

Silvia Tortorelli

List of Publications by Year in descending order

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Version: 2024-02-01

20
papers

1,063
citations

623734

14
h-index

752698

20
g-index

20
all docs

20
docs citations

20
times ranked

1274
citing authors

#	ARTICLE	IF	CITATIONS
1	Clinical validation of cutoff target ranges in newborn screening of metabolic disorders by tandem mass spectrometry: A worldwide collaborative project. <i>Genetics in Medicine</i> , 2011, 13, 230-254.	2.4	308
2	Newborn screening for X-linked adrenoleukodystrophy (X-ALD): Validation of a combined liquid chromatography-tandem mass spectrometric (LC-MS/MS) method. <i>Molecular Genetics and Metabolism</i> , 2009, 97, 212-220.	1.1	154
3	Precision newborn screening for lysosomal disorders. <i>Genetics in Medicine</i> , 2018, 20, 847-854.	2.4	99
4	Combined liquid chromatography-tandem mass spectrometry as an analytical method for high throughput screening for X-linked adrenoleukodystrophy and other peroxisomal disorders: Preliminary findings. <i>Molecular Genetics and Metabolism</i> , 2006, 89, 185-187.	1.1	97
5	Recent developments and new applications of tandem mass spectrometry in newborn screening. <i>Current Opinion in Pediatrics</i> , 2004, 16, 427-433.	2.0	65
6	Making the case for objective performance metrics in newborn screening by tandem mass spectrometry. <i>Mental Retardation and Developmental Disabilities Research Reviews</i> , 2006, 12, 255-261.	3.6	52
7	Acute liver failure in neonates with undiagnosed hereditary fructose intolerance due to exposure from widely available infant formulas. <i>Molecular Genetics and Metabolism</i> , 2018, 123, 428-432.	1.1	40
8	The critical role of psychosine in screening, diagnosis, and monitoring of Krabbe disease. <i>Genetics in Medicine</i> , 2020, 22, 1108-1118.	2.4	39
9	Continuous Age- and Sex-Adjusted Reference Intervals of Urinary Markers for Cerebral Creatine Deficiency Syndromes: A Novel Approach to the Definition of Reference Intervals. <i>Clinical Chemistry</i> , 2015, 61, 760-768.	3.2	38
10	Simultaneous Testing for 6 Lysosomal Storage Disorders and X-Adrenoleukodystrophy in Dried Blood Spots by Tandem Mass Spectrometry. <i>Clinical Chemistry</i> , 2016, 62, 1248-1254.	3.2	37
11	Incorporation of Second-Tier Biomarker Testing Improves the Specificity of Newborn Screening for Mucopolysaccharidosis Type I. <i>International Journal of Neonatal Screening</i> , 2020, 6, 10.	3.2	32
12	Biochemical Diagnosis of Acute Hepatic Porphyrin: Updated Expert Recommendations for Primary Care Physicians. <i>American Journal of the Medical Sciences</i> , 2021, 362, 113-121.	1.1	24
13	A Comparative Effectiveness Study of Newborn Screening Methods for Four Lysosomal Storage Disorders. <i>International Journal of Neonatal Screening</i> , 2020, 6, 44.	3.2	23
14	The Combined Impact of CLIR Post-Analytical Tools and Second Tier Testing on the Performance of Newborn Screening for Disorders of Propionate, Methionine, and Cobalamin Metabolism. <i>International Journal of Neonatal Screening</i> , 2020, 6, 33.	3.2	19
15	Multiplex testing for the screening of lysosomal storage disease in urine: Sulfatides and glycosaminoglycan profiles in 40 cases of sulfatiduria. <i>Molecular Genetics and Metabolism</i> , 2020, 129, 106-110.	1.1	10
16	A new galactose treatment monitoring index for PGM1-CDG. <i>Journal of Inherited Metabolic Disease</i> , 2021, 44, 1263-1271.	3.6	10
17	The Clinical Impact of CLIR Tools toward Rapid Resolution of Post-Newborn Screening Confirmatory Testing for X-Linked Adrenoleukodystrophy in California. <i>International Journal of Neonatal Screening</i> , 2020, 6, 62.	3.2	5
18	Immune dysfunction in MGAT2-CDG: A clinical report and review of the literature. <i>American Journal of Medical Genetics, Part A</i> , 2021, 185, 213-218.	1.2	5

#	ARTICLE	IF	CITATIONS
19	The low excretor phenotype of glutaric acidemia type I is a source of false negative newborn screening results and challenging diagnoses. <i>JIMD Reports</i> , 2021, 60, 67-74.	1.5	4
20	Biochemical phenotype and its relationship to treatment in 16 individuals with PCCB c.1606A>G (p.Asn536Asp) variant propionic acidemia. <i>Molecular Genetics and Metabolism</i> , 2020, 131, 316-324.	1.1	2