

## Case study

### Cataract in hypoparathyroidism: a case study.

#### ABSTRACT:

Ophthalmologic involvement in hypoparathyroidism can affect each of the components of the eye. However, it has a particular tropism for the lens, the prevalence of cataract reaches approximately 50% of hypoparathyroid subjects. Cataracts are sometimes indicative of these affections. Its detection is a constant concern in the monitoring of patients, because it constitutes a severity index of the disease, and is likely to benefit from a specific management.

In this case, we want to describe the aspect of cataract secondary to hypoparathyroidism, which is an etiology rarely encountered in ophthalmology, also describe its pathogenesis and report the other ophthalmological manifestations linked to this etiology which reflect a certain element of severity in order to detect and prevent them.

**Keywords** : Cataract, hypoparathyroidism, hypocalcemia.

#### 1. INTRODUCTION :

Insufficient production or action of parathyroid hormone determines a situation of chronic calcipenia which is clinically expressed in the form of acute or chronic signs of neuromuscular hyperexcitability or tetany.

The ophthalmologic involvement of hypoparathyroidism is likely to alter each of the constituent parts of the eye. However, it has a particular tropism for the lens, the prevalence of cataract reaches approximately 50% of hypoparathyroid subjects (1). Cataracts are sometimes indicative of these affections. Its detection is a constant concern in the monitoring of patients, because it constitutes a severity index of the disease, and is likely to benefit from a specific management. Big questions persist as to its pathogenesis.

#### 2. CASE REPORT:

We report the case of a 37-year-old patient followed for a familial hypercalciuric hypocalcemia as well as for a renal lithiasis and who consults for a progressive decrease in visual acuity in his right eye.

On physical examination, the patient had stable vital signs. The ophthalmological examination revealed a visual acuity of 20/32 in his right eye increasing to 20/20 after correction, and a visual acuity of 20/20 in the left eye.

The Intraocular pressure is normal in both eyes and the oculomotor examination is without abnormality.

The slit lamp examination is normal in the left eye and revealed in the right eye:

In diffused lighting: a clear cornea, a normal anterior chamber and a dusty cataract (**Figure 1**). In section: the opacities are radiary in an arc of a circle and sit at the level of the epinuclei (**Figure 2**). The dilated fundus is normal in both eyes.

Considering his good corrected visual acuity, the patient is adapted in corrective glasses.

### 3. DISCUSSION:

Hypoparathyroidism is a deficiency of parathyroid hormone often caused by autoimmune disease, genetics, iatrogenic damage or removal of glands during thyroidectomy or parathyroidectomy, but the cause is often still imperfectly determined (**1**). Measurement of parathyroid hormone (PTH) levels is necessary for diagnosis. Treatment includes calcium and vitamin D supplementation.

Familial hypercalciuric hypocalcemia or Autosomal dominant hypocalcemia (ADH) is most often due to activating mutations of the CaSR (calcium-sensing receptor) gene encoding the calcium-sensitive receptor leading to hypocalcaemia associated with an unsuitable parathyroid hormone level (low or normal), as during hypoparathyroidism, but also an abnormally high calciuria due to hypocalcemia.

The most common ophthalmologic consequence of hypoparathyroidism is cataract. although this mechanism remains obscure (**2**). Cataract appears to be authentically linked to chronic calcipenia, but also undoubtedly involves, as in diabetes mellitus, disorders of the hydration of the crystalline lenses.

Experimentally as early as 1926 had been obtained in rats subjected to phases of hypo- and normocalcemia, concentric layers of opacities and clarities (**3**). Lens opacities have also been obtained in rats on a calcium and hyperphosphorus diet (**4**).

Cataract lesions therefore appear frequent in adults with genetic hypoparathyroidism (like our case) that were recognized late. Conversely, compared to 2064 controlled subjects, the risk of cataract and its age of onset did not differ significantly in the 688 subjects who developed post-surgical hypoparathyroidism and subjected to vitamin-calcium treatment (**5**), even though traditionally the prevalence of cataract affects approximately 50% of hypoparathyroid patients (**1**).

This endocrine cataract can be unilateral or bilateral and also nuclear and / or cortical and / or subcapsular (**6**). The evolution of cataracts is typically slow in cases of idiopathic hypoparathyroidism like our case and more rapid in cases of acute hypoparathyroidism (**7**).

The severity of cataracts appears to be related to both the severity and the duration of the hypocalcemia with hyperphosphatemia (**8**). Cataracts have been shown to occur in 28% of patients with hypocalcaemia for more than 4 years (**9**).

