Familial thyroid dyshormonogenesis

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Familial thyroid dyshormonogenesis. ORPHA:95716

Familial thyroid dyshormonogenesis is a type of primary congenital hypothyroidism (see this term), a permanent thyroid hormone deficiency that is present from birth, which results from inborn errors of thyroid hormone synthesis.