

Beth K Potter

List of Publications by Year in Descending Order

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

82
papers

1,246
citations

19
h-index

32
g-index

86
ext. papers

1,571
ext. citations

3.9
avg, IF

4.12
L-index

#	Paper	IF	Citations
82	Families' healthcare experiences for children with inherited metabolic diseases: protocol for a mixed methods cohort study.. <i>BMJ Open</i> , 2022 , 12, e055664	3	0
81	Blood metals and vitamin D status in a pregnancy cohort: A bidirectional biomarker analysis.. <i>Environmental Research</i> , 2022 , 211, 113034	7.9	0
80	Patient Engagement in a Multi-Stakeholder Workshop to Plan the Collection of Patient-Oriented Outcomes for Children with Inherited Metabolic Diseases.. <i>Healthcare Quarterly (Toronto, Ont)</i> , 2022 , 24, 81-85	0.5	0
79	Establishing a core outcome set for mucopolysaccharidoses (MPS) in children: study protocol for a rapid literature review, candidate outcomes survey, and Delphi surveys. <i>Trials</i> , 2021 , 22, 816	2.8	0
78	A Retrospective Cohort Study Investigating the Impact of Maternal Pre-Pregnancy Body Mass Index on Pediatric Health Service Utilization. <i>Journal of Obstetrics and Gynaecology Canada</i> , 2021 , 43, 1267-1273 ³	1.3	0
77	Family Experiences with Care for Children with Inherited Metabolic Diseases in Canada: A Cross-Sectional Survey. <i>Patient</i> , 2021 , 1	3.7	0
76	Core Outcome Sets for Medium-Chain Acyl-CoA Dehydrogenase Deficiency and Phenylketonuria. <i>Pediatrics</i> , 2021 , 148,	7.4	2
75	Association between newborn screening analyte profiles and infant mortality. <i>Journal of Maternal-Fetal and Neonatal Medicine</i> , 2021 , 34, 835-838	2	0
74	Developments in evidence creation for treatments of inborn errors of metabolism. <i>Journal of Inherited Metabolic Disease</i> , 2021 , 44, 88-98	5.4	6
73	Stakeholder perspectives on clinical research related to therapies for rare diseases: therapeutic misconception and the value of research. <i>Orphanet Journal of Rare Diseases</i> , 2021 , 16, 26	4.2	3
72	Screening for depression in children and adolescents: a protocol for a systematic review update. <i>Systematic Reviews</i> , 2021 , 10, 24	3	2
71	Health services use by children identified as heterozygous hemoglobinopathy mutation carriers via newborn screening. <i>BMC Pediatrics</i> , 2021 , 21, 296	2.6	0
70	Patient and family engagement in the development of core outcome sets for two rare chronic diseases in children. <i>Research Involvement and Engagement</i> , 2021 , 7, 66	4.4	0
69	Parental psychosocial aspects and stressors involved in the management of inborn errors of metabolism. <i>Molecular Genetics and Metabolism Reports</i> , 2020 , 25, 100654	1.8	3
68	Economic Evaluation of Cannabinoid Oil for Dravet Syndrome: A Cost-Utility Analysis. <i>Pharmacoeconomics</i> , 2020 , 38, 971-980	4.4	4
67	Neurologists' perspectives on medical cannabis for pediatric drug-resistant epilepsy in Canada: A qualitative interview study. <i>Seizure: the Journal of the British Epilepsy Association</i> , 2020 , 78, 118-126	3.2	4
66	Outcomes in pediatric studies of medium-chain acyl-coA dehydrogenase (MCAD) deficiency and phenylketonuria (PKU): a review. <i>Orphanet Journal of Rare Diseases</i> , 2020 , 15, 12	4.2	7