The presence of cataract in a subject with hypoparathyroidism is an indication of severity of the disease. It imposes:

- ✓ The search for other trophic disorders: evaluation of the dentition, dental enamel, search for calcifications or ossifications of the soft parts, detection of nephrocalcinosis and nephrolithiasis, calcifications of the central gray nuclei or more diffuse of the brain (Fahr syndrome), ideally by abdominal and brain computed tomography (CT) scans without injection.

- ✓ The intensification of therapeutic management to obtain a metabolic balance as perfect as possible, normalizing calcium levels, phosphatemia, phosphocalcic blood product, calciuria, at the cost of vitamin-calcium therapies, sometimes thiazides, or even recombinant PTH.

Other ocular manifestations related to hypoparathyroidism have been described in the literature:

- ✓ **Papilledema:** Much less often reported, papilledema appears mainly in children and is observed in both hypo- and pseudo-hypoparathyroidism (7- 9). Bilateral, often revealing, it is actually reversible with correction of hypocalcemia. Papilledema is associated with alterations in axoplasmic conduction associated with hypocalcemia and stasis (10 - 11).
- ✓ Weiss's sign and spasm of the oculomotor muscles: Neuromuscular hyperexcitability linked to hypocalcemia is traditionally demonstrated by Weiss's sign: the percussion of the external angle of the orbit determines contracture of the orbicularis and of fact a zipper of the eyelids. In a 13-year-old adolescent girl with pseudo-hypoparathyroidism, irregular spasmodic eye movements, predominantly horizontally, were also reported during cataract surgery. The signs were precipitated by a state of respiratory alkalosis associated with hypocalcemia, and corrected with continuous infusion of calcium gluconate (12).
- ✓ Palpebral, orbital and sclera calcifications or ossifications: They have also been reported, forming part of the calcium deposits or ossifications of the soft tissues, characteristic of these situations (13).
- ✓ Keratoconjunctivitis: An exemplary inventory of the ocular signs of APECED syndrome or Autoimmune Polyendocrinopathy Candidiasis Ectodermal Dystrophia (a condition, transmitted in an autosomal recessive manner which is typically characterized by Whitaker's triad: hypoparathyroidism, adrenal insufficiency, recurrent mucocutaneous candidiasis), evolution and prognosis was carried out in Helsinki, reported in 2000. In a series of 69 patients, keratoconjunctivitis was present in 17 subjects (25% of cases) (14).
- ✓ More rarely: sicca syndrome, iridocyclitis, severe retinal sufferings and optic atrophies (15).

Our patient has only a unilateral cataract, the opacities are radiary in an arc of a circle and sit at the level of the epinuclei, the rest of his ophthalmological examination did not reveal any other characteristic signs of hypoparathyroidism described above, hence the interest of close monitoring in order to detect and prevent them.

#### 4. CONCLUSION:

Diversity of expressions and mechanisms characterize the ophthalmologic complications associated with hypoparathyroidism. However, the most common ophthalmologic consequence of this etiology is cataract.

These manifestations reflect a certain element of severity. It is important to detect and prevent them.

#### CONSENT

All authors declare that written informed consent was obtained from the patient for publication of this case report and preserved by the authors.

