## Giancarlo La Marca

List of Publications by Year in descending order

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61984 106344 5,373 136 43 65 citations h-index g-index papers 139 139 139 7108 docs citations times ranked citing authors all docs

#	Article	IF	CITATIONS
1	Clinical validation of cutoff target ranges in newborn screening of metabolic disorders by tandem mass spectrometry: A worldwide collaborative project. Genetics in Medicine, 2011, 13, 230-254.	2.4	308
2	Guidelines for diagnosis and management of the cobalaminâ€related remethylation disorders cblC, cblD, cblE, cblF, cblG, cblJ and MTHFR deficiency. Journal of Inherited Metabolic Disease, 2017, 40, 21-48.	3.6	206
3	The â€~headache tree' via umbellulone and TRPA1 activates the trigeminovascular system. Brain, 2012, 135, 376-390.	7.6	163
4	The pathophysiology of retinopathy of prematurity: an update of previous and recent knowledge. Acta Ophthalmologica, 2014, 92, 2-20.	1.1	146
5	Role of the Adrenergic System in a Mouse Model of Oxygen-Induced Retinopathy: Antiangiogenic Effects of $\hat{I}^2$ -Adrenoreceptor Blockade. , 2011, 52, 155.		141
6	Odorant-Binding Proteins and Chemosensory Proteins in Pheromone Detection and Release in the Silkmoth Bombyx mori. Chemical Senses, $2011$ , $36$ , $335-344$ .	2.0	134
7	Neonatal Screening in Europe Revisited: An ISNS Perspective on the Current State and Developments Since 2010. International Journal of Neonatal Screening, 2021, 7, 15.	3.2	118
8	Enhanced interpretation of newborn screening results without analyte cutoff values. Genetics in Medicine, 2012, 14, 648-655.	2.4	117
9	Proteinuria Impairs Podocyte Regeneration by Sequestering Retinoic Acid. Journal of the American Society of Nephrology: JASN, 2013, 24, 1756-1768.	6.1	116
10	Expanded newborn screening by mass spectrometry: New tests, future perspectives. Mass Spectrometry Reviews, 2016, 35, 71-84.	5.4	107
11	Rapid 2nd-Tier Test for Measurement of 3-OH-Propionic and Methylmalonic Acids on Dried Blood Spots: Reducing the False-Positive Rate for Propionylcarnitine during Expanded Newborn Screening by Liquid Chromatography–Tandem Mass Spectrometry. Clinical Chemistry, 2007, 53, 1364-1369.	3.2	104
12	Oleuropein aglycone protects against pyroglutamylated-3 amyloid-ß toxicity: biochemical, epigenetic and functional correlates. Neurobiology of Aging, 2015, 36, 648-663.	3.1	91
13	Rapid assay of topiramate in dried blood spots by a new liquid chromatography-tandem mass spectrometric method. Journal of Pharmaceutical and Biomedical Analysis, 2008, 48, 1392-1396.	2.8	87
14	Conventional and long-circulating liposomes of artemisinin: preparation, characterization, and pharmacokinetic profile in mice. Journal of Liposome Research, 2011, 21, 237-244.	3.3	87
15	Oral Propranolol for Retinopathy of Prematurity: Risks, Safety Concerns, and Perspectives. Journal of Pediatrics, 2013, 163, 1570-1577.e6.	1.8	80
16	Phenobarbital for neonatal seizures in hypoxic ischemic encephalopathy: A pharmacokinetic study during whole body hypothermia. Epilepsia, 2011, 52, 794-801.	5.1	79
17	Oleuropein, the Main Polyphenol of Olea europaea Leaf Extract, Has an Anti-Cancer Effect on Human BRAF Melanoma Cells and Potentiates the Cytotoxicity of Current Chemotherapies. Nutrients, 2018, 10, 1950.	4.1	79
18	Dominulin A and B: Two new antibacterial peptides identified on the cuticle and in the venom of the social paper wasp Polistes dominulus using MALDI-TOF, MALDI-TOF/TOF, and ESI-ion trap. Journal of the American Society for Mass Spectrometry, 2006, 17, 376-383.	2.8	78