65	Evaluation of the quality of clinical data collection for a pan-Canadian cohort of children affected by inherited metabolic diseases: lessons learned from the Canadian Inherited Metabolic Diseases Research Network. <i>Orphanet Journal of Rare Diseases</i> , 2020 , 15, 89	4.2	4
64	Cannabis-based products for pediatric epilepsy: An updated systematic review. <i>Seizure: the Journal of the British Epilepsy Association</i> , 2020 , 75, 18-22	3.2	16
63	Barriers in accessing medical cannabis for children with drug-resistant epilepsy in Canada: A qualitative study. <i>Epilepsy and Behavior</i> , 2020 , 111, 107120	3.2	5
62	External validation of ELASTIC NET regression models including newborn metabolomic markers for postnatal gestational age estimation in East and South-East Asian infants. <i>Gates Open Research</i> , 2020 , 4, 164	2.4	1
61	Health-care providers' perspectives on uncertainty generated by variant forms of newborn screening targets. <i>Genetics in Medicine</i> , 2020 , 22, 566-573	8.1	4
60	What is in a Name? Parent, Professional and Policy-Maker Conceptions of Consent-Related Language in the Context of Newborn Screening. <i>Public Health Ethics</i> , 2019 , 12, 158-175	1.8	
59	Screening for depression in women during pregnancy or the first year postpartum and in the general adult population: a protocol for two systematic reviews to update a guideline of the Canadian Task Force on Preventive Health Care. <i>Systematic Reviews</i> , 2019 , 8, 27	3	15
58	Decision Models for Assessing the Cost Effectiveness of Treatments for Pediatric Drug-Resistant Epilepsy: A Systematic Review of Economic Evaluations. <i>Pharmacoeconomics</i> , 2019 , 37, 1261-1276	4.4	8
57	Health services use among children diagnosed with medium-chain acyl-CoA dehydrogenase deficiency through newborn screening: a cohort study in Ontario, Canada. <i>Orphanet Journal of Rare Diseases</i> , 2019 , 14, 70	4.2	3
56	Cost-effectiveness of cannabinoids for pediatric drug-resistant epilepsy: protocol for a systematic review of economic evaluations. <i>Systematic Reviews</i> , 2019 , 8, 75	3	6
55	A quality assessment of Health Management Information System (HMIS) data for maternal and child health in Jimma Zone, Ethiopia. <i>PLoS ONE</i> , 2019 , 14, e0213600	3.7	19
54	Incidental screen positive findings in a prospective cohort study in Matlab, Bangladesh: insights into expanded newborn screening for low-resource settings. <i>Orphanet Journal of Rare Diseases</i> , 2019 , 14, 25	4.2	6
53	Health Care for Mitochondrial Disorders in Canada: A Survey of Physicians. <i>Canadian Journal of Neurological Sciences</i> , 2019 , 46, 717-726	1	4
52	A systematic review of the association between coping strategies and quality of life among caregivers of children with chronic illness and/or disability. <i>BMC Pediatrics</i> , 2019 , 19, 215	2.6	17
51	Association between newborn screening analytes and hypoxic ischemic encephalopathy. <i>Scientific Reports</i> , 2019 , 9, 15704	4.9	4
50	External validation of postnatal gestational age estimation using newborn metabolic profiles in Matlab, Bangladesh. <i>ELife</i> , 2019 , 8,	8.9	10
49	Family History Taking in Pediatric Practice: A Qualitative Interview Study. <i>Public Health Genomics</i> , 2019 , 22, 110-118	1.9	2
48	Utilization of key preventive measures for pregnancy complications and malaria among women in Jimma Zone, Ethiopia. <i>BMC Public Health</i> , 2019 , 19, 1443	4.1	4

