Toxic multinodular goitre: a surprising finding

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DESCRIPTION
A 16-year-old healthy adolescent boy was referred to the paediatric endocrinology clinic because of multiple thyroid nodules detected by cervical ultrasound, in the context of cervical lymphadenopathies. There was no family history of thyroid disease. He denied recent infections, asthenia, weight loss, sweating, palpitations, mood or sleep disturbances, dysphagia or dysphonia. At physical examination, an enlarged, irregular and fibroelastic thyroid, with a predominant right lobe, was identified. The remaining examination was normal.

The analytical profile was thyroid stimulating hormone (TSH) <0.01 uUI/mL (normal 0.5–4.8 uUI/mL), free triiodothyronine (FT3) 7.27 pg/mL (normal 2.3–4.2 pg/mL) and free thyroxine (FT4) 2.02 ng/dL (normal 0.8–2.3 ng/dL). Thyroid antibodies were negative. Cervical ultrasound revealed an enlarged right thyroid lobe due to the presence of multiple mixed nodules with similar characteristics: the biggest one, 5 cm diameter, predominantly cystic with a peripheral ring of solid isoechoic parenchyma (figure 1A); the second characteristically mixed nodule, 2.2 cm diameter (figure 1B) and the third one, 1.2 cm diameter, predominantly solid with small cystic areas (figure 1C). Thyroid scintigraphy showed hyperactive multinodular goitre with predominance of autonomous hot nodules on the right lobe, suggestive of toxic multinodular goitre (TMNG) (figure 2A–C). A fine needle aspiration biopsy of thyroid predominantly solid nodule, with ultrasound guidance, was performed to exclude malignancy but the cytological examination was non-diagnostic. The treatment options were discussed, with patient and family, and the surgical approach was followed due to the non-diagnostic result of biopsy and the high volume goitre. The patient started oral thiamazole to achieve euthyroidism and preparing for surgery.

Thyroid nodules are a frequent incidental finding with an incidence between 9.4% and 27.0%.1 In contrast to adults, TMNG is an uncommon thyroid disease in paediatric age. The presence of hyperthyroidism determines the need for a definitive therapy in multinodular goitre, and a correct evaluation must be established before deciding between surgery or radioactive iodine. The diagnosis of TMNG (overt or subclinical) should always be excluded in patients with physical examination or ultrasound findings consistent with nodular thyroid disease.2 3

Learning points
► Thyroid nodules are relatively common in adolescents, usually asymptomatic and often incidentally detected. While multinodular goitre has almost always a good prognosis, malignancy needs to be excluded and a large nodule increases its probability.
► The first line investigation should include a thyroid ultrasound to evaluate the nodules’ characteristics and a thyroid stimulating hormone (TSH) determination. In the presence of a suppressed TSH, thyroid scintigraphy is indicated to differentiate between hyperfunctioning (hot) and hypofunctioning (cold) nodule.
► When hyperthyroidism is caused by an autonomous thyroid nodule, a definitive treatment is necessary such as radioactive iodine therapy or surgery. In children and adolescents, surgical resection must be considered in face of suspicious ultrasound findings, clinical presentation, nodule size >4 cm and/or compressive symptoms.

Figure 1 Cervical ultrasound revelling multiples nodules in the right thyroid lobe. (A) Predominantly cystic nodule, (B) characteristic mixed nodule and (C) predominantly solid nodule.

Figure 2 Thyroid scintigraphy showing hyperactive multinodular goitre with hot nodules dominant on the right lobe. Scintigraphic features suggestive of toxic multinodular goitre. (A) Anterior, (B) right anterior oblique and (C) left anterior oblique.

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REFERENCES