Consanguineous marriages

Journal of Community Genetics 3, 185-192

DOI: 10.1007/s12687-011-0072-y

Citation Report

#	Article	IF	CITATIONS
2	Variability in laboratory reporting practices for regions of homozygosity indicating parental relatedness as identified by SNP microarray testing. Genetics in Medicine, 2012, 14, 971-976.	2.4	21
3	What can be offered to couples at (possibly) increased genetic risk?. Journal of Community Genetics, 2012, 3, 167-174.	1.2	6
4	Editorial: genetic aspects of preconception consultation in primary care. Journal of Community Genetics, 2012, 3, 155-157.	1.2	2
5	Challenges in the care for consanguineous couples: an exploratory interview study among general practitioners and midwives. BMC Family Practice, 2012, 13, 105.	2.9	10
6	The first five years of a preventive programme for haemoglobinopathies in Northeastern Iraq. Journal of Medical Screening, 2013, 20, 171-176.	2.3	16
7	Consanguinity, endogamy, and genetic disorders in Tunisia. Journal of Community Genetics, 2013, 4, 273-284.	1.2	79
8	β- And α-Thalassemia Intermedia in Basra, Southern Iraq. Hemoglobin, 2013, 37, 553-563.	0.8	8
9	Bone marrow transplantation for thalassemia from alternative related donors: improved outcomes with a new approach. Blood, 2013, 122, 2751-2756.	1.4	44
10	Training Community Health Workers to discuss cousin marriage and genetic risk with migrant women in the Netherlands (Family, Health and Genetics, G3-project phase I). European Journal of Public Health, 2014, 24, .	0.3	0
11	Specific Aspects of Consanguinity: Some Examples from the Tunisian Population. Human Heredity, 2014, 77, 167-174.	0.8	21
12	A long-term follow up of premarital counseling in the Israeli Arab population. Journal of Community Genetics, 2014, 5, 377-381.	1.2	4
13	Consanguineous marriage and reproductive risk: attitudes and understanding of ethnic groups practising consanguinity in Western society. European Journal of Human Genetics, 2014, 22, 452-457.	2.8	34
14	Wilson′s disease: An endocrine revelation. Indian Journal of Endocrinology and Metabolism, 2014, 18, 855.	0.4	17
15	Variable approaches to genetic counseling for microarray regions of homozygosity associated with parental relatedness. American Journal of Medical Genetics, Part A, 2014, 164, 87-98.	1.2	8
16	Understanding Utilization of Outpatient Clinics for Children with Special Health Care Needs in Southern Israel. Maternal and Child Health Journal, 2014, 18, 1831-1845.	1.5	4
17	Consanguinity profile in the Gaza Strip of Palestine: Large-scale community-based study. European Journal of Medical Genetics, 2014, 57, 90-94.	1.3	21
18	The prevalence of isolated growth hormone deficiency among children of short stature in Jordan and its relationship with consanguinity. Clinical Endocrinology, 2014, 81, 876-882.	2.4	8
19	Immunoglobulin genes in Andalusia (Spain). Genetic diversity in the Mediterranean space. Comptes Rendus - Biologies, 2014, 337, 646-656.	0.2	9

