

# A case of paediatric GH-secreting pituitary adenoma apoplexy

C. Partenope<sup>1</sup>, D. Gallo<sup>1</sup>, S. Recupero<sup>1</sup>, C. Baldoli<sup>2</sup>, P. Mortini<sup>3</sup>, M. Losa<sup>3</sup>, G. Weber<sup>1</sup>,  
G. Barera<sup>1</sup>, G. Pozzobon<sup>1</sup>



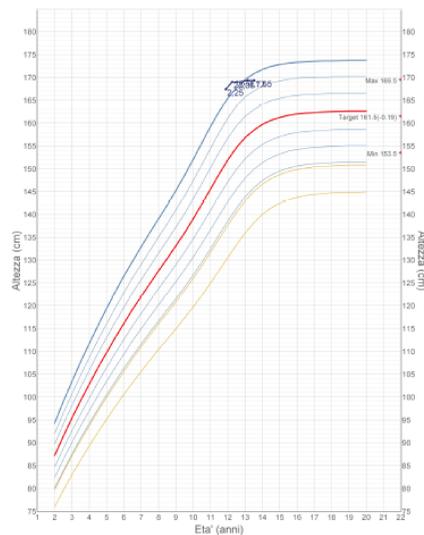
<sup>1</sup>Pediatric Department, San Raffaele Hospital, Milan, Italy  
<sup>2</sup>Neuroradiology Department, San Raffaele Hospital, Milan, Italy  
<sup>3</sup>Neurosurgery Department, San Raffaele Hospital, Milan, Italy

## Background

Paediatric pituitary adenomas comprise rare but challenging pathologies in children and adolescents related to their endocrine and neurological features. Pituitary apoplexy is a clinical syndrome caused by hemorrhage or infarction of the pituitary gland and is predominantly seen in patients with pituitary adenomas.

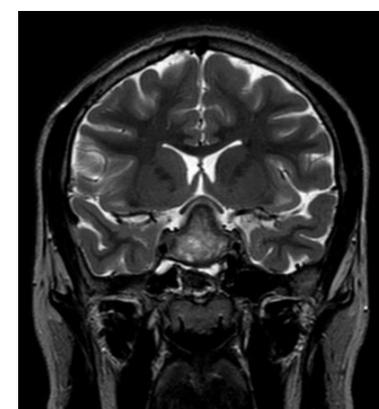
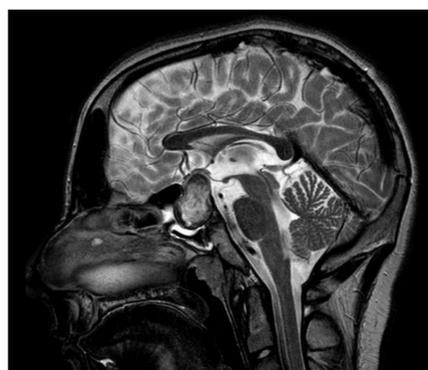
## Methods

A 11-year-old girl presented with fever and sudden headache with vomiting. She abruptly developed bilateral hemianopia. On suspicion of infectious meningitis, lumbar puncture and brain MRI were performed on admission.



## I.N. , ♀

- The physical exam was normal, without neurological abnormalities except for visual disturbances.
- Auxological parameters were >95th percentile (height 2.25 SDS, weight 1.85 SDS).
- Biochemistry laboratory and cerebrospinal fluid analysis were normal.
- Brain MRI showed a 33 mm-sellar and suprasellar mass with pituitary stalk and optic chiasm dislocation and third ventricle compression; it has heterogeneous signal features and contrast enhancement, highly suggestive of hemorrhagic transformation of a pre-existing pituitary adenoma.
- The endocrine investigations demonstrated central hypothyroidism, hypocortisolism, Growth Hormone (GH) deficiency and diabetes insipidus.



- The lesion was surgically removed with a transphenoidal approach.
- The histological examination and immunohistochemical staining were compatible with somatotroph GH-secreting adenoma, diffuse necrosis and hemorrhage areas.
- In the postoperative period, multiple pituitary hormone replacement therapy (hydrocortisone, desmopressin, estrogen and L-thyroxine) was started with good response. GH substitution was started 7 months later.

## Conclusions

- Pituitary apoplexy in children and adolescents is a rare entity that requires rapid and adequate treatment to prevent a life-threatening situation.
- Pituitary failure may develop with the effect of adenoma itself or following surgical excision.
- Clinical and laboratory signals may be suggestive and MRI neuroimaging is fundamental for diagnosis.

## References

1. F. Guaraldi et al., "Paediatric Pituitary Adenomas: A Decade of Change", *Horm Res Paediatr* 2014;81:145–155
2. H. Zijlker et al., "Pituitary Adenoma Apoplexy in an Adolescent: A Case Report and Review of the Literature", *Clin Res Pediatr Endocrinol* 2017;9(3):265-273
3. M. Ozcetin et al., "A pediatric case of pituitary macroadenoma presenting with pituitary apoplexy and cranial nerve involvement: case report", *Turk Pediatri Ars* 2016; 51: 162-5

