# CASE 1: UNCOMMON EVOLUTION OF HYPOPITUITARISM AFTER PROLACTINOMA APOPLEXY

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## PATIENT HISTORY

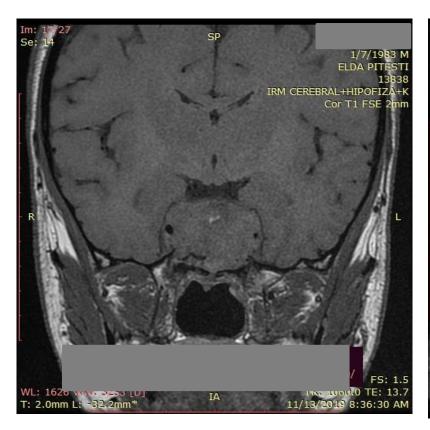


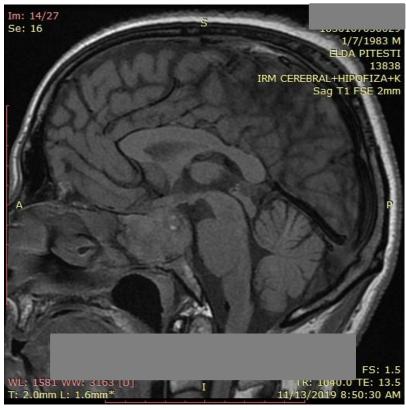
•	Summer	of 2019	low appetite
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<ul> <li>No:</li> </ul>	/ 4 <sup>th</sup> , 2019	headache	admitted to local hospital N	SAID
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- Nov 6<sup>th</sup>, 2019 double vision
- Nov 10<sup>th</sup>, 2019 severe headache transferred to regional hospital head CT
- Nov 13<sup>th</sup>, 2019 pituitary apoplexy transferred to our hospital



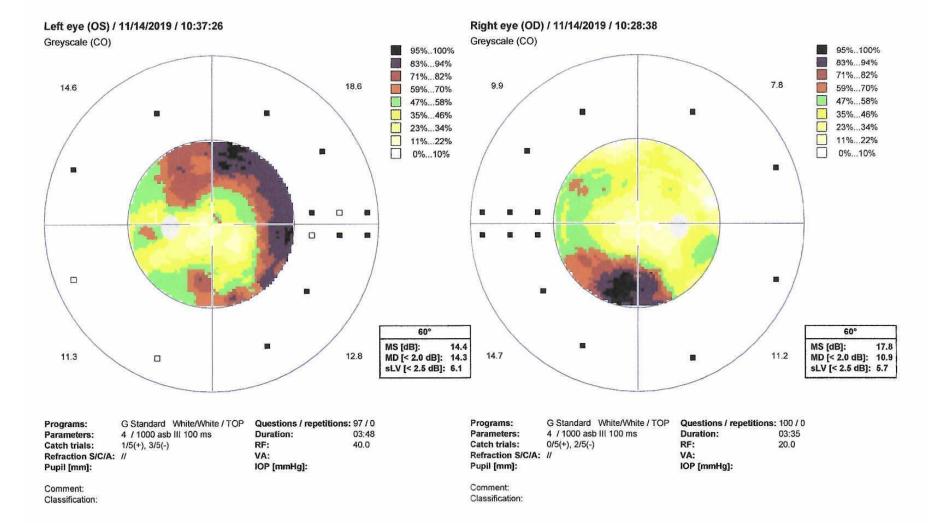




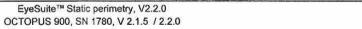


### 1 4 NOV 2019













# What should we do next?

Surgery?
Medical treatment?



Dexketoprofen

Acetaminophen

Dexamethasone

Manitol

Cabergoline 2 mg/wk

Prednisone 5mg/d

Levothyroxine 50 μg/d



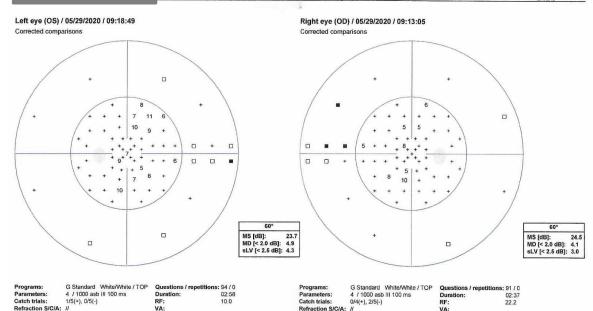






2 weeks later

2 9 MAI 2020



**OCTOPUS®** 

IOP [mmHg]:

