Murray Potter

List of Publications by Year in descending order

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1683934 1719901 8 49 5 7 citations h-index g-index papers 8 8 8 83 docs citations times ranked citing authors all docs

#	Article	IF	CITATIONS
1	Outcomes in pediatric studies of medium-chain acyl-coA dehydrogenase (MCAD) deficiency and phenylketonuria (PKU): a review. Orphanet Journal of Rare Diseases, 2020, 15, 12.	1.2	15
2	Health services use among children diagnosed with medium-chain acyl-CoA dehydrogenase deficiency through newborn screening: a cohort study in Ontario, Canada. Orphanet Journal of Rare Diseases, 2019, 14, 70.	1.2	9
3	Severe cystic degeneration and intractable seizures in a newborn with molybdenum cofactor deficiency type B. Molecular Genetics and Metabolism Reports, 2019, 18, 11-13.	0.4	7
4	Health Care for Mitochondrial Disorders in Canada: A Survey of Physicians. Canadian Journal of Neurological Sciences, 2019, 46, 717-726.	0.3	6
5	Implementation of Epilepsy Multigene Panel Testing in Ontario, Canada. Canadian Journal of Neurological Sciences, 2020, 47, 61-68.	0.3	6
6	I-Cell Disease (Mucolipidosis II) Presenting as Neonatal Fractures: A Case for Continued Monitoring of Serum Parathyroid Hormone Levels. Clinical Pediatric Endocrinology, 2008, 17, 81-85.	0.4	5
7	Analysis of serum tranexamic acid in patients undergoing open heart surgery. Clinical Biochemistry, 2021, 87, 74-78.	0.8	1
8	Vomiting and seizure following circumcision in an infant. Paediatrics and Child Health, 2019, 24, 146-147.	0.3	0