DISEASE:
Hypothyroidism due to TSH receptor mutations

NAME:	Hypothyroidism due to TSH receptor mutations
DESCRIPTION:	A type of primary congenital hypothyroidism, a permanent thyroid hormone deficiency that is present from birth due to thyroid resistance to TSH.
ORPHACODE:	90673
XREF(S):	Orphanet OMIM ICD-10
ANALYTE(S):	TSHR
CREATED:	13 May 2019 - 01:02
CHANGED:	22 Jun 2023 - 16:14

RELATED CONTENT

Related Genetic Tests

- Hyperthyroidism (familial gestational or familial nonautoimmune, hypothyroidism, thyrotropin) - TSHR
- Thyroid disgenesis (38 genes)

Related Laboratories

- Centre de Génétique Humaine - Erasme ULB
- Centrum Medische Genetica - UZ Brussel VUB

Related Analytes

- thyroid stimulating hormone receptor

Related Gene Panels

- Thyroid disgenesis (38 genes) - VUB

Source URL: http://gentest.healthdata.be/disease/3174