#	Article	IF	CITATIONS
20	Diagnostic Exome Sequencing to Elucidate the Genetic Basis of Likely Recessive Disorders in Consanguineous Families. Human Mutation, 2014, 35, 1203-1210.	2.5	75
21	Prevalence of consanguineous marriages and associated factors among Israeli Bedouins. Journal of Community Genetics, 2014, 5, 395-398.	1.2	29
22	Generalized odontodysplasia in a 5-year-old patient with Hallermann-Streiff syndrome: clinical aspects, cone beam computed tomography findings, and conservative clinical approach. Oral Surgery, Oral Medicine, Oral Pathology and Oral Radiology, 2014, 118, e58-e64.	0.4	11
23	Modelling the constraints on consanguineous marriage when fertility declines. Demographic Research, 0, 30, 277-312.	3.0	18
24	Genetics of consanguineous marriage: Impact and importance of counseling. Journal of Pediatric Genetics, 2015, 01, 217-220.	0.7	6
26	Diagnosis and treatment of late-onset Pompe disease in the Middle East and North Africa region: consensus recommendations from an expert group. BMC Neurology, 2015, 15, 205.	1.8	28
27	Migration und Schwangerschaft: Aspekte der PrÄøention und Versorgung. Public Health Forum, 2015, 23, 64-66.	0.2	0
28	A Registerâ€Based Study of Diseases With an Autosomal Recessive Origin in Small Children in <scp>D</scp> enmark According to Maternal Country of Origin. Paediatric and Perinatal Epidemiology, 2015, 29, 351-359.	1.7	5
29	Effect of consanguinity on birth defects in Saudi women: Results from a nested caseâ€control study. Birth Defects Research Part A: Clinical and Molecular Teratology, 2015, 103, 100-104.	1.6	28
30	Cooperation Drives Competition among Tsimane Women in the Bolivian Amazon. , 2015, , .		0
31	Impact of consanguinity on cardio-metabolic health and other diseases: findings from an Afro-Indian tribal community. Journal of Community Genetics, 2015, 6, 129-135.	1.2	4
32	Exploring the Effectiveness of Mandatory Premarital Screening and Genetic Counselling Programmes for \hat{l}^2 -Thalassaemia in the Middle East: A Scoping Review. Public Health Genomics, 2015, 18, 193-203.	1.0	76
33	NADf Chip, a Two-Color Microarray for Simultaneous Screening of Multigene Mutations Associated with Hearing Impairment in North African Mediterranean Countries. Journal of Molecular Diagnostics, 2015, 17, 155-161.	2.8	20
34	A study of consanguineous marriage as a risk factor for developing comitant strabismus. Journal of Community Genetics, 2015, 6, 177-180.	1.2	10
35	Continuous decrease of consanguineous marriages among arabs in israel. American Journal of Human Biology, 2015, 27, 94-98.	1.6	29
36	Homozygous/Compound Heterozygous Triadin Mutations Associated With Autosomal-Recessive Long-QT Syndrome and Pediatric Sudden Cardiac Arrest. Circulation, 2015, 131, 2051-2060.	1.6	92
37	Irrigation and human niche construction. An example of socio-spatial organisation in the Zerqa Triangle, Jordan. Water History, 2015, 7, 441-454.	1.3	4
38	Diagnostic tools and strategies for assessing disease progression in Alkaptonuria. Expert Opinion on Orphan Drugs, 2015, 3, 705-717.	0.8	1

