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Familial hyperthyroidism due to mutations in TSH receptor

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. <u>Familial hyperthyroidism due to mutations in TSH receptor</u>. ORPHA:424

Familial non-autoimmune autosomal dominant hyperthyroidism (FNAH) is a rare hyperthyroidism (see this term) characterized by mild to severe hyperthyroidism, presence of goiter, absence of features of autoimmunity, frequent relapses while on treatment and a positive family history.

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