Leigh Syndrome: Spectrum of Molecular Defects and Cl

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Citation Report

#	Article	IF	Citations
1	A boy with a progressive neurologic decline harboring two coexisting mutations in KMT2D and VPS13D. Brain and Development, 2023, , .	1.1	0
2	Mitochondrial Chronic Progressive External Ophthalmoplegia. Brain Sciences, 2024, 14, 135.	2.3	0