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19	Topiramate concentrations in neonates treated with prolonged whole body hypothermia for hypoxic ischemic encephalopathy. Epilepsia, 2009, 50, 2355-2361.	5.1	75
20	A Chaperone Enhances Blood $\hat{l}_{\pm}$ -Glucosidase Activity in Pompe Disease Patients Treated With Enzyme Replacement Therapy. Molecular Therapy, 2014, 22, 2004-2012.	8.2	75
21	Hematopoietic Stem- and Progenitor-Cell Gene Therapy for Hurler Syndrome. New England Journal of Medicine, 2021, 385, 1929-1940.	27.0	75
22	Serum Levels of Acyl-Carnitines along the Continuum from Normal to Alzheimer's Dementia. PLoS ONE, 2016, 11, e0155694.	2.5	72
23	New Strategy for the Screening of Lysosomal Storage Disorders: The Use of the Online Trapping-and-Cleanup Liquid Chromatography/Mass Spectrometry. Analytical Chemistry, 2009, 81, 6113-6121.	6.5	65
24	Tandem mass spectrometry, but not T-cell receptor excision circle analysis, identifies newborns with late-onset adenosine deaminase deficiency. Journal of Allergy and Clinical Immunology, 2013, 131, 1604-1610.	2.9	65
25	Oral Topiramate in Neonates with Hypoxic Ischemic Encephalopathy Treated with Hypothermia: A Safety Study. Journal of Pediatrics, 2010, 157, 361-366.	1.8	64
26	Are Patients with Potential Celiac Disease Really Potential? The Answer of Metabonomics. Journal of Proteome Research, 2011, 10, 714-721.	3.7	64
27	Neonatal screening for severe combined immunodeficiency caused by an adenosine deaminase defect: AÂreliable and inexpensive method using tandem mass spectrometry. Journal of Allergy and Clinical Immunology, 2011, 127, 1394-1399.	2.9	63
28	<i>N</i> -Carbamylglutamate in Emergency Management of Hyperammonemia in Neonatal Acute Onset Propionic and Methylmalonic Aciduria. Neonatology, 2010, 97, 286-290.	2.0	60
29	A TRPA1 antagonist reverts oxaliplatin-induced neuropathic pain. Scientific Reports, 2013, 3, 2005.	3.3	58
30	The inclusion of succinylacetone as marker for tyrosinemia type I in expanded newborn screening programs. Rapid Communications in Mass Spectrometry, 2008, 22, 812-818.	1.5	57
31	Diagnosis of immunodeficiency caused by a purine nucleoside phosphorylase defect by using tandem mass spectrometry on dried blood spots. Journal of Allergy and Clinical Immunology, 2014, 134, 155-159.e3.	2.9	56
32	Mass spectrometry in clinical chemistry: the case of newborn screening. Journal of Pharmaceutical and Biomedical Analysis, 2014, 101, 174-182.	2.8	55
33	Adenosine kinase deficiency: expanding the clinical spectrum and evaluating therapeutic options. Journal of Inherited Metabolic Disease, 2016, 39, 273-283.	3.6	55
34	New developments in the treatment of hyperammonemia: emerging use of carglumic acid. International Journal of General Medicine, 2011, 4, 21.	1.8	54
35	Lactate Rewires Lipid Metabolism and Sustains a Metabolic–Epigenetic Axis in Prostate Cancer. Cancer Research, 2022, 82, 1267-1282.	0.9	52
36	Rapid quantitation of globotriaosylceramide in human plasma and urine: a potential application for monitoring enzyme replacement therapy in Anderson-Fabry disease. Rapid Communications in Mass Spectrometry, 2002, 16, 1507-1514.	1.5	51

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37	A new rapid micromethod for the assay of phenobarbital from dried blood spots by LCâ€ŧandem mass spectrometry. Epilepsia, 2009, 50, 2658-2662.	5.1	50
