

THYROID-RELATED BONE LESIONS: clinical illustration



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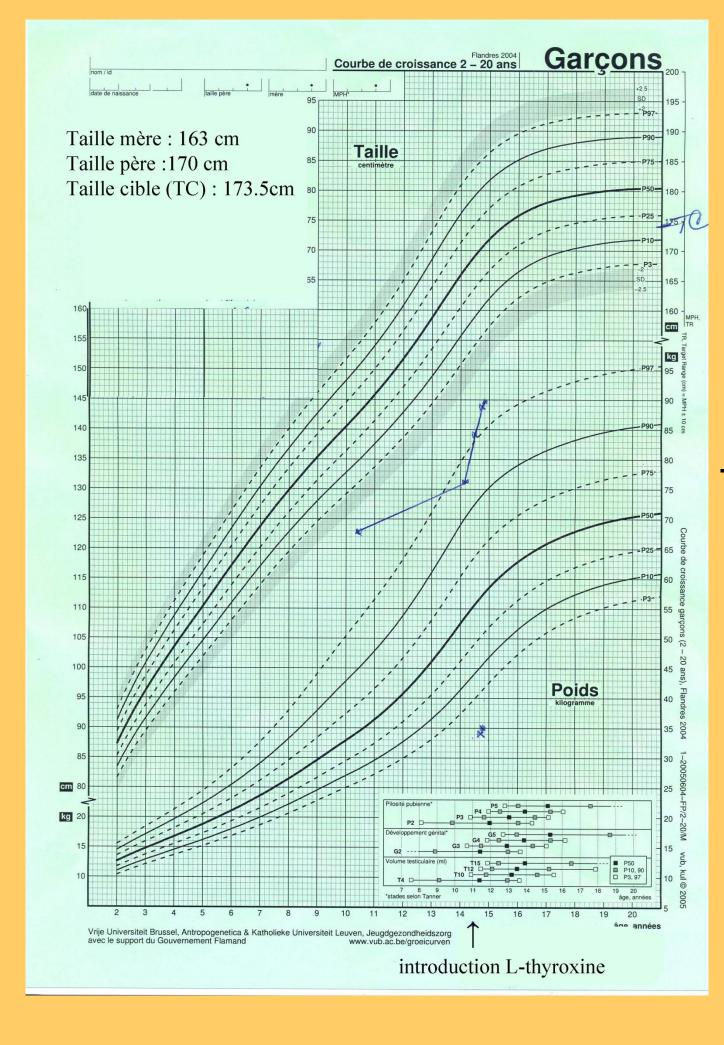
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INTRODUCTION:

Primary hypothyroidism is a common endocrine disease. Although the diagnosis is often simple, some clinical presentations are rarer. The association of short stature, epiphysis bone lesions and severe retarded bone age in children should lead to the diagnosis of severe, long-standing acquired hypothyroidism.

CLINICAL CASE:

- A 14-year-old boy presents bilateral hip pain that have been evolving for 3 days (already intermittent for 2 years)
- Pain increase with mobilization and weight bearing.
- No fever and no history of trauma.



- Clinical examination:
 - -internal and external rotation of hips provoked pain
 - -no neurological deficiency.
 - -very short proportionate stature (133cm)
 - -abnormal phenotypic features of the patient : relatively macrocephaly and macroorchidism

Additionnal tests:

- Biology:
 - no inflammatory syndrome;
- low level of T3 and T4 hormones;
- very high TSH level in serum (312mUI/l);
- no antibodies against thyroid;
- gonadotrophin, cortisol normals;
- growth hormones (IGF-1 and IGF-BP3) a little bit reduced;
- → diagnosis of severe acquired hypothyroidism

