

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

*Źródło

orphanet