38	Study protocol: safety and efficacy of propranolol in newborns with Retinopathy of Prematurity (PROP-ROP): ISRCTN18523491. BMC Pediatrics, 2010, 10, 83.	1.7	50
39	Serum Amino Acid Profiles in Normal Subjects and in Patients with or at Risk of Alzheimer Dementia. Dementia and Geriatric Cognitive Disorders Extra, 2017, 7, 143-159.	1.3	50
40	Safety and efficacy of topiramate in neonates with hypoxic ischemic encephalopathy treated with hypothermia (NeoNATI): a feasibility study. Journal of Maternal-Fetal and Neonatal Medicine, 2018, 31, 973-980.	1.5	50
41	Development of an UPLC–MS/MS method for the determination of antibiotic ertapenem on dried blood spots. Journal of Pharmaceutical and Biomedical Analysis, 2012, 61, 108-113.	2.8	49
42	Reticular dysgenesis–associated AK2 protects hematopoietic stem and progenitor cell development from oxidative stress. Journal of Experimental Medicine, 2015, 212, 1185-1202.	8.5	49
43	î² <sub>3</sub> â€Adrenoceptor as a potential immunoâ€suppressor agent in melanoma. British Journal of Pharmacology, 2019, 176, 2509-2524.	5.4	49
44	Rapid assay of rufinamide in dried blood spots by a new liquid chromatography–tandem mass spectrometric method. Journal of Pharmaceutical and Biomedical Analysis, 2011, 54, 192-197.	2.8	48
45	Delayed-onset adenosine deaminase deficiency: Strategies for an early diagnosis. Journal of Allergy and Clinical Immunology, 2012, 130, 991-994.	2.9	44
46	Hypocitrullinemia in expanded newborn screening by LC–MS/MS is not a reliable marker for ornithine transcarbamylase deficiency. Journal of Pharmaceutical and Biomedical Analysis, 2009, 49, 1292-1295.	2.8	43
47	Eye drop propranolol administration promotes the recovery of oxygen-induced retinopathy in mice. Experimental Eye Research, $2013, 111, 27-35$ .	2.6	43
48	Detection of honeybee venom in envenomed tissues by direct MALDI MSI. Journal of the American Society for Mass Spectrometry, 2009, 20, 112-123.	2.8	42
49	Rapid and sensitive LC–MS/MS method for the analysis of antibiotic linezolid on dried blood spot. Journal of Pharmaceutical and Biomedical Analysis, 2012, 67-68, 86-91.	2.8	41
50	Safety and Tolerability of Antiepileptic Drug Treatment in Children with Epilepsy. Drug Safety, 2012, 35, 519-533.	3.2	41
51	Synergy between the pharmacological chaperone 1â€deoxygalactonojirimycin and the human recombinant alphaâ€galactosidase A in cultured fibroblasts from patients with Fabry disease. Journal of Inherited Metabolic Disease, 2012, 35, 513-520.	3.6	40
52	The inclusion of ADA-SCID in expanded newborn screening by tandem mass spectrometry. Journal of Pharmaceutical and Biomedical Analysis, 2014, 88, 201-206.	2.8	40
53	Ultrasensitive detection of cancer biomarkers by nickel-based isolation of polydisperse extracellular vesicles from blood. EBioMedicine, 2019, 43, 114-126.	6.1	40
54	Therapeutic drug monitoring of carbamazepine and its metabolite in children from dried blood spots using liquid chromatography and tandem mass spectrometry. Journal of Pharmaceutical and Biomedical Analysis, 2015, 109, 164-170.	2.8	39

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55	LC-MS/MS Method for Simultaneous Determination on a Dried Blood Spot of Multiple Analytes Relevant for Treatment Monitoring in Patients with Tyrosinemia Type I. Analytical Chemistry, 2012, 84, 1184-1188.	6.5	37
56	Newborn screening for homocystinurias: Recent recommendations versus current practice. Journal of Inherited Metabolic Disease, 2019, 42, 128-139.	3.6	37