#	Article	IF	Citations
39	Perspective: Encourage the innovators. Nature, 2015, 528, S7-S7.	27.8	14
40	Dysferlinopathy in Iran: Clinical and genetic report. Journal of the Neurological Sciences, 2015, 359, 256-259.	0.6	12
41	The Provision of Medical and Health Genetics and Genomics in the Developing World., 2016,, 285-294.		4
42	The Effect of Consanguineous Marriage on Mental Health among the Students of the Shahrekord University of Medical Sciences. Journal of Clinical and Diagnostic Research JCDR, 2016, 10, GC01-GC04.	0.8	6
43	Epidemiology of Transfusion Transmitted Infection among Patients with $\langle i \rangle \hat{l}^2 \langle i \rangle$. Thalassaemia Major in Pakistan. Journal of Blood Transfusion, 2016, 2016, 1-5.	3.3	35
44	Segregation of Incomplete Achromatopsia and Alopecia Due to PDE6H and LPAR6 Variants in a Consanguineous Family from Pakistan. Genes, 2016, 7, 41.	2.4	8
45	A unique <i>LAMB3 < /i> splice-site mutation with founder effect from the Balkans causes lethal epidermolysis bullosa in several European countries. British Journal of Dermatology, 2016, 175, 721-727.</i>	1.5	12
46	CHANGES IN MARRIAGE PATTERNS AMONG THE ARAB COMMUNITY IN ISRAEL OVER A 60-YEAR PERIOD. Journal of Biosocial Science, 2016, 48, 283-287.	1.2	19
47	Stillbirth and congenital anomalies in migrants in Europe. Best Practice and Research in Clinical Obstetrics and Gynaecology, 2016, 32, 50-59.	2.8	22
48	Meckel–Gruber syndrome: prevalence from a hospital-based study in Oman. Journal of Maternal-Fetal and Neonatal Medicine, 2016, 29, 3696-3698.	1.5	0
49	The Present and Future Global Burden of the Inherited Disorders of Hemoglobin. Hematology/Oncology Clinics of North America, 2016, 30, 327-341.	2.2	63
50	Characterizing the genetic structure of a forensic DNA database using a latent variable approach. Forensic Science International: Genetics, 2016, 23, 130-149.	3.1	4
51	Prevalence of Childhood Blindness and Ocular Morbidity in a Rural Pediatric Population in Southern India: The Pavagada Pediatric Eye Disease Study-1. Ophthalmic Epidemiology, 2016, 23, 185-192.	1.7	41
52	Arranged marriages in people with epilepsy: A pilot knowledge, attitudes and practices survey from India. International Journal of Epilepsy, 2016, 03, 075-079.	0.5	0
53	Whole Exome Screening Identifies Novel and Recurrent WISP3 Mutations Causing Progressive Pseudorheumatoid Dysplasia in Jammu and Kashmir-India. Scientific Reports, 2016, 6, 27684.	3.3	13
54	Responding to the increased genetic risk associated with customary consanguineous marriage among minority ethnic populations: lessons from local innovations in England. Journal of Community Genetics, 2016, 7, 215-228.	1.2	13
55	Consanguinity and associated perinatal outcomes, including stillbirth. Australian and New Zealand Journal of Obstetrics and Gynaecology, 2016, 56, 599-604.	1.0	13
56	Community engagement and education: addressing the needs of South Asian families with genetic disorders. Journal of Community Genetics, 2016, 7, 317-323.	1.2	7

#	Article	IF	CITATIONS
57	Differential impact of consanguineous marriages on autosomal recessive diseases in <pre><scp>T</scp>unisia</pre> . American Journal of Human Biology, 2016, 28, 171-180.	1.6	21
58	Assessing women's knowledge and attitudes toward cord blood banking: policy and ethical implications for Jordan. Transfusion, 2016, 56, 2052-2061.	1.6	9
59	Wide disparity of clinical genetics services and EU rare disease research funding across Europe. Journal of Community Genetics, 2016, 7, 119-126.	1.2	20
60	Consanguinity and autosomal recessive neuromuscular disorders. Developmental Medicine and Child Neurology, 2016, 58, 796-797.	2.1	2
61	A review of consanguinity in Ireland—estimation of frequency and approaches to mitigate risks. Irish Journal of Medical Science, 2016, 185, 17-28.	1.5	3
62	Paediatric orphan lung diseases in Asia. Lancet Respiratory Medicine, the, 2016, 4, 174-175.	10.7	1
63	Identification of six novel mutations in Iranian patients with maple syrup urine disease and their in silico analysisâ€∢. Mutation Research - Fundamental and Molecular Mechanisms of Mutagenesis, 2016, 786, 34-40.	1.0	12
64	ConStruct 1.0: An R Script to distinguish between substructure and consanguinity within a population using multilocus microsatellite data. Cogent Biology, 2016, 2, 1128317.	1.7	1
65	Consanguinity and the Risk of Hashimoto's Thyroiditis. Thyroid, 2017, 27, 390-395.	4.5	2
66	Loss-of-Function Variants in MYLK Cause Recessive Megacystis Microcolon Intestinal Hypoperistalsis Syndrome. American Journal of Human Genetics, 2017, 101, 123-129.	6.2	67
67	Endogamous marriage and the prevalence of hemoglobin E in ethnic groups of northern Thailand. Asian Pacific Journal of Tropical Medicine, 2017, 10, 414-417.	0.8	0
68	A multicenter clinical exome study in unselected cohorts from a consanguineous population of Saudi Arabia demonstrated a high diagnostic yield. Molecular Genetics and Metabolism, 2017, 121, 91-95.	1.1	68
69	Genetic Predisposition to Sporadic Congenital Hearing Loss in a Pediatric Population. Scientific Reports, 2017, 7, 45973.	3.3	28
70	The influence of family violence and child marriage on unmet need for family planning in Jordan. Journal of Family Planning and Reproductive Health Care, 2017, 43, 105-112.	0.8	19
71	Under-Five Child Mortality and Morbidity Associated with Consanguineous Child Marriage in Pakistan: Retrospective Analysis using Pakistan Demographic and Health Surveys, 1990–91, 2006–07, 2012–13. Maternal and Child Health Journal, 2017, 21, 1095-1104.	1.5	12
72	Prevalence of visual and hearing impairment in adults with intellectual disabilities in the southwestern Iran: a cross sectional study. International Journal on Disability and Human Development, 2017, 16, .	0.2	0
73	From genomes to genomic medicine: enabling personalized and precision medicine in the Middle East. Personalized Medicine, 2017, 14, 377-382.	1.5	17
74	The risk ratio for development of hereditary sensorineural hearing loss in consanguineous marriage offspring. International Journal of Pediatric Otorhinolaryngology, 2017, 101, 7-10.	1.0	10