Pupil [mm]:

Classification:

Comment:

EyeSuite™ Static perimetry, V2.2.0 OCTOPUS 900, SN 1780, V 2.1.5 / 2.2.0

Pupil [mm]:

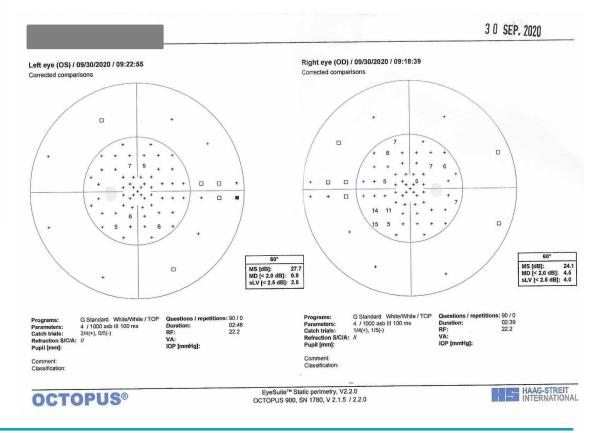
Classification:

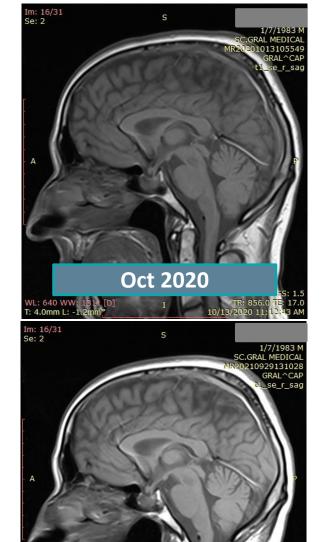
Comment:



IOP [mmHg]:

