

David T Miller

List of Publications by Year in descending order

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Version: 2024-02-01

37
papers

5,797
citations

361413
20
h-index

345221
36
g-index

40
all docs

40
docs citations

40
times ranked

9651
citing authors

#	ARTICLE	IF	CITATIONS
1	Clinical validity assessment of genes frequently tested on intellectual disability/autism sequencing panels. <i>Genetics in Medicine</i> , 2022, 24, 1899-1908.	2.4	9
2	ACMG SF v3.1 list for reporting of secondary findings in clinical exome and genome sequencing: A policy statement of the American College of Medical Genetics and Genomics (ACMG). <i>Genetics in Medicine</i> , 2022, 24, 1407-1414.	2.4	119
3	Growth, development, and phenotypic spectrum of individuals with deletions of 2q33.1 involving <i>SATB2</i> . <i>Clinical Genetics</i> , 2021, 99, 547-557.	2.0	13
4	Clinical Syndromic Phenotypes and the Potential Role of Genetics in Pulmonary Vein Stenosis. <i>Children</i> , 2021, 8, 128.	1.5	5
5	Recommendations for reporting of secondary findings in clinical exome and genome sequencing, 2021 update: a policy statement of the American College of Medical Genetics and Genomics (ACMG). <i>Genetics in Medicine</i> , 2021, 23, 1391-1398.	2.4	145
6	ACMG SF v3.0 list for reporting of secondary findings in clinical exome and genome sequencing: a policy statement of the American College of Medical Genetics and Genomics (ACMG). <i>Genetics in Medicine</i> , 2021, 23, 1381-1390.	2.4	356
7	Genotype-phenotype correlations and novel molecular insights into the <i>DHX30</i> -associated neurodevelopmental disorders. <i>Genome Medicine</i> , 2021, 13, 90.	8.2	16
8	Chromosomal microarray analysis, including constitutional and neoplastic disease applications, 2021 revision: a technical standard of the American College of Medical Genetics and Genomics (ACMG). <i>Genetics in Medicine</i> , 2021, 23, 1818-1829.	2.4	18
9	Response to McGurk et al. <i>Genetics in Medicine</i> , 2021, , .	2.4	0
10	Genetics of human malignant peripheral nerve sheath tumors. <i>Neuro-Oncology Advances</i> , 2020, 2, i50-i61.	0.7	34
11	Insufficient Evidence for "Autism-Specific" Genes. <i>American Journal of Human Genetics</i> , 2020, 106, 587-595.	6.2	110
12	Points to consider for reporting of germline variation in patients undergoing tumor testing: a statement of the American College of Medical Genetics and Genomics (ACMG). <i>Genetics in Medicine</i> , 2020, 22, 1142-1148.	2.4	59
13	Systematic evidence-based review: outcomes from exome and genome sequencing for pediatric patients with congenital anomalies or intellectual disability. <i>Genetics in Medicine</i> , 2020, 22, 986-1004.	2.4	53
14	Genomics of MPNST (GeM) Consortium: Rationale and Study Design for Multi-Omic Characterization of NF1-Associated and Sporadic MPNSTs. <i>Genes</i> , 2020, 11, 387.	2.4	16
15	Meta-analysis and multidisciplinary consensus statement: exome sequencing is a first-tier clinical diagnostic test for individuals with neurodevelopmental disorders. <i>Genetics in Medicine</i> , 2019, 21, 2413-2421.	2.4	378
16	Response to Knoppers et al.. <i>Genetics in Medicine</i> , 2019, 21, 2403.	2.4	0
17	Expanding the clinical phenotype of individuals with a 3-bp in-frame deletion of the <i>NF1</i> gene (c.2970_2972del): an update of genotype-phenotype correlation. <i>Genetics in Medicine</i> , 2019, 21, 867-876.	2.4	62
18	Patient re-contact after revision of genomic test results: points to consider—a statement of the American College of Medical Genetics and Genomics (ACMG). <i>Genetics in Medicine</i> , 2019, 21, 769-771.	2.4	91

