorticoids]		CNS (Hypothalamus)
LH	• Stimulates development of follicles • Stimulates the production of sperm in the seminiferous tubules of the testes	Ovulation is triggered by GnRH (Hypothalamus)	
FSH	Stimulates development of ovaries		
TSH	Stimulates thyrocytes production by the interstitial cells of the testes	Orbit (Hypothalamus)	
MSH	Promotes distribution of melanin granules		

Classification

	Hormones	Clinical Features
Secretory (70%)	Chromophobe S (~50%)	Females: infertility, amenorrhea, gynaecomastia Males: Hypogonadism, impotence, sterility, libido, gynaecomastia
	Macroadenoma	
	Acidophilic (20%)	Giantism Acromegaly (adult) Gigantism (child)
Nonsecreting (25%)	Basodiphilic (5%) ACTH, FSH & TSH	Cushing disease, FSH & TSH tumors

Clinical presentation

Localized mass effects

• Chiasmal syndromes

- Compression of other adjacent structures

~ Cavernous sinus (paresis of 3rd, 4th or 6th CN causing disorders of extraocular motility)

- ~ Hypopituitarism (direct pressure, vascular damage)

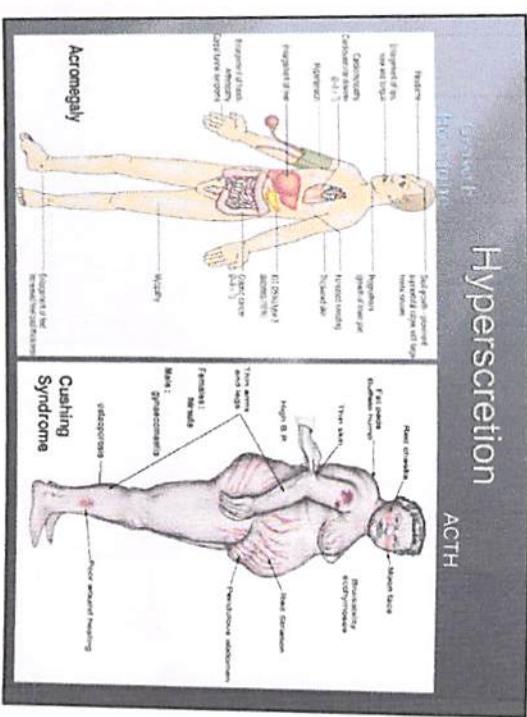
~ Papilloedema (raised ICP, very rare)

Endocrine effects

- Hypersecretion

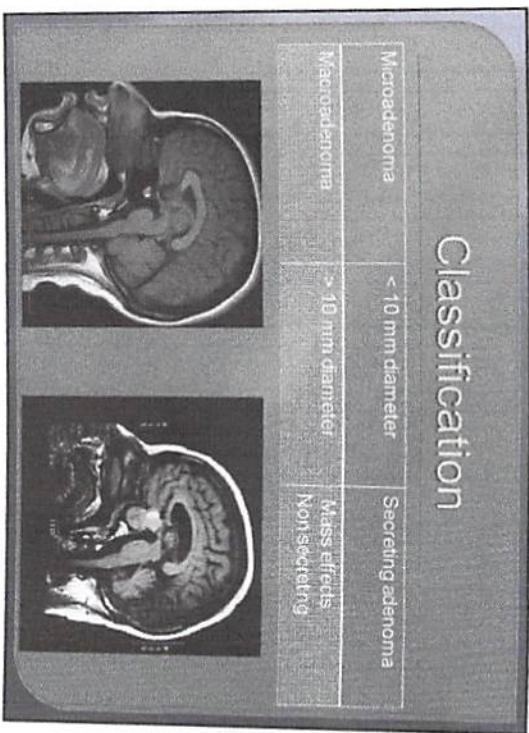
Hyperscretion

ACTH



Classification

Microadenoma	< 10 mm diameter	Secretory adenoma
Macroadenoma	> 10 mm diameter	Mass effects
		Nonsecreting



Hyperscretion Prolactin

Female

- Infertility-amenorrhea- galactorrhea

Male

- Hypogonadism, impotence, sterility, ↓ libido, gynecomastia, galactorrhea



Hyposecretion

Growth hormone deficit

- In children: dwarfism
- In adults: weakness, overweight, reduced cardiac output, low blood sugar levels, and reduced exercise tolerance

TSH deficit

Hypothyroidism

- Underactive adrenal gland which causes low blood pressure, hypoglycemia, fatigue, weight loss, vomiting, and low stress tolerance