#	Article	IF	CITATIONS
75	Molecular epidemiology of junctional epidermolysis bullosa: discovery of novel and frequent <i>LAMB3</i> mutations in Chilean patients with diagnostic significance. British Journal of Dermatology, 2017, 176, 1090-1092.	1.5	13
76	First report of pediatric hematopoietic stem cell transplantation activities in the eastern mediterranean region from 1984 to 2011: on behalf of the pediatric cancer working committee of the eastern mediterranean blood and marrow transplantation group. Bone Marrow Transplantation, 2017, 52. 120-125.	2.4	8
77	Family History of Early Infant Death Correlates with Earlier Age at Diagnosis But Not Shorter Time to Diagnosis for Severe Combined Immunodeficiency. Frontiers in Immunology, 2017, 8, 808.	4.8	34
78	Morquio's Syndrome: A Case Report of Two Siblings. Case Reports in Dentistry, 2017, 2017, 1-4.	0.5	2
79	Oral and Periodontal Diseases in Consanguineous Marriages. , 2017, , .		1
80	Consanguineous Marriage and the Psychopathology of Progeny. JAMA Psychiatry, 2018, 75, 438.	11.0	11
81	Assessing Intimate Partner Abuse: Associated Factors and Health Consequences among Jordanian Women. Issues in Mental Health Nursing, 2018, 39, 344-352.	1.2	11
82	Single Nucleotide Polymorphism-Based Noninvasive Prenatal Testing: Experience in India. Journal of Obstetrics and Gynecology of India, 2018, 68, 462-470.	0.9	7
83	Global Hearing Loss Prevention. Otolaryngologic Clinics of North America, 2018, 51, 575-592.	1.1	65
84	Genetic characterisation of 19 autosomal STR loci in a population sample from the Southeastern Anatolia Region of Turkey. Annals of Human Biology, 2018, 45, 148-159.	1.0	1
85	Mapping autosomal recessive intellectual disability: combined microarray and exome sequencing identifies 26 novel candidate genes in 192 consanguineous families. Molecular Psychiatry, 2018, 23, 973-984.	7.9	147
86	Women with Inherited Bleeding Disorders in Different Cultural Settings. , 0, , 225-233.		0
87	Unravelling the genetic architecture of autosomal recessive epilepsy in the genomic era. Journal of Neurogenetics, 2018, 32, 295-312.	1.4	7
88	Cousin Marriage Is Not Choice: Muslim Marriage and Underdevelopment. AEA Papers and Proceedings American Economic Association, 2018, 108, 353-357.	1.2	13
89	Prevalence of hereditary cancer susceptibility syndromes in children with cancer in a highly consanguineous population. Cancer Epidemiology, 2018, 55, 88-95.	1.9	13
90	Autozygosity mapping of methylmalonic acidemia associated genes by short tandem repeat markers facilitates the identification of five novel mutations in an Iranian patient cohort. Metabolic Brain Disease, 2018, 33, 1689-1697.	2.9	5
91	Reproductive choices: a qualitative study of Dutch Moroccan and Turkish consanguineously married women's perspectives on preconception carrier screening. BMC Women's Health, 2018, 18, 79.	2.0	9
92	Long-term exposure to ambient air pollution and autism spectrum disorder in children: A case-control study in Tehran, Iran. Science of the Total Environment, 2018, 643, 1216-1222.	8.0	49

