DISEASE:
Hypothyroidism due to deficient transcription factors involved in pituitary development or function

NAME:	Hypothyroidism due to deficient transcription factors involved in pituitary development or function
DESCRIPTION:	Hypothyroidism due to mutations in transcription factors involved in pituitary development or function is a type of central congenital hypothyroidism (see this term), a permanent thyroid deficiency that is present from birth, characterized by low levels of thyroid hormones caused by disorders in the development or function of the pituitary.
ORPHACODE:	226307
XREF(S):	Orphanet ICD-10
ANALYTE(S):	POU1F1 PROP1 HESX1 LHX3 LHX4
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RELATED CONTENT

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- Hypogonadotropic hypogonadism (33 genes)
- Short Stature (gene panel)

Related Laboratories

- Centre de Génétique-Institut de Pathologie et de Génétique (IPG)
- Centrum Medische Genetica - UZ Brussel VUB

Related Analytes

- HESX homeobox 1
- LIM homeobox 3
- LIM homeobox 4
- POU class 1 homeobox 1
- PROP paired-like homeobox 1

Related Gene Panels

- Hypogonadotropic Hypogonadism/Kallmann (61 genes) - ULG
- Hypogonadotropic hypogonadism (33 genes) - VUB
- Short Stature (46 genes) - IPG

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