

Opis choroby *

Definicja

A rare, genetic, primary orthostatic disorder characterized by dizziness, palpitations, fatigue, blurred vision and tachycardia following postural change from a supine to an upright position, in the absence of hypotension. A syncope with transient cognitive impairment and dyspnea may also occur. The norepinephrine transporter deficiency leads to abnormal uptake and high plasma concentrations of norepinephrine.

Dane

Klasyfikacja

Choroba

Synonimy

Familial orthostatic tachycardia due to
norepinephrine transporter deficiency
Rodzinny częstoskurcz ortostatyczny
spowodowany niedoborem transportera
norepinefryny
Zespół częstoskurcza posturalnego
spowodowanego niedoborem NET
Orthostatic intolerance due to NET deficiency
POTS due to NET deficiency

Kod ORPHA

443236

Kod OMIM

604715

Kod ICD10

I95.1

Kod ICD11

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[*Źródło](#)

orphanet