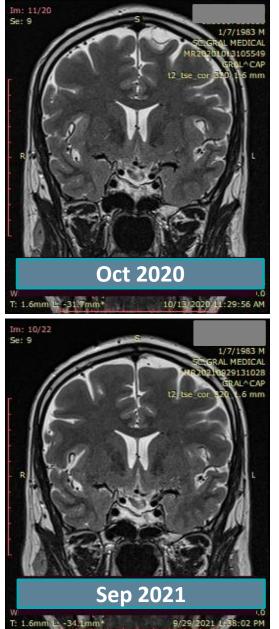




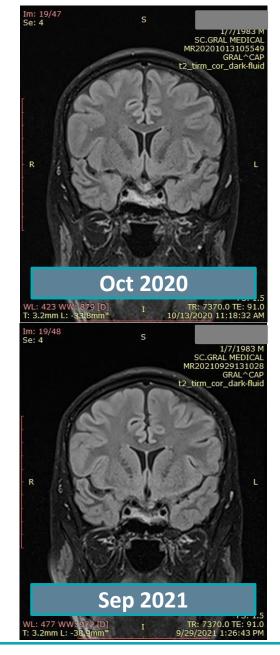


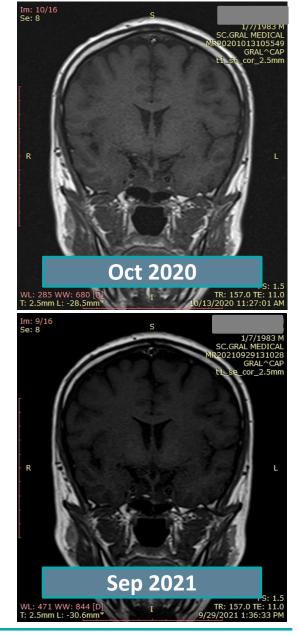
**Sep 2021** 

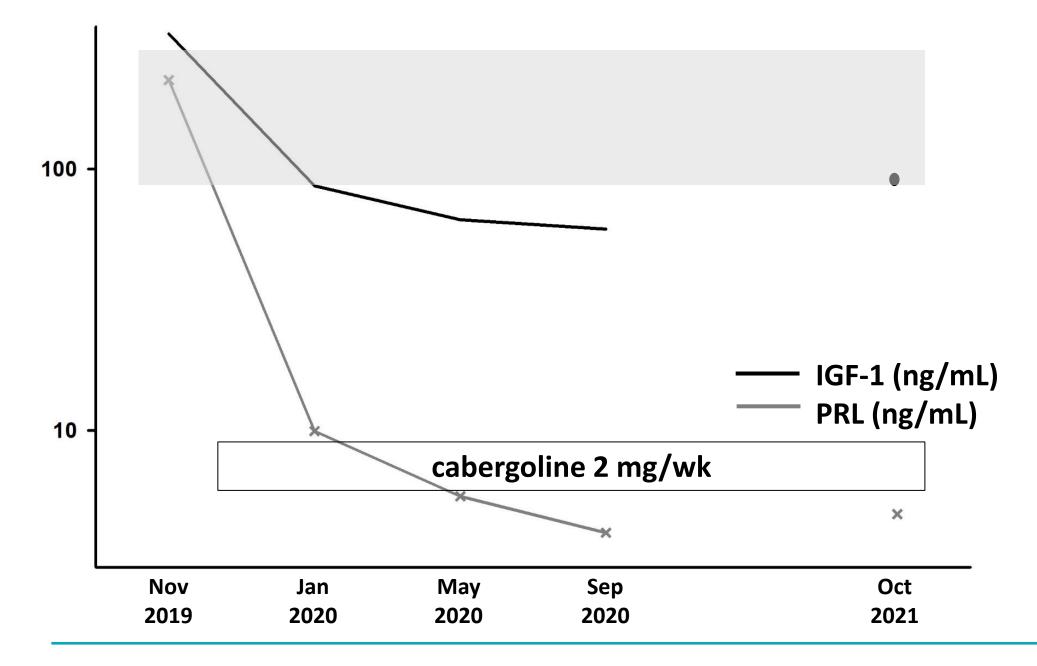
WL: 485 WW: 99L[D] T: 4.0mm L: 1.5mm\*



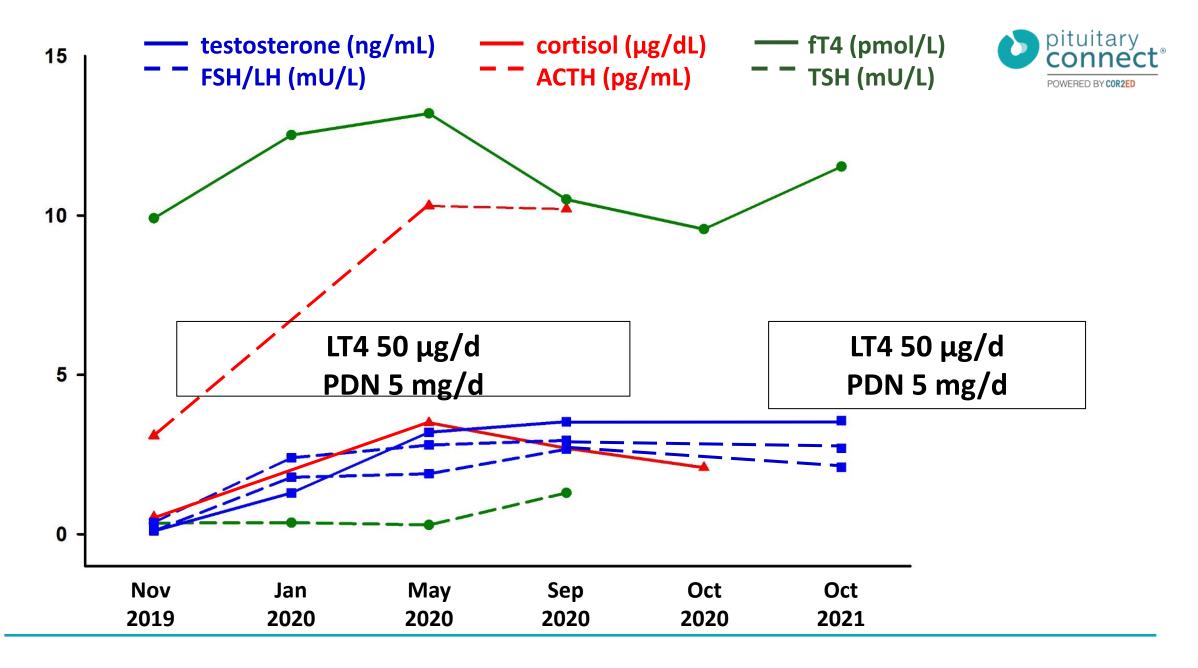
T: 1.6mm L -34.1mm\*













Should we continue cabergoline?

Should we continue substitution treatment?

# **CONCLUSIONS**



- Common hypopituitarism:  $GH \rightarrow FSH/LH \rightarrow TSH \rightarrow ACTH$
- Pituitary apoplexy