Joann N Bodurtha

List of Publications by Year
in descending order

42 papers	645 citations	686830 13 h-index	610482 24 g-index
43 all docs	43 docs citations	43 times ranked	1292 citing authors

#	Article	IF	Citations
1	The natural history of OTOF-related auditory neuropathy spectrum disorders: a multicenter study. Human Genetics, 2022, 141, 853-863.	1.8	7
2	â€œItâ€™s a Little Different for Menâ€â€”Sponsorship and Gender in Academic Medicine: a Qualitative Study. Journal of General Internal Medicine, 2021, 36, 1-8.	1.3	35
3	The 2019 US medical genetics workforce: a focus on clinical genetics. Genetics in Medicine, 2021, 23, 1458-1464.	1.1	70
4	Demographic and socioeconomic trends in DNA banking utilization in the USA. Journal of Community Genetics, 2021, 12, 593-602.	0.5	1
5	Response to Biesecker etÂal.. American Journal of Human Genetics, 2021, 108, 1807-1808.	2.6	3
6	Linkage-specific deubiquitylation by OTUD5 defines an embryonic pathway intolerant to genomic variation. Science Advances, 2021, 7, .	4.7	25
7	Lung cancer and family-centered concerns. Supportive Care in Cancer, 2020, 28, 497-505.	1.0	2
8	Regional models of genetic services in the United States. Genetics in Medicine, 2020, 22, 381-388.	1.1	10
9	A structured genetics rotation for pediatric residents: an important educational opportunity. Genetics in Medicine, 2020, 22, 793-796.	1.1	9
10	Post Graduate Education: Is Genomics Included on Board Certification Exams?. Medical Science Educator, 2020, 30, 1379-1382.	0.7	1
11	Gene Therapy. Pediatrics in Review, 2020, 41, 606-608.	0.2	1
12	Identifying Gender Disparities and Barriers to Measuring the Status of Female Faculty: The Experience of a Large School of Medicine. Journal of Women's Health, 2019, 28, 1569-1575.	1.5	17
13	Factors affecting quality of life in children and adolescents with hypermobile Ehlersâ€Danlos syndrome/hypermobility spectrum disorders. American Journal of Medical Genetics, Part A, 2019, 179, 561-569.	0.7	39
14	Uncertainty, hope, and coping efficacy among mothers of children with Duchenne/Becker muscular dystrophy. Clinical Genetics, 2019, 95, 677-683.	1.0	13
15	Family health history and genetic servicesâ€”the East Baltimore community stakeholder interview project. Journal of Community Genetics, 2019, 10, 219-227.	0.5	8
16	Highâ€Risk Palliative Care Patientsâ€™ Knowledge and Attitudes about Hereditary Cancer Testing and DNA Banking. Journal of Genetic Counseling, 2018, 27, 834-843.	0.9	6
17	Juvenile Idiopathic Arthritis Associated with Combined JPâ€“HHT Syndrome: A Novel Phenotype Associated with a Novel Variant in SMAD4. Journal of Pediatric Genetics, 2018, 07, 078-082.	0.3	3
18	A Cost Analysis of Universal versus Targeted Cholesterol Screening in Pediatrics. Journal of Pediatrics, 2018, 196, 201-207.e2.	0.9	9

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19	Lung cancer and family-centered patient concerns. Supportive Care in Cancer, 2018, 26, 3047-3053.	1.0	4
20	The Wills Eye Handbook of Ocular Genetics, 1st Edition. Journal of Pediatric Genetics, 2018, 07, 191-192.	0.3	1
21	Pain and sleep quality in children with nonâ€vascular Ehlersâ€“Danlos syndromes. American Journal of Medical Genetics, Part A, 2018, 176, 1858-1864.	0.7	12
22	The Role of Palliative Medicine in Assessing Hereditary Cancer Risk. American Journal of Hospice and Palliative Medicine, 2018, 35, 1490-1497.	0.8	5
23	PhenX measures for phenotyping rare genetic conditions. Genetics in Medicine, 2017, 19, 834-837.	1.1	2
24	Response to Laissue et al.. Genetics in Medicine, 2017, 19, 1380-1380.	1.1	0
25	Milestones for medical students completing a clinical genetics elective. Genetics in Medicine, 2017, 19, 236-239.	1.1	5
26	Whole-exome sequencing in the molecular diagnosis of individuals with congenital anomalies of the kidney and urinary tract and identification of a new causative gene. Genetics in Medicine, 2017, 19, 412-420.	1.1	73
27	Essential Mednotesâ€”Comprehensive Medical Reference & Review for USMLE II and MCCQE I. Journal of Pediatric Genetics, 2017, 06, 128-128.	0.3	0
28	Five Pediatric Cancers â€“ Update on Genetic Implications. Current Pediatric Reviews, 2017, 13, 42-48.	0.4	3
29	De novo missense variants in PPP1CB are associated with intellectual disability and congenital heart disease. Human Genetics, 2016, 135, 1399-1409.	1.8	40
30	Genotypeâ€“phenotype correlation of congenital anomalies in multiple congenital anomalies hypotonia seizures syndrome (MCAHS1)/PIGNâ€related epilepsy. American Journal of Medical Genetics, Part A, 2016, 170, 77-86.	0.7	41
31	Incontinentia Pigmenti with Persistent Hypercalcemia: Case Report. Pediatric Dermatology, 2016, 33, e315-7.	0.5	3
32	Family Ties: The Role of Family Context in Family Health History Communication About Cancer. Journal of Health Communication, 2016, 21, 346-355.	1.2	19
33	Trends in Unmet Need for Genetic Counseling Among Children With Special Health Care Needs, 2001â€“2010. Academic Pediatrics, 2015, 15, 544-550.	1.0	7
34	Horizontal integration of OMIM across the medical school preclinical curriculum for early reinforcement of clinical genetics principles. Genetics in Medicine, 2015, 17, 158-163.	1.1	6
35	Hereditary Renal Hypouricemia: A New Role for Allopurinol?. American Journal of Medicine, 2014, 127, e3-e4.	0.6	47
36	Patient-reported hereditary breast and ovarian cancer in a primary care practice. Journal of Community Genetics, 2014, 5, 179-183.	0.5	10

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37	The KinFact Intervention â€“ A Randomized Controlled Trial to Increase Family Communication About Cancer History. Journal of Women's Health, 2014, 23, 806-816.	1.5	28
38	Genomics and Perinatal Care. New England Journal of Medicine, 2012, 366, 64-73.	13.9	40
39	46 Chromosomes and Me. Journal of Genetic Counseling, 2012, 21, 173-174.	0.9	2
40	Physicians' Current Practices and Opportunities for DNA Banking of Dying Patients With Cancer. Journal of Oncology Practice, 2011, 7, 183-187.	2.5	20
41	Exploring Hereditary Cancer Among Dying Cancer Patientsâ€”A Cross-Sectional Study of Hereditary Risk and Perceived Awareness of DNA Testing and Banking. Journal of Genetic Counseling, 2010, 19, 497-525.	0.9	18
42	Successful Treatment of Refractory Orthostatic Intolerance (OI) With Droxidopa. Clinical Pediatrics, 0, , 000992282210926.	0.4	0