ACTH deficit

Hypoadrenocorticism

ADH deficit

- Diabetes Insipidus

Visual field defect in pituitary adenoma

Chiasmal Syndromes

- Blurred vision
- Headache
- Diplopia
- Colour desaturation
- Visual field defect
- Optic atrophy
- Post fixation blindness
- Visual hallucination
- See-Saw nystagmus

Bitemporal hemianopia

Classic defect in pituitary adenoma

Occurs in central chiasmal defect

Superotemporal field affected first

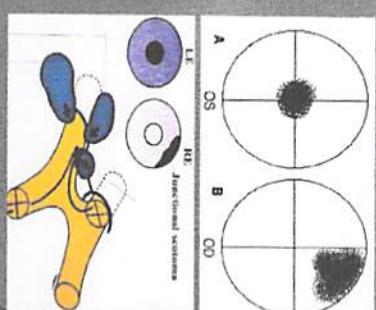
Lower temporal field defect



Junctional scotoma

Central scotoma in one eye with superotemporal visual field loss in the other eye

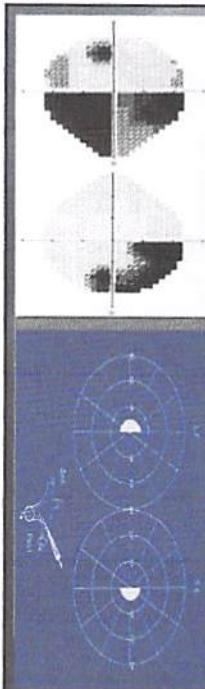
Caused by compression to anterior loop to the decussating nasal fibers in posterior optic nerve
(von Wilbrand's knee)