57	Implementing tandem mass spectrometry as a routine tool for characterizing the complete purine and pyrimidine metabolic profile in urine samples. Journal of Mass Spectrometry, 2006, 41, 1442-1452.	1.6	36
58	Solid Olive Residues:Â Insight into Their Phenolic Composition. Journal of Agricultural and Food Chemistry, 2005, 53, 8963-8969.	<b>5.</b> 2	34
59	Propranolol concentrations after oral administration in term and preterm neonates. Journal of Maternal-Fetal and Neonatal Medicine, 2013, 26, 833-840.	1.5	34
60	Pre- and post-dialysis quantitative dosage of thymidine in urine and plasma of a MNGIE patient by using HPLC-ESI-MS/MS. Journal of Mass Spectrometry, 2006, 41, 586-592.	1.6	33
61	Propranolol 0.2% Eye Micro-Drops for Retinopathy of Prematurity: A Prospective Phase IIB Study. Frontiers in Pediatrics, 2019, 7, 180.	1.9	31
62	Development of Strategies to Decrease False Positive Results in Newborn Screening. International Journal of Neonatal Screening, 2020, 6, 84.	3.2	31
63	Highâ€performance liquid chromatography/electrospray ionization tandem mass spectrometric investigation of stilbenoids in cell cultures of <i>Vitis vinifera</i> L., cv. Malvasia. Rapid Communications in Mass Spectrometry, 2010, 24, 2065-2073.	1.5	29
64	Food supplements of Tribulus terrestris L.: An HPLC-ESI-MS method for an estimation of the saponin content. Chromatographia, 2003, 57, 581-592.	1.3	28
65	Falsely elevated C4-carnitine as expression of glutamate formiminotransferase deficiency in tandem mass spectrometry newborn screening. Journal of Mass Spectrometry, 2006, 41, 263-265.	1.6	28
66	Safety and efficacy of topiramate in neonates with hypoxic ischemic encephalopathy treated with hypothermia (NeoNATI). BMC Pediatrics, 2012, 12, 144.	1.7	28
67	Fabry disease: polymorphic haplotypes and a novel missense mutation in the <i>GLA</i> gene. Clinical Genetics, 2012, 81, 224-233.	2.0	28
68	Sudden unexpected fatal encephalopathy in adults with OTC gene mutations-Clues for early diagnosis and timely treatment. Orphanet Journal of Rare Diseases, 2014, 9, 105.	2.7	28
69	Heptadecanoylcarnitine (C17) a novel candidate biomarker for newborn screening of propionic and methylmalonic acidemias. Clinica Chimica Acta, 2015, 450, 342-348.	1.1	27
70	Clinical relevance of short-chain acyl-CoA dehydrogenase (SCAD) deficiency: Exploring the role of new variants including the first SCAD-disease-causing allele carrying a synonymous mutation. BBA Clinical, 2016, 5, 114-119.	4.1	27
71	Children who develop type $1$ diabetes early in life show low levels of carnitine and amino acids at birth: does this finding shed light on the etiopathogenesis of the disease?. Nutrition and Diabetes, 2013, 3, e94-e94.	3.2	26
72	Electrospray ionisation tandem mass spectrometric investigation of phenylpropanoids and secoiridoids from solid olive residue. Rapid Communications in Mass Spectrometry, 2006, 20, 2013-2022.	1.5	25

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73	A rapid liquid chromatography tandem mass spectrometry-based method for measuring propranolol on dried blood spots. Journal of Pharmaceutical and Biomedical Analysis, 2013, 78-79, 34-38.	2.8	25
74	Propranolol 0.1% eye micro-drops in newborns with retinopathy of prematurity: a pilot clinical trial. Pediatric Research, 2017, 81, 307-314.	2.3	24
75	Barth syndrome presenting with acute metabolic decompensation in the neonatal period. Journal of Inherited Metabolic Disease, 2006, 29, 684-684.	3.6	22
