DISEASE:
Familial gestational hyperthyroidism

NAME:	Familial gestational hyperthyroidism
DESCRIPTION:	A rare genetic hyperthyroidism characterized by hyperemesis gravidarum associated with hyperthyroidism due to hypersensitivity of the thyrotropin receptor to chorionic gonadotropin, in the absence of abnormally high serum chorionic gonadotropin levels. Clinical manifestations include severe nausea, vomiting, weight loss, tachycardia, excessive sweating, and hand tremor, but no signs of ophthalmopathy.
ORPHACODE:	99819
XREF(S):	Orphanet ICD-10 ICD-10 MeSH OMIM
ANALYTE(S):	TSHR
CREATED:	13 May 2019 - 01:02
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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