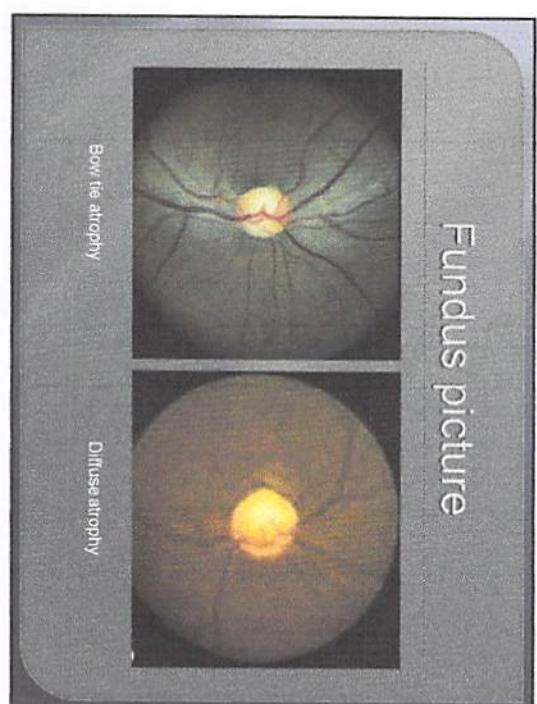
Incongruous homonymous hemianopia

Occurs in optic tract lesion

Compressing only the macular fibers

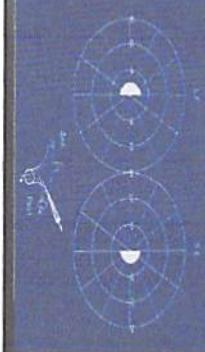


Fundus picture

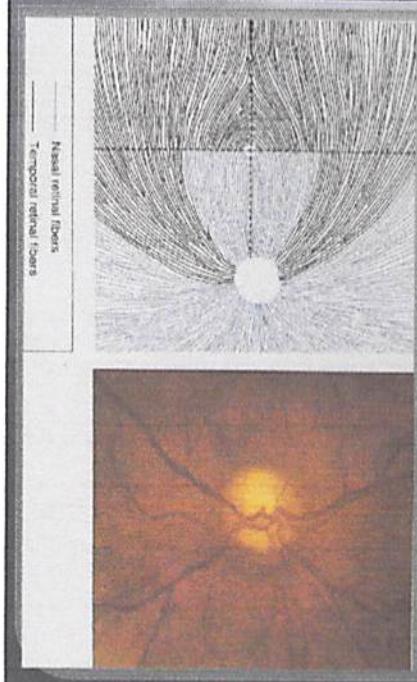


Bitemporal central scotoma

Occurs in post. Chiasmal lesion



Bow tie atrophy



Ocular Examination

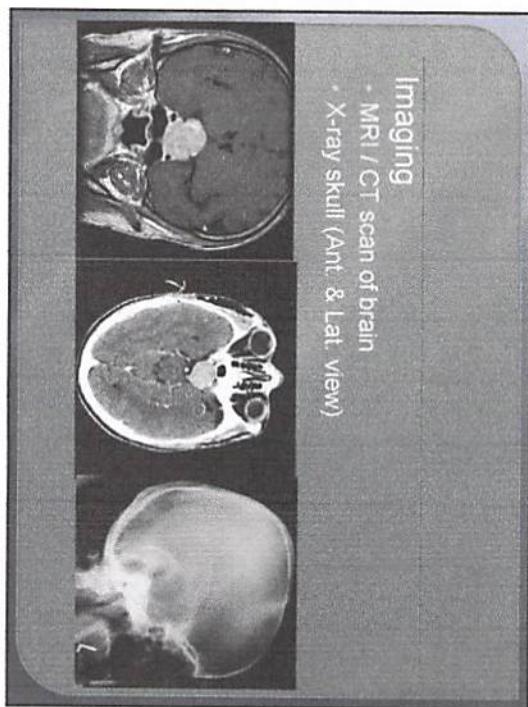
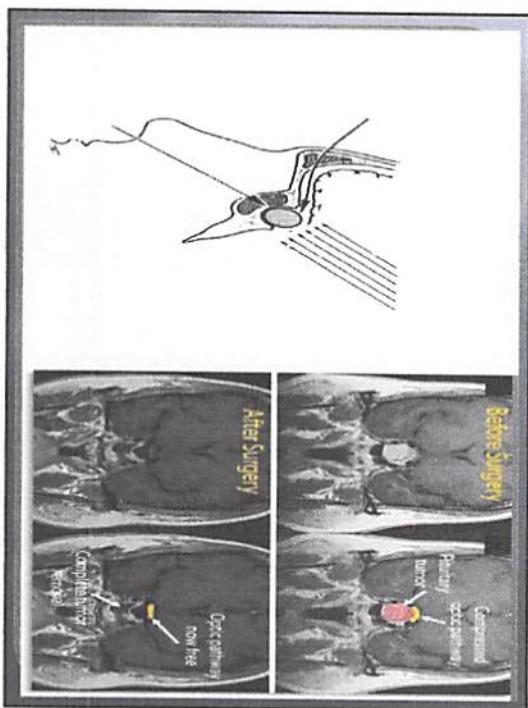
- Visual acuity
- RAPD
- Color vision
- Ocular motility
- Fundus examination

Pituitary apoplexy

- Large adenoma leading to haemorrhage or infarction of pituitary gland
- Occur in pregnancy
- Compresses hypophysial portal vessels
- Presentation: hyperacute chiasmal syndrome
- Treatment: high dose steroid / surgery

Investigations

- Endocrinological evaluation
 - Serum prolactin
 - FSH
 - TSH
 - GH
 - Insulin stress test
- Ocular investigation
 - Visual Field Analysis
 - Hass / Lees chart



Treatment

Referral to Endocrinologist & Neuro-surgeon

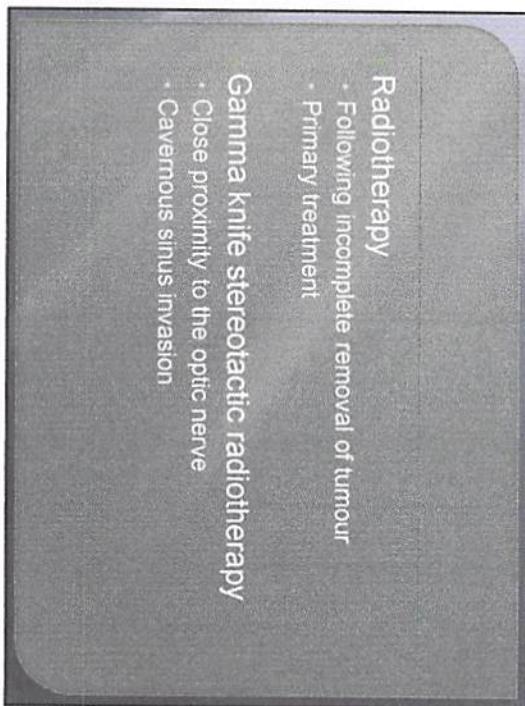
Observation

Medical therapy

- Dopamine agonists – Cabergoline/Bromocriptine

Surgery

- When mass causing severe compression
- Endoscopic - Transsphenoidal, transfrontal
- Craniotomy
- Visual recovery is tri-phasic



