Opis choroby *

Definicja

A rare, genetic, congenital carbohydrate intolerance disorder characterized by lack of endogenous sucrase activity, marked reduction in isomaltase activity, and moderate decrease in maltase activity, and clinically manifesting with diarrhea, abdominal pain and bloating, failure to thrive.

Dane

Klasyfikacja
Choroba

Synonimy
CSID
CSID
Niedolerancja disacharydów
Wrodzona nietolerancja sacharozy
Wrodzone złe wchłanianie sacharazy-izomaltazy
Congenital sucrose intolerance
Disaccharide intolerance

Kod ORPHA
35122

Kod OMIM
222900

Kod ICD10
E74.3

Kod ICD11
5C61.2

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*Źródło

orphanet