Congenital Sucrase-Isomaltase Deficiency

National Cancer Institute

Source

National Cancer Institute. Congenital Sucrase-Isomaltase Deficiency. NCI Thesaurus. Code C128190.

An autosomal recessive genetic disorder caused by mutations in the SI gene, encoding sucrase-isomaltase, intestinal. The condition is characterized by malabsorption and osmotic diarrhea.