Opis choroby *

Definicja

A rare, hereditary, hematologic disease characterized by an increase in hemoglobin, hematocrit and erythrocyte mass resulting in plethora or ruddy complexion, headache, dizziness, tinnitus and exertional dyspnea. In some cases, thrombophlebitis and arthralgia have also been reported.

Dane

Klasyfikacja
Choroba

Synonimy
Autosomal recessive secondary erythrocytosis not associated with VHL gene
Autosomalna recesywna wtórna erytrocytoza niezwiązana z genem VHL
Autosomalna recesywna wtórna erytrocytoza, nie typ Chuvasha
Autosomalna recesywna wtórna nadkrwistość, nie typ Chuvasha
Autosomal recessive secondary erythrocytosis, non-Chuvash type
Autosomal recessive secondary polycythemia, non-Chuvash type

Kod ORPHA
247378

Kod OMIM
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Kod ICD10
D75.1

Kod ICD11
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*Źródło

orphanet