

Coenzyme Q10 and Pyridoxal Phosphate Deficiency Is a Mucopolysaccharidosis Type III

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#	ARTICLE	IF	CITATIONS
1	Secondary coenzyme Q 10 deficiencies in oxidative phosphorylation (OXPHOS) and non-OXPHOS disorders. <i>Mitochondrion</i> , 2016, 30, 51-58.	1.6	70
2	Molecular diagnosis of coenzyme Q ₁₀ deficiency: an update. <i>Expert Review of Molecular Diagnostics</i> , 2018, 18, 491-498.	1.5	33
3	Plasma coenzyme Q10 status is impaired in selected genetic conditions. <i>Scientific Reports</i> , 2019, 9, 793.	1.6	27
4	Putative adjunct therapies to target mitochondrial dysfunction and oxidative stress in phenylketonuria, lysosomal storage disorders and peroxisomal disorders. <i>Expert Opinion on Orphan Drugs</i> , 2020, 8, 431-444.	0.5	0
5	Oxidative Stress in Mucopolysaccharidoses: Pharmacological Implications. <i>Molecules</i> , 2021, 26, 5616.	1.7	12
6	The Inflammation in the Cytopathology of Patients With Mucopolysaccharidoses- Immunomodulatory Drugs as an Approach to Therapy. <i>Frontiers in Pharmacology</i> , 2022, 13, .	1.6	10
7	Coenzyme Q10: Role in Less Common Age-Related Disorders. <i>Antioxidants</i> , 2022, 11, 2293.	2.2	1
8	Mucopolysaccharidoses: Cellular Consequences of Glycosaminoglycans Accumulation and Potential Targets. <i>International Journal of Molecular Sciences</i> , 2023, 24, 477.	1.8	13