

Michael J Field

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/1175740/publications.pdf>

Version: 2024-02-01

21
papers

391
citations

1040056

9
h-index

794594

19
g-index

22
all docs

22
docs citations

22
times ranked

966
citing authors

#	ARTICLE	IF	CITATIONS
1	De Novo Loss-of-Function Mutations in SETD5, Encoding a Methyltransferase in a 3p25 Microdeletion Syndrome Critical Region, Cause Intellectual Disability. American Journal of Human Genetics, 2014, 94, 618-624.	6.2	96
2	Bayesian approach to determining penetrance of pathogenic SDH variants. Journal of Medical Genetics, 2018, 55, 729-734.	3.2	44
3	Early Detection of Fragile X Syndrome: Applications of a Novel Approach for Improved Quantitative Methylation Analysis in Venous Blood and Newborn Blood Spots. Clinical Chemistry, 2014, 60, 963-973.	3.2	43
4	Lessons learnt from implementation of a Lynch syndrome screening program for patients with gynaecological malignancy. Pathology, 2017, 49, 457-464.	0.6	34
5	Streamlined genetic education is effective in preparing women newly diagnosed with breast cancer for decision making about treatment-focused genetic testing: a randomized controlled noninferiority trial. Genetics in Medicine, 2017, 19, 448-456.	2.4	30
6	Maternal attitudes to newborn screening for fragile X syndrome. American Journal of Medical Genetics, Part A, 2013, 161, 301-311.	1.2	24
7	Incomplete silencing of full mutation alleles in males with fragile X syndrome is associated with autistic features. Molecular Autism, 2019, 10, 21.	4.9	20
8	Relationships between UBE3A and SNORD116 expression and features of autism in chromosome 15 imprinting disorders. Translational Psychiatry, 2020, 10, 362.	4.8	14
9	Feasibility of Screening for Chromosome 15 Imprinting Disorders in 16,579 Newborns by Using a Novel Genomic Workflow. JAMA Network Open, 2022, 5, e2141911.	5.9	14
10	MEN4, the MEN1 Mimicker: A Case Series of three Phenotypically Heterogenous Patients With Unique CDKN1B Mutations. Journal of Clinical Endocrinology and Metabolism, 2022, 107, 2339-2349.	3.6	14
11	Psychological outcomes and surgical decisions after genetic testing in women newly diagnosed with breast cancer with and without a family history. European Journal of Human Genetics, 2018, 26, 972-983.	2.8	12
12	Surveillance Improves Outcomes for Carriers of SDHB Pathogenic Variants: A Multicenter Study. Journal of Clinical Endocrinology and Metabolism, 2022, 107, e1907-e1916.	3.6	11
13	Rapid detection of BRCA1/2 recurrent mutations in Chinese breast and ovarian cancer patients with multiplex SNaPshot genotyping panels. Oncotarget, 2018, 9, 7832-7843.	1.8	9
14	Current perspectives on medical education in China. Medical Education, 2006, 40, 938-939.	2.1	8
15	EFFECT OF ATRIAL NATRIURETIC PEPTIDE ON CELLULAR ELEMENT CONCENTRATIONS IN RAT PROXIMAL TUBULES: EVIDENCE FOR INHIBITION OF THE SODIUM PUMP. Clinical and Experimental Pharmacology and Physiology, 1994, 21, 775-780.	1.9	5
16	Are We Ready for Fragile X Newborn Screening Testing? Lessons Learnt from a Feasibility Study. International Journal of Neonatal Screening, 2018, 4, 9.	3.2	4
17	Skin rash, a kidney mass and a family mystery dating back to World War II. Medical Journal of Australia, 2014, 201, 58-60.	1.7	3
18	Implementing gene curation for hereditary cancer susceptibility in Australia: achieving consensus on genes with clinical utility. Journal of Medical Genetics, 2021, 58, 853-858.	3.2	3

#	ARTICLE	IF	CITATIONS
19	Role of insulin-like growth factor binding proteins in human post-nephrectomy proximal tubule cells. Journal of Physiology, 1998, 508, 587-595.	2.9	1
20	IDMOD: An Australian microsimulation model of lifetime economic and social factors in familial intellectual disability. , 2020, 13, 52-66.		1
21	Acute infusion of amphotericin B: Proximal tubular effects. Nephrology, 1996, 2, 399-403.	1.6	0