

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

Kod ORPHA	Kod OMIM	Kod ICD10
35122	222900	E74.3

Kod ICD11
5C61.2

*Źródło

orphanet