47	Cannabis-based products for pediatric epilepsy: A systematic review. <i>Epilepsia</i> , 2019 , 60, 6-19	6.4	39
46	Mental Health Screening and Differences in Access to Care among Prisoners. <i>Canadian Journal of Psychiatry</i> , 2018 , 63, 692-700	4.8	6
45	Decision curve analysis as a framework to estimate the potential value of screening or other decision-making aids. <i>International Journal of Methods in Psychiatric Research</i> , 2018 , 27,	4.3	2
44	Mental Health Screening, Treatment, and Institutional Incidents: A Propensity Score Matched Analysis of Long-Term Outcomes of Screening. <i>International Journal of Forensic Mental Health</i> , 2018 , 17, 133-144	1	
43	Association Between Newborn Metabolic Profiles and Pediatric Kidney Disease. <i>Kidney International Reports</i> , 2018 , 3, 691-700	4.1	6
42	T-cell receptor excision circle levels and safety of paediatric immunization: A population-based self-controlled case series analysis. <i>Human Vaccines and Immunotherapeutics</i> , 2018 , 14, 1378-1391	4.4	1
41	Using a meta-narrative literature review and focus groups with key stakeholders to identify perceived challenges and solutions for generating robust evidence on the effectiveness of treatments for rare diseases. <i>Orphanet Journal of Rare Diseases</i> , 2018 , 13, 104	4.2	12
40	Attitudes of undergraduate university women towards HPV vaccination: a cross-sectional study in Ottawa, Canada. <i>BMC Women's Health</i> , 2018 , 18, 134	2.9	10
39	Mental health treatment patterns following screening at intake to prison. <i>Journal of Consulting and Clinical Psychology</i> , 2018 , 86, 15-23	6.5	7
38	Rural and urban disparities in the care of Canadian patients with inflammatory bowel disease: a population-based study. <i>Clinical Epidemiology</i> , 2018 , 10, 1613-1626	5.9	19
37	Migraine and Mental Health in a Population-Based Sample of Adolescents. <i>Canadian Journal of Neurological Sciences</i> , 2017 , 44, 44-50	1	16
36	Psychosocial Response to Uncertain Newborn Screening Results for Cystic Fibrosis. <i>Journal of Pediatrics</i> , 2017 , 184, 165-171.e1	3.6	19
35	Postnatal Prediction of Gestational Age Using Newborn Fetal Hemoglobin Levels. <i>EBioMedicine</i> , 2017 , 15, 203-209	8.8	19
34	Performance of a postnatal metabolic gestational age algorithm: a retrospective validation study among ethnic subgroups in Canada. <i>BMJ Open</i> , 2017 , 7, e015615	3	9
33	False-Positive Newborn Screening for Cystic Fibrosis and Health Care Use. <i>Pediatrics</i> , 2017 , 140,	7.4	20
32	Establishing core outcome sets for phenylketonuria (PKU) and medium-chain Acyl-CoA dehydrogenase (MCAD) deficiency in children: study protocol for systematic reviews and Delphi surveys. <i>Trials</i> , 2017 , 18, 603	2.8	6
31	A secondary benefit: the reproductive impact of carrier results from newborn screening for cystic fibrosis. <i>Genetics in Medicine</i> , 2017 , 19, 403-411	8.1	7
30	Using newborn screening analytes to identify cases of neonatal sepsis. <i>Scientific Reports</i> , 2017 , 7, 18020	4.9	15