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19	Genotype-Phenotype Correlation in NF1: Evidence for a More Severe Phenotype Associated with Missense Mutations Affecting NF1 Codons 844-848. <i>American Journal of Human Genetics</i> , 2018, 102, 69-87.	6.2	144
20	School liaison program supporting children with neurofibromatosis type 1: a model of care for children with chronic disease. <i>Genetics in Medicine</i> , 2018, 20, 785-788.	2.4	3
21	Yield of additional genetic testing after chromosomal microarray for diagnosis of neurodevelopmental disability and congenital anomalies: a clinical practice resource of the American College of Medical Genetics and Genomics (ACMG). <i>Genetics in Medicine</i> , 2018, 20, 1105-1113.	2.4	57
22	Response to Biesecker. <i>Genetics in Medicine</i> , 2017, 19, 605.	2.4	0
23	Recommendations for reporting of secondary findings in clinical exome and genome sequencing, 2016 update (ACMG SF v2.0): a policy statement of the American College of Medical Genetics and Genomics. <i>Genetics in Medicine</i> , 2017, 19, 249-255.	2.4	1,398
24	<i>BRAT1</i> mutations present with a spectrum of clinical severity. <i>American Journal of Medical Genetics, Part A</i> , 2016, 170, 2265-2273.	1.2	34
25	A Case of HDR Syndrome and Ichthyosis: Dual Diagnosis by Whole-Genome Sequencing of Novel Mutations in <i>GATA3</i> and <i>STS</i> Genes. <i>Journal of Clinical Endocrinology and Metabolism</i> , 2016, 101, 837-840.	3.6	9
26	A Clinician's perspective on clinical exome sequencing. <i>Human Genetics</i> , 2016, 135, 643-654.	3.8	33
27	Classifying Germline Sequence Variants in the Era of Next-Generation Sequencing. <i>Clinical Chemistry</i> , 2016, 62, 799-806.	3.2	0
28	GenomeConnect: Matchmaking Between Patients, Clinical Laboratories, and Researchers to Improve Genomic Knowledge. <i>Human Mutation</i> , 2015, 36, 974-978.	2.5	56
29	Advances in Genetic Discovery and Implications for Counseling of Patients and Families with Autism Spectrum Disorders. <i>Current Genetic Medicine Reports</i> , 2014, 2, 124-134.	1.9	7
30	Advances in Genetic Diagnosis of Autism Spectrum Disorders. <i>Current Pediatrics Reports</i> , 2014, 2, 71-81.	4.0	3
31	Treating the Whole Person With Autism: The Proceedings of the Autism Speaks National Autism Conference. <i>Current Problems in Pediatric and Adolescent Health Care</i> , 2014, 44, 26-47.	1.7	12
32	Oligonucleotide Microarrays for Clinical Diagnosis of Copy Number Variation and Zygosity Status. <i>Current Protocols in Human Genetics</i> , 2012, 74, Unit8.12.	3.5	11
33	Chromosomal microarray testing influences medical management. <i>Genetics in Medicine</i> , 2011, 13, 770-776.	2.4	107
34	Consensus Statement: Chromosomal Microarray Is a First-Tier Clinical Diagnostic Test for Individuals with Developmental Disabilities or Congenital Anomalies. <i>American Journal of Human Genetics</i> , 2010, 86, 749-764.	6.2	2,325
35	Genetic Testing for Developmental Delay: Keep Searching for an Answer. <i>Clinical Chemistry</i> , 2009, 55, 827-830.	3.2	3
36	Oligonucleotide Microarrays for Clinical Diagnosis of Copy Number Variation. <i>Current Protocols in Human Genetics</i> , 2008, 58, Unit 8.12.	3.5	6

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37	Atherosclerosis. Journal of the American College of Cardiology, 2007, 49, 1589-1599.	2.8	63