76	Delivery of doxorubicin across the blood–brain barrier by ondansetron pretreatment: a study in vitro and in vivo. Cancer Letters, 2014, 353, 242-247.	7.2	22
77	Dried blood spot assay for the quantification of phenytoin using Liquid Chromatography-Mass Spectrometry. Clinica Chimica Acta, 2015, 440, 31-35.	1.1	22
78	Fatal Malonyl CoA Decarboxylase Deficiency Due to Maternal Uniparental Isodisomy of the Telomeric End of Chromosome 16. Annals of Human Genetics, 2007, 71, 705-712.	0.8	21
79	Detection of doxorubicin hydrochloride accumulation in the rat brain after morphine treatment by mass spectrometry. Cancer Chemotherapy and Pharmacology, 2011, 67, 1333-1340.	2.3	20
80	Rapid determination of orotic acid in urine by a fast liquid chromatography/tandem mass spectrometric method. Rapid Communications in Mass Spectrometry, 2003, 17, 788-793.	1.5	19
81	Simultaneous determination of creatine and guanidinoacetate in plasma by liquid chromatography–tandem mass spectrometry (LC–MS/MS). Journal of Pharmaceutical and Biomedical Analysis, 2011, 56, 792-798.	2.8	18
82	Strategies for reducing the incidence of skin complications in newborns treated with whole-body hypothermia. Journal of Maternal-Fetal and Neonatal Medicine, 2012, 25, 2115-2121.	1.5	18
83	Aminoacylase I deficiency due to <scp>ACY1 mRNA</scp> exon skipping. Clinical Genetics, 2014, 86, 367-372.	2.0	18
84	Lipoic-Based TRPA1/TRPV1 Antagonist to Treat Orofacial Pain. ACS Chemical Neuroscience, 2015, 6, 380-385.	3.5	18
85	LC-MS/MS method for simultaneous quantification of heparan sulfate and dermatan sulfate in urine by butanolysis derivatization. Clinica Chimica Acta, 2019, 488, 98-103.	1.1	18
86	Estimating the integrity of aged DNA samples by CE. Electrophoresis, 2009, 30, 3986-3995.	2.4	17
87	Leukocyte and Dried Blood Spot Arylsulfatase A Assay by Tandem Mass Spectrometry. Analytical Chemistry, 2020, 92, 6341-6348.	6.5	17
88	Pharmacological modulation of blood-brain barrier increases permeability of doxorubicin into the rat brain. American Journal of Cancer Research, 2013, 3, 424-32.	1.4	17
89	Successful Propranolol Treatment of a Kaposiform Hemangioendothelioma Apparently Resistant to Propranolol. Pediatric Blood and Cancer, 2016, 63, 1290-1292.	1.5	16
90	The successful inclusion of succinylacetone as a marker of tyrosinemia type I in Tuscany newborn screening program. Rapid Communications in Mass Spectrometry, 2009, 23, 3891-3893.	1.5	15

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91	Measurement of succinyl-carnitine and methylmalonyl-carnitine on dried blood spot by liquid chromatography-tandem mass spectrometry. Clinica Chimica Acta, 2014, 429, 30-33.	1.1	15
92	Vacuolated PAS-positive lymphocytes as an hallmark of Pompe disease and other myopathies related to impaired autophagy. Journal of Cellular Physiology, 2018, 233, 5829-5837.	4.1	15
93	Pharmacokinetics and local safety profile of propranolol eye drops in rabbits. Pediatric Research, 2014, 76, 378-385.	2.3	14
94	Rapid diagnosis of medium chain Acyl Co-A dehydrogenase(MCAD) deficiency in a newborn by liquid chromatography/tandem mass spectrometry. Rapid Communications in Mass Spectrometry, 2003, 17, 2688-2692.	1.5	13
95	Thermal inactivation of SARS COVID-2 virus: Are steam inhalations a potential treatment?. Life Sciences, 2021, 265, 118801.	4.3	12
96	Analysis of Organic Acids and Acylglycines for the Diagnosis of Related Inborn Errors of Metabolism by GC- and HPLC-MS. Methods in Molecular Biology, 2011, 708, 73-98.	0.9	12