29	Rural and Urban Residence During Early Life is Associated with Risk of Inflammatory Bowel Disease: A Population-Based Inception and Birth Cohort Study. <i>American Journal of Gastroenterology</i> , 2017 , 112, 1412-1422	0.7	57
28	Child and family experiences with inborn errors of metabolism: a qualitative interview study with representatives of patient groups. <i>Journal of Inherited Metabolic Disease</i> , 2016 , 39, 139-47	5.4	19
27	Translating rare-disease therapies into improved care for patients and families: what are the right outcomes, designs, and engagement approaches in health-systems research?. <i>Genetics in Medicine</i> , 2016 , 18, 117-23	8.1	30
26	The use of relative incidence ratios in self-controlled case series studies: an overview. <i>BMC Medical Research Methodology</i> , 2016 , 16, 126	4.7	14
25	Experiences of caregivers of children with inherited metabolic diseases: a qualitative study. <i>Orphanet Journal of Rare Diseases</i> , 2016 , 11, 168	4.2	20
24	The health system impact of false positive newborn screening results for medium-chain acyl-CoA dehydrogenase deficiency: a cohort study. <i>Orphanet Journal of Rare Diseases</i> , 2016 , 11, 12	4.2	24
23	Accurate prediction of gestational age using newborn screening analyte data. <i>American Journal of Obstetrics and Gynecology</i> , 2016 , 214, 513.e1-513.e9	6.4	25
22	Consent for newborn screening: parents and health-care professionals' experiences of consent in practice. <i>European Journal of Human Genetics</i> , 2016 , 24, 1530-1534	5.3	16
21	Yield and Efficiency of Mental Health Screening: A Comparison of Screening Protocols at Intake to Prison. <i>PLoS ONE</i> , 2016 , 11, e0154106	3.7	8
20	Attitudes to incorporating genomic risk assessments into population screening programs: the importance of purpose, context and deliberation. <i>BMC Medical Genomics</i> , 2016 , 9, 25	3.7	7
19	Education and parental involvement in decision-making about newborn screening: understanding goals to clarify content. <i>Journal of Genetic Counseling</i> , 2015 , 24, 400-8	2.5	8
18	Scoping review of patient- and family-oriented outcomes and measures for chronic pediatric disease. <i>BMC Pediatrics</i> , 2015 , 15, 7	2.6	16
17	Metabolic Clinic Atlas: Organization of Care for Children with Inherited Metabolic Disease in Canada. <i>JIMD Reports</i> , 2015 , 21, 15-22	1.9	3
16	Benefits and burdens of newborn screening: public understanding and decision-making. <i>Personalized Medicine</i> , 2014 , 11, 593-607	2.2	8
15	Metabolomics of prematurity: analysis of patterns of amino acids, enzymes, and endocrine markers by categories of gestational age. <i>Pediatric Research</i> , 2014 , 75, 367-73	3.2	33
14	Seasonal variation in rates of emergency room visits and acute admissions following recommended infant vaccinations in Ontario, Canada: a self-controlled case series analysis. <i>Vaccine</i> , 2014 , 32, 7148-53	4.1	2
13	Achieving the "triple aim" for inborn errors of metabolism: a review of challenges to outcomes research and presentation of a new practice-based evidence framework. <i>Genetics in Medicine</i> , 2013 , 15, 415-22	8.1	24
12	Variability in the clinical management of fatty acid oxidation disorders: results of a survey of Canadian metabolic physicians. <i>Journal of Inherited Metabolic Disease</i> , 2012 , 35, 115-23	5.4	19

11	Factors associated with knowledge of and satisfaction with newborn screening education: a survey of mothers. <i>Genetics in Medicine</i> , 2012 , 14, 963-70	8.1	22
10	Newborn screening education on the internet: a content analysis of North American newborn screening program websites. <i>Journal of Community Genetics</i> , 2011 , 2, 127-34	2.5	16
9	Reporting guidelines for survey research: an analysis of published guidance and reporting practices. <i>PLoS Medicine</i> , 2010 , 8, e1001069	11.6	212
8	The first three years of screening for medium chain acyl-CoA dehydrogenase deficiency (MCADD) by newborn screening ontario. <i>BMC Pediatrics</i> , 2010 , 10, 82	2.6	18
7	Exploring informed choice in the context of prenatal testing: findings from a qualitative study. <i>Health Expectations</i> , 2008 , 11, 355-65	3.7	40
6	Guidance for considering ethical, legal, and social issues in health technology assessment: application to genetic screening. <i>International Journal of Technology Assessment in Health Care</i> , 2008 , 24, 412-22	1.8	22
5	Newborn blood spot screening in four countries: stakeholder involvement. <i>Journal of Public Health Policy</i> , 2008 , 29, 121-42	2.9	14
4	Socioeconomic status and non-fatal injuries among Canadian adolescents: variations across SES and injury measures. <i>BMC Public Health</i> , 2005 , 5, 132	4.1	34
3	Is there value in using physician billing claims along with other administrative health care data to document the burden of adolescent injury? An exploratory investigation with comparison to self-reports in Ontario, Canada. <i>BMC Health Services Research</i> , 2005 , 5, 15	2.9	9
2	A comparison of measures of socioeconomic status for adolescents in a Canadian national health survey. <i>Chronic Diseases in Canada</i> , 2005 , 26, 80-9		10
1	Does a relationship exist between body weight, concerns about weight, and smoking among adolescents? An integration of the literature with an emphasis on gender. <i>Nicotine and Tobacco Research</i> , 2004 , 6, 397-425	4.9	144