

DISEASE:
Genetic transient congenital hypothyroidism

NAME:	Genetic transient congenital hypothyroidism
DESCRIPTION:	Genetic transient congenital hypothyroidism is a rare, thyroid disease characterized by a gene mutation induced, temporary deficiency of thyroid hormones at birth, which later reverts to normal with or without replacement therapy in the first few months or years of life.
ORPHACODE:	226316
XREF(S):	Orphanet
ANALYTE(S):	DUOX2
CREATED:	13 May 2019 - 01:02
CHANGED:	22 Jun 2023 - 16:14

Source URL: <http://gentest.healthdata.be/disease/1503>

RELATED CONTENT

Related Genetic Tests

- Thyroid disgenesis (38 genes)

Related Laboratories

- Centrum Medische Genetica - UZ Brussel VUB

Related Analytes

- dual oxidase 2

Related Gene Panels

- Thyroid disgenesis (38 genes) - VUB

Source URL: <http://gentest.healthdata.be/disease/1503>