Therapeutic Modalities Summary

	Surgery	Radiotherapy	Medical
Non-functioning adenoma	Craniotomy Transsphenoidal	External beam Stereotactic radiosurgery	Medication Lifestyle changes
Pituitary adenoma	Transsphenoidal Radiosurgery	External beam Stereotactic radiosurgery	Medication Lifestyle changes
Acromegaly	Transsphenoidal Radiosurgery	External beam Stereotactic radiosurgery	Medication Lifestyle changes

Follow-up after treatment

Medical therapy –

- Monthly for large tumors or during pregnancy for tumors of any size
- 6 month intervals in microadenoma for 1 year, then yearly

Surgery –

- Immediately postoperatively
- 4-6 weeks postoperatively
- 4 months intervals for a year
- Yearly for 5 years
- Every 2 years

CONCLUSIONS

Pituitary adenomas occurs with a wide spectrum of clinical features

Should be managed between different specialists

Neuro-ophthalmological manifestations are frequent and varied

Physicians must be aware about these in order to refer patients to ophthalmologist for early diagnosis and treatment

Radiotherapy –

- At the midpoint and end of radiotherapy
- 3 months interval for a year
- 6 months interval for a year
- Yearly

Evaluation –

- Visual acuity
- Fundoscopy
- Visual field
- Imaging

Case Report

30-year-old male with painless progressive vision loss.

Chief Complaint: Painless central vision loss in the left eye.

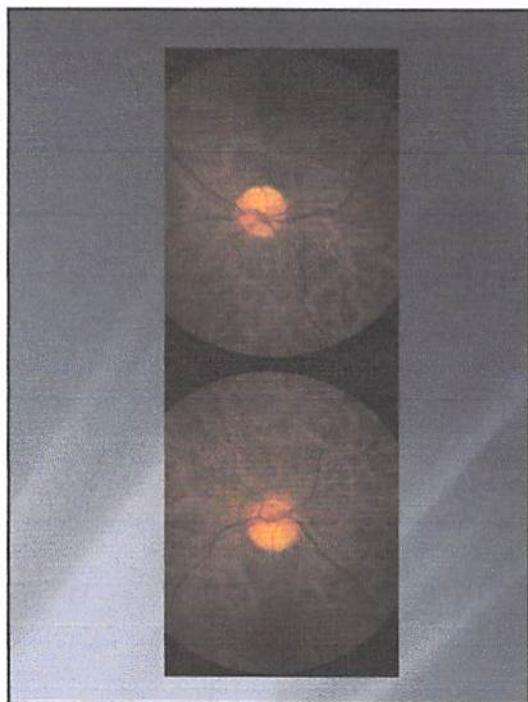
History of Present Illness: This adult female presented to the comprehensive ophthalmology clinic with painless central vision loss in the left eye, recently dates back on an employee screening examination. She had noted difficulty with reading and an opaque "white light" in her central vision of the left eye.

Past Ocular History: Negative for surgery or trauma	Medications: Eye drops, systemic:
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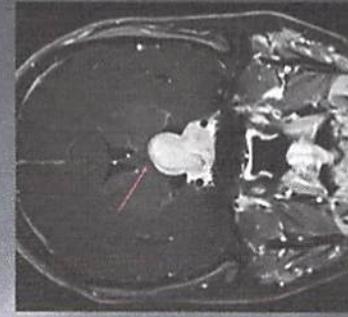
Past Medical History: Hypothyroidism	Review of Systems: Negative, including pain, sight changes, mood changes, diabetes, or loss of libido.
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Ocular Examination

- Visual acuity without correction: 20/25 right eye (OD), 20/125 with eccentric fixation left eye (OS)
- Intraocular pressure: 10 mmHg OD, 12 mmHg OS
- Pupils: 5 → 3 mm in both eyes (OU), brisk reaction OU, 0.3 log-unit relative afferent pupillary defect OS
- Confrontation visual fields: temporal loss OD, central scotoma OS
- Anterior segment: 1+ nuclear sclerosis OU, otherwise normal



This figure shows the fundus photographs of the patient. The top image shows the optic nerve head and surrounding retinal vessels. The bottom image shows the macula and surrounding retinal tissue. These findings are consistent with a diagnosis of papilledema and maculopathy, respectively.



This figure shows an axial MRI scan of the brain. The optic nerves and optic chiasm are visible. There is no evidence of structural abnormalities, such as tumors or aneurysms, which can cause papilledema. The findings are consistent with a diagnosis of papilledema due to increased intracranial pressure.

