Anne-Marie Laberge

List of Publications by Year
in descending order

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	The clinical application of genome-wide sequencing for monogenic diseases in Canada: Position Statement of the Canadian College of Medical Geneticists. Journal of Medical Genetics, 2015, 52, 431-437.	1.5	187
2	BCL11B mutations in patients affected by a neurodevelopmental disorder with reduced type 2 innate lymphoid cells. Brain, 2018, 141, 2299-2311.	3.7	81
3	Experiences of caregivers of children with inherited metabolic diseases: a qualitative study. Orphanet Journal of Rare Diseases, 2016, 11, 168.	1.2	38
4	The health system impact of false positive newborn screening results for medium-chain acyl-CoA dehydrogenase deficiency: a cohort study. Orphanet Journal of Rare Diseases, 2016, 11, 12.	1.2	38
5	A Novel Recurrent COL5A1 Genetic Variant Is Associated With a Dysplasia-Associated Arterial Disease Exhibiting Dissections and Fibromuscular Dysplasia. Arteriosclerosis, Thrombosis, and Vascular Biology, 2020, 40, 2686-2699.	1.1	30
6	Child and family experiences with inborn errors of metabolism: a qualitative interview study with representatives of patient groups. Journal of Inherited Metabolic Disease, 2016, 39, 139-147.	1.7	26
7	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. Journal of Obstetrics and Gynaecology Canada, 2019, 41, 782-791.	0.3	25
8	The value of non-invasive prenatal testing: preferences of Canadian pregnant women, their partners, and health professionals regarding NIPT use and access. BMC Pregnancy and Childbirth, 2019, 19, 22.	0.9	23
9	Scoping review of patient- and family-oriented outcomes and measures for chronic pediatric disease. BMC Pediatrics, 2015, 15, 7.	0.7	20
10	Cross-cultural perspectives on decision making regarding noninvasive prenatal testing: A comparative study of Lebanon and Quebec. AJOB Empirical Bioethics, 2018, 9, 99-111.	0.8	19
11	Homozygous/compound heterozygote RYR1 gene variants: Expanding the clinical spectrum. American Journal of Medical Genetics, Part A, 2019, 179, 386-396.	0.7	19
12	SecondaryÂfindings from next-generation sequencing: what does actionable in childhood really mean?. Genetics in Medicine, 2019, 21, 124-132.	1.1	18
13	Long-term outcomes of the "Genetics in Primary Care" faculty development initiative. Family Medicine, 2009, 41, 266-70.	0.3	18
14	When to test fetuses for RASopathies? Proposition from a systematic analysis of 352 multicenter cases and a postnatal cohort. Genetics in Medicine, 2021, 23, 1116-1124.	1.1	17
15	Use of Factor V Leiden genetic testing in practice and impact on management. Genetics in Medicine, 2009, 11, 750-756.	1.1	16
16	Obstetric and cardiac outcomes in women with Marfan syndrome and an aortic root diameter â‰¥ 45mm. European Journal of Obstetrics, Gynecology and Reproductive Biology, 2018, 230, 68-72.	0.5	14
17	Improving recommendations for genomic medicine: building an evolutionary process from clinical practice advisory documents to guidelines. Genetics in Medicine, 2019, 21, 2431-2438.	1.1	13
18	Genetic Testing in Thoracic Aortic Diseaseâ€"When, Why, and How?. Canadian Journal of Cardiology, 2016, 32, 131-134.	0.8	12

#	Article	IF	Citations
19	Genetic burden linked to founder effects in Saguenayâ€“Lac-Saint-Jean illustrates the importance of genetic screening test availability. Journal of Medical Genetics, 2021, 58, 653-665.	1.5	12
20	Experience of carrier couples identified through a populationâ€based carrier screening pilot program for four founder autosomal recessive diseases in Saguenayâ€“Lacâ€Saintâ€Jean. Prenatal Diagnosis, 2018, 38, 67-74.	1.1	12
21	Diagnostic and Therapeutic Misconception: Parental Expectations and Perspectives Regarding Genetic Testing for Developmental Disorders. Journal of Autism and Developmental Disorders, 2019, 49, 363-375.	1.7	11
22	Retrospective analysis of fetal vertebral defects: Associated anomalies, etiologies, and outcome. American Journal of Medical Genetics, Part A, 2020, 182, 664-672.	0.7	11
23	Implementation challenges for an ethical introduction of noninvasive prenatal testing: a qualitative study of healthcare professionalsâ€™ views from Lebanon and Quebec. BMC Medical Ethics, 2020, 21, 15.	1.0	11
24	Noninvasive Prenatal Testing: Views of Canadian Pregnant Women and Their Partners Regarding Pressure and Societal Concerns. AJOB Empirical Bioethics, 2021, 12, 53-62.	0.8	10
25	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
26	Metabolically healthy obesity in children enrolled in the <scp>CANadian</scp> Pediatric Weight management Registry (<scp>CANPWR</scp>): An exploratory secondary analysis of baseline data. Clinical Obesity, 2022, 12, e12490.	1.1	9
27	Pre-implantation Genetic Diagnosis: The Road Forward in Canada. Journal of Obstetrics and Gynaecology Canada, 2019, 41, 68-71.	0.3	7
28	The Serious Factor in Expanded Prenatal Genetic Testing. American Journal of Bioethics, 2022, 22, 23-25.	0.5	7
29	Paediatricians underuse recommended genetic tests in children with global developmental delay. Paediatrics and Child Health, 2018, 23, e156-e162.	0.3	5
30	A qualitative study of women and partners from Lebanon and Quebec regarding an expanded scope of noninvasive prenatal testing. BMC Pregnancy and Childbirth, 2021, 21, 54.	0.9	5
31	Screening Children for Familial Aortopathies: Tread With Caution. Canadian Journal of Cardiology, 2016, 32, 60-65.	0.8	4
32	Toward Broader Genetic Contextualism: Genetic Testing Enters the Age of Evidence-Based Medicine. American Journal of Bioethics, 2019, 19, 77-79.	0.5	4
33	Recommending inclusion of HFE C282Y homozygotes in the ACMG actionable gene list: cop-out or stealth move toward population screening?. Genetics in Medicine, 2018, 20, 400-402.	1.1	3
34	Variability in How Canadian Pediatric Weight Management Clinics Deliver Care: Evidence from the CANadian Pediatric Weight Management Registry. Childhood Obesity, 2021, 17, 420-426.	0.8	3
35	Expanded Prenatal Testing: Maintaining a Non-Directive Approach to Promote Reproductive Autonomy. American Journal of Bioethics, 2022, 22, 39-42.	0.5	3
36	Individual and family characteristics associated with health indicators at entry into multidisciplinary pediatric weight management: findings from the CANadian Pediatric Weight management Registry (CANPWR). International Journal of Obesity, 2021, , .	1.6	2

#	Article	IF	Citations
37	Family Experiences with Care for Children with Inherited Metabolic Diseases in Canada: A Cross-Sectional Survey. Patient, 2022, 15, 171-185.	1.1	1
38	MG-114â€…First 2 years of experience of an integrated multidisciplinary clinic for adults with aortopathies in a canadian context. Journal of Medical Genetics, 2015, 52, A3.1-A3.	1.5	0