Congenital Disorder of Glycosylation Type Ic

National Cancer Institute

Source

National Cancer Institute. Congenital Disorder of Glycosylation Type Ic. NCI Thesaurus. Code C126869.

A congenital disorder of glycosylation sub-type caused by mutation(s) in the ALG6 gene, encoding dolichyl pyrophosphate Man9GlcNAc2 alpha-1,3-glucosyltransferase.