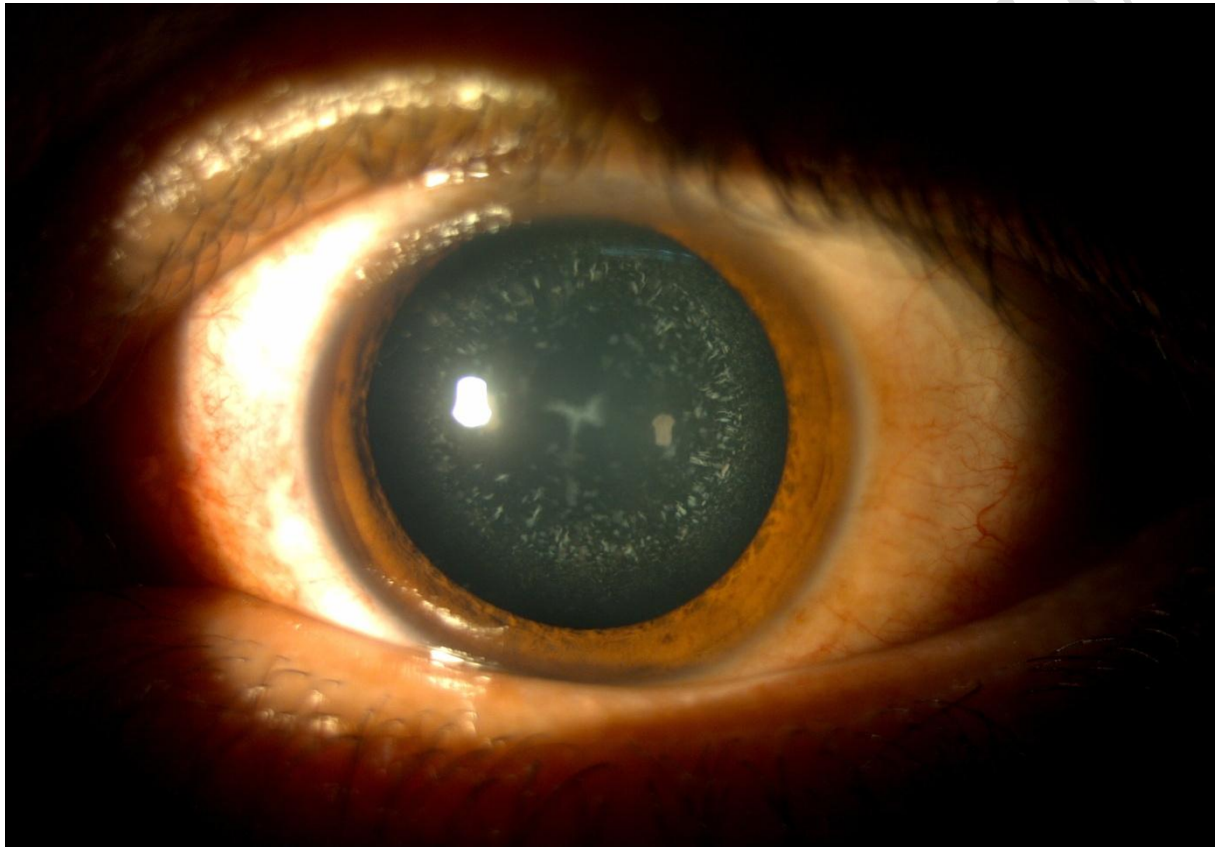
### **ETHICAL APPROVAL**

As per international standard or university standard written ethical approval has been collected and preserved by the authors.