97	Synthesis of the essential core of the human glycosylphosphatidylinositol (GPI) anchor. Bioorganic Chemistry, 2011, 39, 88-93.	4.1	11
98	Newborn Screening for Tyrosinemia Type I: Further Evidence that Succinylacetone Determination on Blood Spot Is Essential. JIMD Reports, 2011, 1, 107-109.	1.5	11
99	Reference intervals for orotic acid in urine, plasma and dried blood spot using hydrophilic interaction liquid chromatography–tandem mass spectrometry. Journal of Chromatography B: Analytical Technologies in the Biomedical and Life Sciences, 2012, 883-884, 155-160.	2.3	11
100	Phenotypic and genetic spectrum of ATP6V1A encephalopathy: a disorder of lysosomal homeostasis. Brain, 2022, 145, 2687-2703.	7.6	11
101	High frequency of biotinidase deficiency in Italian population identified by newborn screening. Molecular Genetics and Metabolism Reports, 2020, 25, 100689.	1.1	10
102	Hyperhydroxyprolinaemia: a new case diagnosed during neonatal screening with tandem mass spectrometry. Rapid Communications in Mass Spectrometry, 2005, 19, 863-864.	1.5	9
103	A Pharmacokinetic Study and Correlation with Clinical Response of Rufinamide in Infants with Epileptic Encephalopathies. Pharmacology, 2013, 91, 275-280.	2.2	9
104	Bone status of children born from mothers with autoimmune diseases treated during pregnancy with prednisone and/or low molecular weight heparin. Pediatric Rheumatology, 2014, 12, 47.	2.1	9
105	A successful unrelated peripheral blood stem cell transplantation with reduced intensityâ€conditioning regimen in a patient with lateâ€onset purine nucleoside phosphorylase deficiency. Pediatric Transplantation, 2015, 19, E47-50.	1.0	9
106	Study protocol: safety and efficacy of propranolol 0.2% eye drops in newborns with a precocious stage of retinopathy of prematurity (DROP-ROP-0.2%): a multicenter, open-label, single arm, phase II trial. BMC Pediatrics, 2017, 17, 165.	1.7	9
107	Orange-colored diapers as first sign of Lesch-Nyhan disease in an asymptomatic infant. Pediatric Nephrology, 2010, 25, 2373-2374.	1.7	7
108	Late-Onset N-Acetylglutamate Synthase Deficiency: Report of a Paradigmatic Adult Case Presenting with Headaches and Review of the Literature. International Journal of Molecular Sciences, 2018, 19, 345.	4.1	7

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109	Effect of Beta 3 Adrenoreceptor Modulation on Patency of the Ductus Arteriosus. Cells, 2020, 9, 2625.	4.1	7
110	Interâ€laboratory analytical improvement of succinylacetone and nitisinone quantification from dried blood spot samples. JIMD Reports, 2020, 53, 90-102.	1.5	7
111	Morquio B disease: From pathophysiology towards diagnosis. Molecular Genetics and Metabolism, 2021, 132, 180-188.	1.1	7
112	Multicenter evaluation of use of dried blood spotÂcompared to conventional plasma in measurements of globotriaosylsphingosine (LysoGb3) concentration in 104 Fabry patients. Clinical Chemistry and Laboratory Medicine, 2021, 59, 1516-1526.	2.3	7
113	The successful inclusion of ADA SCID in Tuscany expanded newborn screening program. Clinical Chemistry and Laboratory Medicine, 2021, 59, e401-e404.	2.3	7
114	The diagnostic challenge of mild citrulline elevation at newborn screening. Molecular Genetics and Metabolism, 2022, 135, 327-332.	1.1	7
115	First prenatal molecular diagnosis in a family with holocarboxylase synthetase deficiency. Prenatal Diagnosis, 2005, 25, 1117-1119.	2.3	6