#	Article	IF	CITATIONS
93	The conserved p.Arg108 residue in S1PR2 (DFNB68) is fundamental for proper hearing: evidence from a consanguineous Iranian family. BMC Medical Genetics, 2018, 19, 81.	2.1	10
94	MMP-2: is too low as bad as too high in the cardiovascular system?. American Journal of Physiology - Heart and Circulatory Physiology, 2018, 315, H1332-H1340.	3.2	37
95	Congenital Myasthenic Syndrome: Spectrum of Mutations in an Indian Cohort. Journal of Clinical Neuromuscular Disease, 2018, 20, 14-27.	0.7	18
96	Current trends of consanguineous marriages and its association with socio-demographic variables in Pakistan. International Journal of Reproduction, Contraception, Obstetrics and Gynecology, 2018, 7, 1699.	0.1	13
97	Genetic heterogeneity of congenital hearing impairment in Algerians from the Gharda $ ilde{A}^-$ a province. International Journal of Pediatric Otorhinolaryngology, 2018, 112, 1-5.	1.0	14
98	Prenatal and preconception genetic counseling for consanguinity: Consanguineous couples' expectations, experiences, and perspectives. Journal of Genetic Counseling, 2019, 28, 982-992.	1.6	10
99	75. A FAMILY OF MATCHED PARENT-CHILD HLA HAPLOTYPES: A CASE STUDY FROM BAHRAIN. Reproductive BioMedicine Online, 2019, 39, e73.	2.4	0
100	Papillon-LefÃ'vre Syndrome: Diagnosis, Dental Management, and a Case Report. Case Reports in Dentistry, 2019, 2019, 1-7.	0.5	11
101	Spatial distribution of healthcare access and utilization: do they affect health outcomes in Turkey?. Middle East Development Journal, 2019, 11, 124-163.	0.8	3
102	Chromosomal aberrations in pregnancy and fetal loss: Insight on the effect of consanguinity, review of 1625 cases. Molecular Genetics & Enomic Medicine, 2019, 7, e820.	1.2	10
103	Can parental consanguinity be a risk factor for the occurrence of nonsyndromic oral cleft?. Early Human Development, 2019, 135, 23-26.	1.8	11
104	Frequency of Ambiguous Genitalia in 14,177 Newborns in Turkey. Journal of the Endocrine Society, 2019, 3, 1185-1195.	0.2	14
105	Quantifying the Levels of Knowledge, Attitude, and Practice Associated with Sickle Cell Disease and Premarital Genetic Counseling in 350 Saudi Adults. Advances in Hematology, 2019, 2019, 1-7.	1.0	16
106	The Influence of Consanguinity on Familial Clefting Among Palestinians. Cleft Palate-Craniofacial Journal, 2019, 56, 1072-1079.	0.9	9
107	Keeping it in the family: consanguineous marriage and genetic disorders, from Islamabad to Bradford. BMJ: British Medical Journal, 2019, 365, 11851.	2.3	9
108	Consanguinity and Inbreeding in Health and Disease in North African Populations. Annual Review of Genomics and Human Genetics, 2019, 20, 155-179.	6.2	45
109	Systematic literature review and meta-analysis on the epidemiology of methylmalonic acidemia (MMA) with a focus on MMA caused by methylmalonyl-CoA mutase (mut) deficiency. Orphanet Journal of Rare Diseases, 2019, 14, 84.	2.7	32
110	Spousal diabetes status as a risk factor for incident type 2 diabetes: a prospective cohort study and meta-analysis. Acta Diabetologica, 2019, 56, 619-629.	2.5	16