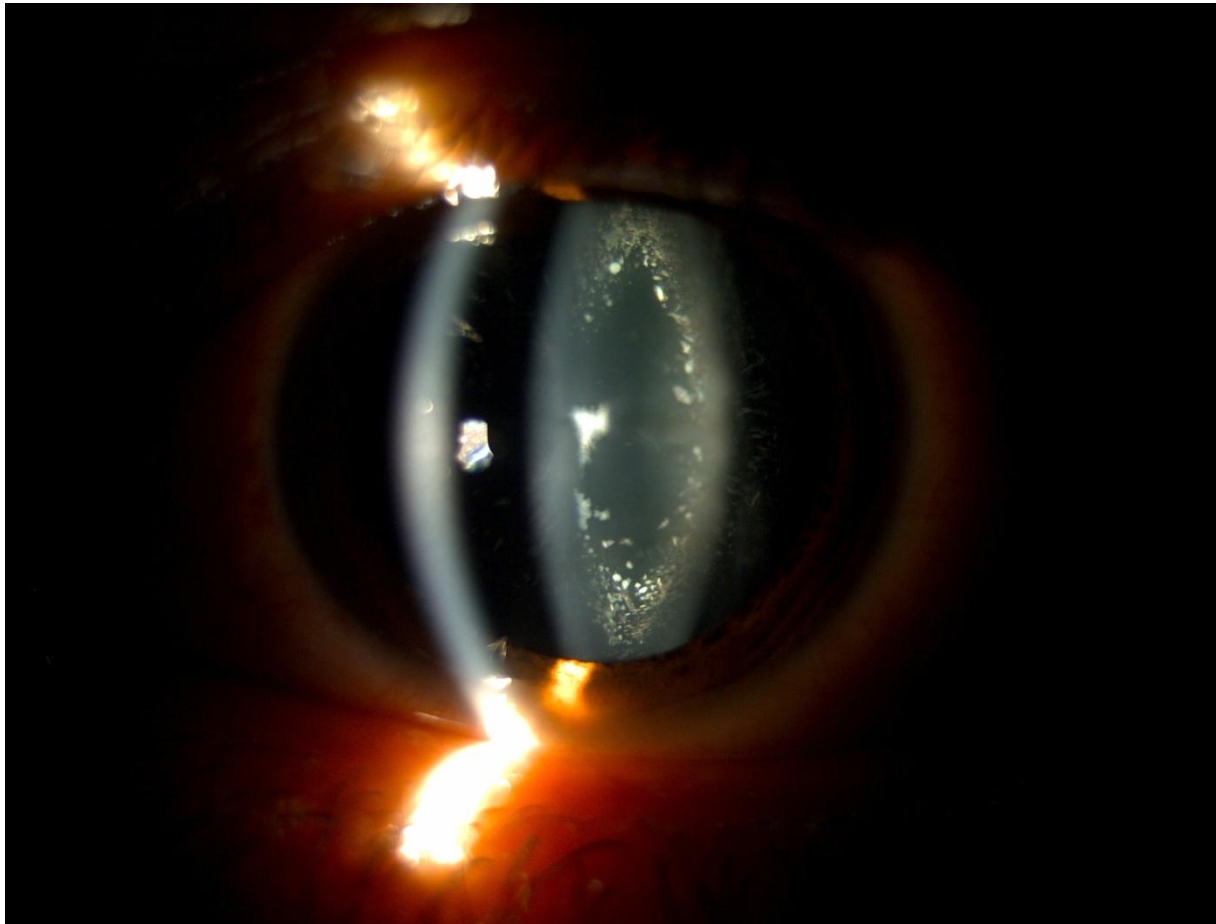
### **COMPETING INTERESTS**

Authors have declared that no competing interests exist.

### **FIGURES:**



**Figure 1:** Slit lamp examination of the right eye showing a dusty cataract in diffused lighting.



**Figure 2:** In section: the opacities are radiary in an arc of a circle and sit at the level of the epinuclei.

#### REFERENCES:

1. J.L. Wémeau, E. Proust-Lemoine, M. Contestin, A. Ryndak, J.F. Rouland Manifestations ophtalmologiques des hypo - et pseudohypoparathyroïdies. *Correspondances en Métabolismes, Hormones, Diabètes et Nutrition* - Vol. XXI - n° 12 - janvier-février 2017
2. Sawicki A. Postoperative hypoparathyroidism: risk of complications. *Pol Tyg Lek* 1991;46:815—7.
3. Goldmann H. 677 Experimentelle Tetaniekatarakt. *Graefes Arch Ophtal* 1929;122:146-97.
4. Von Bahr G. Studies on the etiology and pathology of zonular cataract. *Acta Ophtalm* 1936;4 Suppl X1:1-236.
5. Underbjerg L, Sikjaer T, Mosekilde L, Rejnmark L. Postchirurgical Hypoparathyroidism risk of fractures, psychiatric diseases, cancer, cataract and infections. *J Bone Miner Res* 2014;29(11)2504-10.
6. Arlt W, Fremmery C. Well-being, mood and calcium homeostasis in patients with hypoparathyroidism receiving standard treatment with calcium and vitamin D. *Eur J Endocrinol* 2002;146:215—22.

7. Haviv YS, Safadi R, Zamir E. A rapidly progressive cataract in a patient with autoimmune hypoparathyroidism and acute liver renal failure. *Am J Nephrol* 1999;19:523—6.
8. Stein R, Godel V. Hypocalcemic cataract. *J Pediatr Ophthalmol Strabismus* 1980;17(3):159-61.
9. Mundy GR. Calcium Homeostasis: Hypercalcemia and Hypocalcemia. Martin Dunitz London. 1990:1-16.
10. Fowler WM Jr, LING S. Convulsions and Papilledema in a Child with idiopathic hypoparathyroidism. *Calif Med* 1961;95:180-2.
11. Maheshwari M, Rani RP, Reddy AP. Visual disturbances as a presenting feature of pseudohypoparathyroidism. *Indian J Endocrinol Metab* 2013;17 (Suppl.1):S219-20.
12. Suder RA, Singh M. Pseudohypoparathyroidism: a series of three cases and an usual presentation of ocular tetany. *Anaesthesia* 2006;61(4):394-8.
13. Wong S, Zakov ZN, Albert DM. Scleral and choroidal calcifications in a patient with pseudohypoparathyroidism. *Br J Ophthalmol* 1979;63(3):177-80.
14. Merenmies L, Tarkkanen A. Chronic bilateral keratitis in autoimmune polyendocrinopathy-candidiasis ectodermal dystrophy (APECED). A long-term follow-up and visual prognosis. *Acta Ophthalmol Scand* 2000;78(5):532-5.
15. Chang B, Brosnahan D, McCreery K, Dominguez M, Costigan C. Ocular complications of autoimmune polyendocrinopathy syndrome type 1. *J AAPOS* 2006 ;10(6):515-20.