116	Hypothermia for neonatal hypoxic-ischemic encephalopathy: may an early amplitude-integrated EEG improve the selection of candidates for cooling?. Journal of Maternal-Fetal and Neonatal Medicine, 2012, 25, 2171-2176.	1.5	6
117	PRDX1 gene-related epi-cblC disease is a common type of inborn error of cobalamin metabolism with mono- or bi-allelic MMACHC epimutations. Clinical Epigenetics, 2021, 13, 137.	4.1	6
118	Screening of Lysosomal Storage Disorders: Application of the Online Trapping-and-Cleanup Liquid Chromatography/Mass Spectrometry Method for Mucopolysaccharidosis I. European Journal of Mass Spectrometry, 2013, 19, 497-503.	1.0	5
119	Medium-Chain Acyl-CoA Deficiency: Outlines from Newborn Screening, <i>In Silico </i> Predictions, and Molecular Studies. Scientific World Journal, The, 2013, 2013, 1-8.	2.1	5
120	Upfront Enzyme Replacement via Erythrocyte Transfusions for PNP Deficiency. Journal of Clinical Immunology, 2021, 41, 1112-1115.	3.8	5
121	Liver-Directed Adeno-Associated Virus–Mediated Gene Therapy for Mucopolysaccharidosis Type VI. , 2022, 1, .		5
122	Biotinidase deficiency due to a de novo mutation or gonadal mosaicism in a first child. Clinica Chimica Acta, 2015, 445, 70-72.	1.1	4
123	Reducing the False-Positive Rate for Isovalerylcarnitine in Expanded Newborn Screening. FIRE Forum for International Research in Education, 2016, 4, 232640981666135.	0.7	4
124	Development and validation of a 2nd tier test for identification of purine nucleoside phosphorylase deficiency patients during expanded newborn screening by liquid chromatography-tandem mass spectrometry. Clinical Chemistry and Laboratory Medicine, 2016, 54, 627-32.	2.3	4
125	Development of a fast LC-MS/MS protocol for combined measurement of six LSDs on dried blood spot in a newborn screening program. Journal of Pharmaceutical and Biomedical Analysis, 2019, 165, 135-140.	2.8	4
126	Study protocol: treatment with caffeine of the very preterm infant in the delivery room: a feasibility study. BMJ Open, 2020, 10, e040105.	1.9	4

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127	A new strategy implementing mass spectrometry in the diagnosis of congenital disorders of N-glycosylation (CDG). Clinical Chemistry and Laboratory Medicine, 2021, 59, 165-171.	2.3	4
128	Incessant Automatic Atrial Tachycardia in a Neonate Successfully Treated with Nadolol and Closely Spaced Doses of Flecainide: A Case Report. Pediatric Reports, 2020, 12, 108-113.	1.3	3
129	Serum creatinine during physiological perinatal dehydration may estimate individual nephron endowment. European Journal of Pediatrics, 2018, 177, 1383-1388.	2.7	2
130	Children with special health care needs attending emergency department in Italy: analysis of 3479 cases. Italian Journal of Pediatrics, 2020, 46, 173.	2.6	2
131	Lysosomals. , 2014, , 785-793.		2
132	Early Diagnosis and Treatment of Purine Nucleoside Phosphorylase (PNP) Deficiency through TREC-Based Newborn Screening. International Journal of Neonatal Screening, 2021, 7, 62.	3.2	2
133	Metabolite and thymocyte development defects in ADA-SCID mice receiving enzyme replacement therapy. Scientific Reports, 2021, 11, 23221.	3.3	2
134	Newborn Screening: Are We Ready for It?. Journal of Neuromuscular Diseases, 2015, 2, S10-S10.	2.6	0
135	Reticular dysgenesis–associated AK2 protects hematopoietic stem and progenitor cell development from oxidative stress. Journal of Cell Biology, 2015, 210, 21020IA141.	5.2	O
136	Newborn Screening: Are We Ready for It?. Journal of Neuromuscular Diseases, 2015, 2, S10.	2.6	0