Opis choroby *

Definicja

Hereditary folate malabsorption (HFM) is an inherited disorder of folate transport characterized by a systemic and central nervous system (CNS) folate deficiency manifesting as megaloblastic anemia, failure to thrive, diarrhea and/or oral mucositis, immunologic dysfunction and neurological disorders.

Dane

Klasyfikacja
Choroba

Synonimy
Congenital folate malabsorption
Wrodzone złe wchłanianie kwsau foliowego

Kod ORPHA
90045

Kod OMIM
229050

Kod ICD10
D52.8

Kod ICD11
5C63.1

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*Źródło

orphanet