Opis choroby *

Definicja

Hereditary thrombophilia due to congenital histidine-rich (poly-L) glycoprotein deficiency is a rare, genetic, coagulation disorder characterized by a tendency to develop thrombosis, resulting from decreased histidine-rich glycoprotein (HRG) plasma levels. Manifestations are variable depending on location of thrombosis, but may include headaches, diplopia, progressive pain, limb swelling, itching or ulceration, and brownish skin discoloration, among others.

Dane

Klasyfikacja
Choroba

Synonimy
Hereditary thrombophilia due to congenital HRG deficiency
Dziedziczna trombofilia z powodu wrodzonego niedoboru HRG

Kod ORPHA
217467

Kod OMIM
613116

Kod ICD10
D68.5

Kod ICD11
3B61.0Y

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*Źródło

orphanet