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## Congenital Disorder of Glycosylation Type Ig

National Cancer Institute

## Source

National Cancer Institute. <u>Congenital Disorder of Glycosylation Type Ig.</u> 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.

Qeios ID: 6CVG7Y · https://doi.org/10.32388/6CVG7Y