Congenital Disorder of Glycosylation Type Ig

National Cancer Institute

Source

National Cancer Institute. Congenital Disorder of Glycosylation Type Ig. NCI Thesaurus. Code C126873.

A congenital disorder of glycosylation sub-type caused by mutation(s) in the ALG12 gene, dol-P-Man:Man(7)GlcNAc(2)-PP-Dol alpha-1,6-mannosyltransferase.