

HASHITOXICOSIS: A RARE DIAGNOSIS IN CHILDHOOD

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INTRODUCTION-AIM:

- ✓ Patients with Hashimoto's thyroiditis are often euthyroid or may experience subclinical or true hypothyroidism
- ✓ 5 to 10% of children may present however, a transient phase of hyperthyroidism, called Hashitoxicosis. This is caused by the release of stored, preformed thyroid hormones in blood circulation, due to thyroid gland inflammation
- ✓ Differential diagnosis from Grave's disease may be proved difficult to do and requires detailed history, physical examination, laboratory assessment and systematic follow-up.

Aim: The present three cases that highlight the diagnostic tools for the diagnosis of Hashitoxicosis and its distinction from Grave's disease.

PATIENTS-METHODS

Three female patients were referred to our department:

- ❑ The first 6 1/2 y.o., suffered from Type I diabetes mellitus and during a checkup, severe hypothyroidism was detected due to Hashimoto's thyroiditis. Substitution therapy with L-Thyroxine was given. Subsequently she developed subclinical hyperthyroidism and then returned to hypothyroid status
- ❑ The second 9 6/12 y.o., passed from a hyperthyroidic phase to an euthyroidic one
- ❑ The third 12 4/12 y.o., first experienced hypothyroidism, then subclinical hyperthyroidism and later became euthyroid

RESULTS

- ❖ In all cases, thyroid ultrasound showed mainly structural gland heterogeneity
- ❖ The presence of positive TSI's in the first case, made it even more difficult to diagnose. For this reason, a Tc 99m scintigraphy was required, showing a diffuse and slightly increased intake in both lobes, entity compatible with Hashitoxicosis
- ❖ All three patients showed no evidence of clinical hyperthyroidism and were clinically checked every two weeks
- ❖ Subclinical Hyperthyroidism lasted from 30 to 90 days

	1 st visit	After 2 months	After 7 months	After 10 months	After 12 months
1st patient	1) TSH:201,1 2) FT4: 0,29 3) Anti-TPO: 96 4) Anti-Tg: 1193	1. TSH: 1,49 2. T4: 10,49 3. T4: 25 µg x1	1. TSH: <0,01 2. FT4: 2,50 3. Anti-TPO: 127 4. Anti-Tg:196 5. TSI: positive	1. TSH: 34,63 2. FT4: 0,79 3. T3: 143	
2nd patient	1) TSH: 0,040 2) FT4: 25,72 3) Anti-TPO: 69,7 4) Anti-Tg:226 5) TSI: negative	1. TSH: 12,439 2. FT4: 14,56 3. T4: 50 µg x1	1. TSH: 2,94 2. FT4: 17,78 3. T4: 50x1		
3rd patient	1) TSH: 7,68 2) T4: 7,06 3) T3: 138 4) Anti-TPO: 219 5) Anti-Tg: 826	1. TSH: 1,320 2. FT4: 1,34 3. T4: 50 µg x1		1. TSH: 0,083 2. FT4: 1,34 3. Anti-TPO: 129 4. Anti-Tg: 2655 5. TSI: negative	1. TSH: 1,330 2. FT4: 1,18 3. Anti-TPO: 67,67 4. Anti-Tg: 1024
Units : TSH in mIU /lt FT4 in ng/dl Anti-Tg, Anti-Tpo in IU/lt					

CONCLUSIONS

- Hashitoxicosis is a rare event that needs to be recognized the earliest possible
- Tc ^{99m} scintigraphy is not always necessary for the final diagnosis
- Prospective careful clinical monitoring is essential
- In patients with cardiovascular signs and symptoms, propranolol is recommended

REFERENCES

1. Hashitoxicosis in children: clinical features and natural history. Nabhan ZM, Kreher NC, Eugster EA J Pediatr. 2005;146(4):533
2. Frequency of Hashimoto's thyroiditis antecedents in the history of children and adolescents with graves' disease. Wasniewska M, Corrias A, Arrigo T, Lombardo F, Salerno M, Mussa A, Vigone MC, De Luca F Horm Res Paediatr. 2010;73(6):473. Epub 2010 Apr 15

