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 częstoskurczu 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