

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 kwasu foliowego

Kod ORPHA

90045

Kod OMIM

229050

Kod ICD10

D52.8

Kod ICD11

5C63.1

*Źródło

orphanet