Cardiac complications of congenital disorders of glycos of the literature

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Citation Report

#	Article	IF	Citations
1	Protein N-Glycosylation in Cardiovascular Diseases and Related Risk Factors. Current Cardiovascular Risk Reports, 2018, 12, 1.	0.8	19
2	Recognizable phenotypes in CDG. Journal of Inherited Metabolic Disease, 2018, 41, 541-553.	1.7	68
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14	Keeping an eye on congenital disorders of Oâ€glycosylation: A systematic literature review. Journal of Inherited Metabolic Disease, 2019, 42, 29-48.	1.7	22
15	A mutation in mannoseâ€phosphateâ€dolichol utilization defect 1 reveals clinical symptoms of congenital disorders of glycosylation type I and dystroglycanopathy. JIMD Reports, 2019, 50, 31-39.	0.7	14
16	Functional glycomics: Application to medical science and hepatology. Hepatology Research, 2020, 50, 153-164.	1.8	17
17	Vascular ring anomaly in a patient with phosphomannomutase 2 deficiency: A case report and review of the literature. JIMD Reports, 2020, 56, 27-33.	0.7	3
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20	Deciphering the premature mortality in PIGA-CDG â€“ An untold story. Epilepsy Research, 2021, 170, 106530.	0.8	15
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