DISEASE:
Familial thyroid dyshormonogenesis

NAME:	Familial thyroid dyshormonogenesis
DESCRIPTION:	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.
ORPHACODE:	95716
SYNONYMS:	Thyroid dyshormonogenesis
XREF(S):	Orphanet OMIM OMIM OMIM OMIM OMIM OMIM ICD-10 ICD-10

ANALYTE(S):	SLC5A5 TPO DUOX2 DUOXA2 TG IYD
CREATED:	13 May 2019 - 01:02
CHANGED:	22 Jun 2023 - 16:14

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