

# Anne-Marie Laberge

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/7761938/publications.pdf>

Version: 2024-02-01

38  
papers

743  
citations

623574

14  
h-index

552653

26  
g-index

38  
all docs

38  
docs citations

38  
times ranked

1450  
citing authors

#	ARTICLE	IF	CITATIONS
1	Family Experiences with Care for Children with Inherited Metabolic Diseases in Canada: A Cross-Sectional Survey. <i>Patient</i> , 2022, 15, 171-185.	1.1	1
2	Metabolically healthy obesity in children enrolled in the <sc>CANadian</sc> Pediatric Weight management Registry (<sc>CANPWR</sc>): An exploratory secondary analysis of baseline data. <i>Clinical Obesity</i> , 2022, 12, e12490.	1.1	9
3	The Serious Factor in Expanded Prenatal Genetic Testing. <i>American Journal of Bioethics</i> , 2022, 22, 23-25.	0.5	7
4	Expanded Prenatal Testing: Maintaining a Non-Directive Approach to Promote Reproductive Autonomy. <i>American Journal of Bioethics</i> , 2022, 22, 39-42.	0.5	3
5	Noninvasive Prenatal Testing: Views of Canadian Pregnant Women and Their Partners Regarding Pressure and Societal Concerns. <i>AJOB Empirical Bioethics</i> , 2021, 12, 53-62.	0.8	10
6	A qualitative study of women and partners from Lebanon and Quebec regarding an expanded scope of noninvasive prenatal testing. <i>BMC Pregnancy and Childbirth</i> , 2021, 21, 54.	0.9	5
7	When to test fetuses for RASopathies? Proposition from a systematic analysis of 352 multicenter cases and a postnatal cohort. <i>Genetics in Medicine</i> , 2021, 23, 1116-1124.	1.1	17
8	Genetic burden linked to founder effects in Saguenayâ€“Lac-Saint-Jean illustrates the importance of genetic screening test availability. <i>Journal of Medical Genetics</i> , 2021, 58, 653-665.	1.5	12
9	Variability in How Canadian Pediatric Weight Management Clinics Deliver Care: Evidence from the CANadian Pediatric Weight Management Registry. <i>Childhood Obesity</i> , 2021, 17, 420-426.	0.8	3
10	Individual and family characteristics associated with health indicators at entry into multidisciplinary pediatric weight management: findings from the CANadian Pediatric Weight management Registry (CANPWR). <i>International Journal of Obesity</i> , 2021, , .	1.6	2
11	Retrospective analysis of fetal vertebral defects: Associated anomalies, etiologies, and outcome. <i>American Journal of Medical Genetics, Part A</i> , 2020, 182, 664-672.	0.7	11
12	A Novel Recurrent <i>COL5A1</i> Genetic Variant Is Associated With a Dysplasia-Associated Arterial Disease Exhibiting Dissections and Fibromuscular Dysplasia. <i>Arteriosclerosis, Thrombosis, and Vascular Biology</i> , 2020, 40, 2686-2699.	1.1	30
13	Implementation challenges for an ethical introduction of noninvasive prenatal testing: a qualitative study of healthcare professionalsâ€™ views from Lebanon and Quebec. <i>BMC Medical Ethics</i> , 2020, 21, 15.	1.0	11
14	Secondary findings from next-generation sequencing: what does actionable in childhood really mean?. <i>Genetics in Medicine</i> , 2019, 21, 124-132.	1.1	18
15	Toward Broader Genetic Contextualism: Genetic Testing Enters the Age of Evidence-Based Medicine. <i>American Journal of Bioethics</i> , 2019, 19, 77-79.	0.5	4
16	Improving recommendations for genomic medicine: building an evolutionary process from clinical practice advisory documents to guidelines. <i>Genetics in Medicine</i> , 2019, 21, 2431-2438.	1.1	13
17	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.	1.2	9
18	Canadian Pregnant Women's Preferences Regarding NIPT for Down Syndrome: The Information They Want, How They Want to Get It, and With Whom They Want to Discuss It. <i>Journal of Obstetrics and Gynaecology Canada</i> , 2019, 41, 782-791.	0.3	25

