Michael J Field

List of Publications by Year
in descending order

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	Feasibility of Screening for Chromosome 15 Imprinting Disorders in 16â€¯579 Newborns by Using a Novel Genomic Workflow. JAMA Network Open, 2022, 5, e2141911.	5.9	14
2	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
3	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
4	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
5	Relationships between UBE3A and SNORD116 expression and features of autism in chromosome 15 imprinting disorders. Translational Psychiatry, 2020, 10, 362.	4.8	14
6	IDMOD: An Australian microsimulation model of lifetime economic and social factors in familial intellectual disability. , 2020, 13, 52-66.		1
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	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
9	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
10	Bayesian approach to determining penetrance of pathogenic SDH variants. Journal of Medical Genetics, 2018, 55, 729-734.	3.2	44
11	Are We Ready for Fragile X Newborn Screening Testing?â€”Lessons Learnt from a Feasibility Study. International Journal of Neonatal Screening, 2018, 4, 9.	3.2	4
12	Lessons learnt from implementation of a Lynch syndrome screening program for patients with gynaecological malignancy. Pathology, 2017, 49, 457-464.	0.6	34
13	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
14	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
15	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
16	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
17	Maternal attitudes to newborn screening for fragile X syndrome. American Journal of Medical Genetics, Part A, 2013, 161, 301-311.	1.2	24
18	Current perspectives on medical education in China. Medical Education, 2006, 40, 938-939.	2.1	8

#	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	Acute infusion of amphotericin B: Proximal tubular effects. Nephrology, 1996, 2, 399-403.	1.6	0
21	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