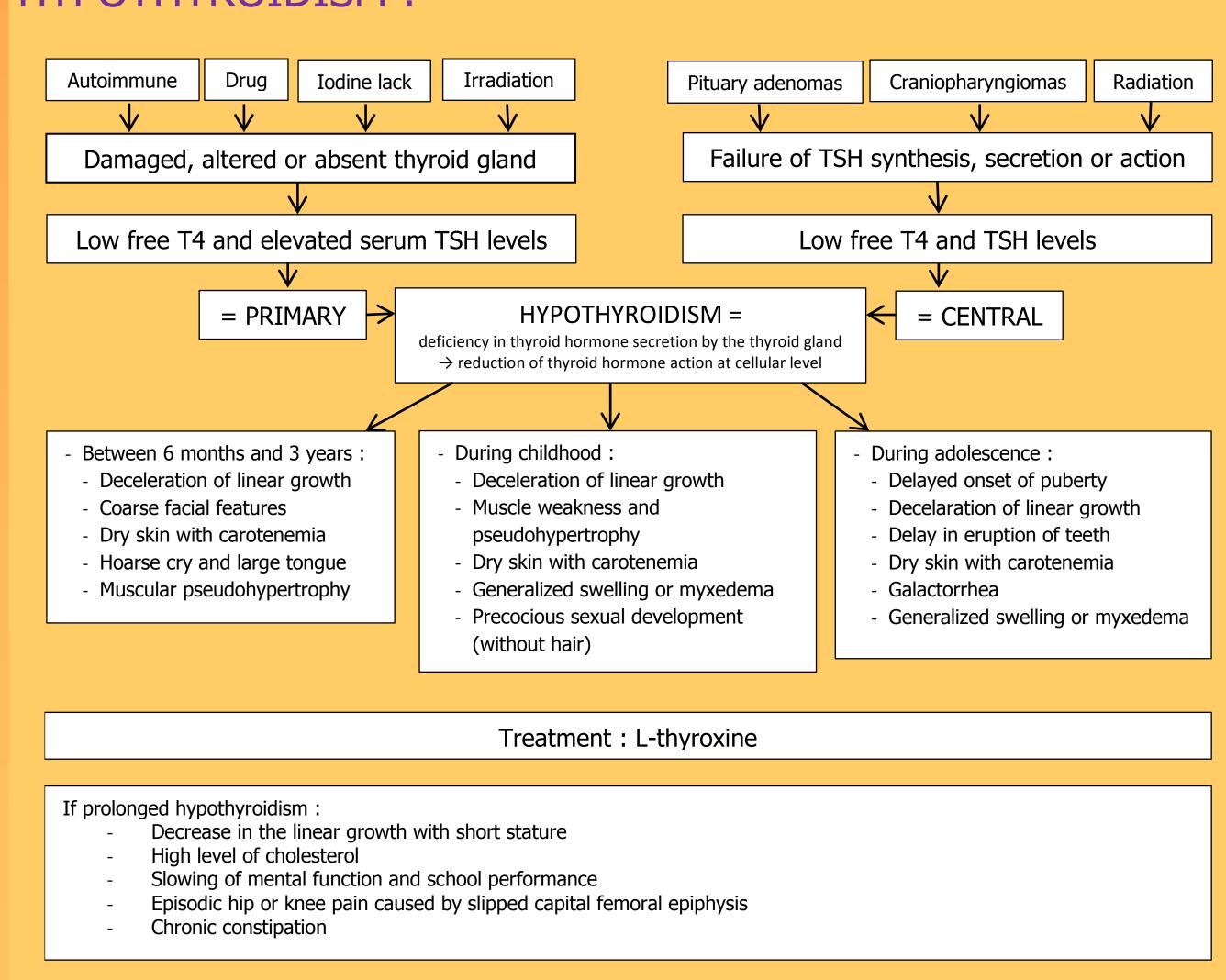
- Radiology:

- condensation and fragmentation of the epiphyseal nuclei bilateraly with irregular aspect of cotyles?
- → diagnosis of severe and bilateral Legg-Calve-Perthes disease (infirmed aftewardr)



- US: atrophic thyroid
- Bone maturation: 6 years (Graulich and Pyle atlas)
- <u>Treatment</u>: L-thyroxine with progressive dosis
- <u>Evolution</u>: favorable with linear growth acceleration

HYPOTHYROIDISM:



Bone-related lesions:

- retarded growth and bone maturation with delayed ossification
- reduced postnatal growth and bone mineralization
- -delayed closure of the cranial sutures in early life
- -greater thickness in cortical bone with ↑ risk of fractures -disturbances of endochondral ossification

THYROID AND BONE:

Actions of T3:

- During bone formation : stimulation of osteoblast proliferation, differentiation and apopotosis
 - ↑ the expression of osteocalcin, type 1 collagen, alkalin phosphatase, IGF1,...
- During bone resorption : † expression of IL6 and PGE2 (differenting factors of osteoclast lineage)
 - synergistic action with PTH and vitamin D
 - activation of RANK, a key step in osteoclastogenesis
 - → anabolism during development and catabolism after bone maturation

Actions of TSH

- negative regulator of bone turnover
- enhanced bone remodeling and osteoporosis

-Mains effects of thyroid disorders on bone :

Hypothyroidism	Hyperthyroidism
Maximum	Minimum
↓	\uparrow
Prolonged (mainly	Reduced (mainly
resorption phase)	formation phase)
↓	↑
\downarrow	↑
↓	↑
↓ and disproportionate	↓ and proportionate
	Maximum Prolonged (mainly resorption phase)

CONCLUSION:

Relationships between endocrine-associated biochemical abnormalities and altered growth plate structure and function are known.

Hypothyroidism in children results in short stature and delayed bone maturation with a fragmentation of the femoral heads, misdiagnosed as Legg-Calve-Perthes disease, as in this clinical case.

Recuperation of the expected weight and of the bone mineralization depends on the duration and severity of the lack of hormones.