Opis choroby *

Definicja

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.

Dane

Klasyfikacja
Choroba

Kod ORPHA
99819

Kod OMIM
603373

Kod ICD10
E05.8

Kod ICD11
5A02.Y

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*Źródło

orphanet