#	ARTICLE	IF	CITATIONS
19	Homozygous/compound heterozygote <i>RYR1</i> gene variants: Expanding the clinical spectrum. <i>American Journal of Medical Genetics, Part A</i> , 2019, 179, 386-396.	0.7	19
20	The value of non-invasive prenatal testing: preferences of Canadian pregnant women, their partners, and health professionals regarding NIPT use and access. <i>BMC Pregnancy and Childbirth</i> , 2019, 19, 22.	0.9	23
21	Pre-implantation Genetic Diagnosis: The Road Forward in Canada. <i>Journal of Obstetrics and Gynaecology Canada</i> , 2019, 41, 68-71.	0.3	7
22	Diagnostic and Therapeutic Misconception: Parental Expectations and Perspectives Regarding Genetic Testing for Developmental Disorders. <i>Journal of Autism and Developmental Disorders</i> , 2019, 49, 363-375.	1.7	11
23	Recommending inclusion of HFE C282Y homozygotes in the ACMG actionable gene list: cop-out or stealth move toward population screening?. <i>Genetics in Medicine</i> , 2018, 20, 400-402.	1.1	3
24	Obstetric and cardiac outcomes in women with Marfan syndrome and an aortic root diameter $\geq$ 45mm. <i>European Journal of Obstetrics, Gynecology and Reproductive Biology</i> , 2018, 230, 68-72.	0.5	14
25	BCL11B mutations in patients affected by a neurodevelopmental disorder with reduced type 2 innate lymphoid cells. <i>Brain</i> , 2018, 141, 2299-2311.	3.7	81
26	Cross-cultural perspectives on decision making regarding noninvasive prenatal testing: A comparative study of Lebanon and Quebec. <i>AJOB Empirical Bioethics</i> , 2018, 9, 99-111.	0.8	19
27	Paediatricians underuse recommended genetic tests in children with global developmental delay. <i>Paediatrics and Child Health</i> , 2018, 23, e156-e162.	0.3	5
28	Experience of carrier couples identified through a population-based carrier screening pilot program for four founder autosomal recessive diseases in Saguenay-Lac-Saint-Jean. <i>Prenatal Diagnosis</i> , 2018, 38, 67-74.	1.1	12
29	Experiences of caregivers of children with inherited metabolic diseases: a qualitative study. <i>Orphanet Journal of Rare Diseases</i> , 2016, 11, 168.	1.2	38
30	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.	1.2	38
31	Genetic Testing in Thoracic Aortic Disease—When, Why, and How?. <i>Canadian Journal of Cardiology</i> , 2016, 32, 131-134.	0.8	12
32	Screening Children for Familial Aortopathies: Tread With Caution. <i>Canadian Journal of Cardiology</i> , 2016, 32, 60-65.	0.8	4
33	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, 139-147.	1.7	26
34	MG-114...First 2 years of experience of an integrated multidisciplinary clinic for adults with aortopathies in a Canadian context. <i>Journal of Medical Genetics</i> , 2015, 52, A3.1-A3.	1.5	0
35	Scoping review of patient- and family-oriented outcomes and measures for chronic pediatric disease. <i>BMC Pediatrics</i> , 2015, 15, 7.	0.7	20
36	The clinical application of genome-wide sequencing for monogenic diseases in Canada: Position Statement of the Canadian College of Medical Geneticists. <i>Journal of Medical Genetics</i> , 2015, 52, 431-437.	1.5	187

#	ARTICLE	IF	CITATIONS
37	Use of Factor V Leiden genetic testing in practice and impact on management. <i>Genetics in Medicine</i> , 2009, 11, 750-756.	1.1	16
38	Long-term outcomes of the "Genetics in Primary Care" faculty development initiative. <i>Family Medicine</i> , 2009, 41, 266-70.	0.3	18