#	ARTICLE	IF	CITATIONS
111	A Syrian Refugee in Iraq Diagnosed as a Case of IL12RB1 Deficiency in Japan Using Dried Blood Spots. Frontiers in Immunology, 2019, 10, 58.	4.8	6
112	DRUG UTILIZATION ASSESSMENT IN PREGNANCY WOMEN: A CROSS SECTIONAL STUDY. International Research Journal of Pharmacy, 2019, 10, 109-113.	0.2	0
113	Preconception Pharmaceutical Care & Rare Diseases. , 2019, 05, .		0
114	Congenital anomalies and associated risk factors in a Saudi population: a cohort study from pregnancy to age 2 years. BMJ Open, 2019, 9, e026351.	1.9	17
115	Genetic peopling of Pakistan: Influence of consanguinity on population structure and forensic evaluation of traces. Forensic Science International: Genetics Supplement Series, 2019, 7, 232-233.	0.3	1
116	Inherited Bleeding Disorders in Women 2e. , 2019, , .		3
117	Molecular diagnostics of disorders of sexual development: an Indian survey and systems biology perspective. Systems Biology in Reproductive Medicine, 2019, 65, 105-120.	2.1	8
118	Prevalence of FVIII inhibitors in severe haemophilia A patients: Effect of treatment and genetic factors in an Indian population. Haemophilia, 2019, 25, 67-74.	2.1	9
119	Pathologies of matrix metalloproteinase-2 underactivity: a perspective on a neglected condition. Canadian Journal of Physiology and Pharmacology, 2019, 97, 486-492.	1.4	6
120	Overcoming bioethical, legal, and hereditary barriers to mitochondrial replacement therapy in the USA. Journal of Assisted Reproduction and Genetics, 2019, 36, 383-393.	2.5	22
121	A case series of hereditary cerebellar ataxias in a highly consanguineous population from Northeast Brazil. Parkinsonism and Related Disorders, 2019, 61, 193-197.	2.2	8
122	Effect of inbreeding on intellectual disability revisited by trio sequencing. Clinical Genetics, 2019, 95, 151-159.	2.0	49
123	User Acceptability of Whole Exome Reproductive Carrier Testing for Consanguineous Couples in Australia. Journal of Genetic Counseling, 2019, 28, 240-250.	1.6	5
124	Enhancing health literacy through co-design: development of culturally appropriate materials on genetic risk and customary consanguineous marriage. Primary Health Care Research and Development, 2019, 20, e2.	1.2	14
125	Genetics of intellectual disability in consanguineous families. Molecular Psychiatry, 2019, 24, 1027-1039.	7.9	131
126	Consanguinity and genetic diseases among the Bedouin population in the Negev. Journal of Community Genetics, 2020, 11, 13-19.	1.2	19
127	Exploring perceptions of consanguineous unions with women from an East London community: analysis of discussion groups. Journal of Community Genetics, 2020, 11, 225-234.	1.2	4
128	Emirati Women's Experiences of Consanguineous Marriage: a Qualitative Exploration of Attitudes, Health Challenges, and Coping Styles. International Journal of Mental Health and Addiction, 2020, 18, 1113-1127.	7.4	9

#	Article	IF	CITATIONS
129	Has the long-predicted decline in consanguineous marriage in India occurred?. Journal of Biosocial Science, 2020, 52, 746-755.	1.2	9
130	Effects of consanguinity in a cohort of subjects with certain genetic disorders in Qatar. Molecular Genetics & Samp; Genomic Medicine, 2020, 8, e1051.	1.2	31
131	Prevalence and pattern of consanguineous marriage among educated married individuals in Riyadh. Journal of Biosocial Science, 2020, 52, 768-775.	1.2	17
132	Clinical spectrum of non-syndromic microphthalmos, anophthalmos and coloboma in the paediatric population: a multicentric study from North India. British Journal of Ophthalmology, 2020, 105, bjophthalmol-2020-316910.	3.9	1
133	Presentation and management of N-acetylglutamate synthase deficiency: a review of the literature. Orphanet Journal of Rare Diseases, 2020, 15, 279.	2.7	5
134	Genetic Spectrum of Syndromic and Non-Syndromic Hearing Loss in Pakistani Families. Genes, 2020, 11, 1329.	2.4	7
135	Whole exome sequencing and homozygosity mapping reveals genetic defects in consanguineous Iranian families with inherited retinal dystrophies. Scientific Reports, 2020, 10, 19413.	3.3	9
136	Outcome associated with EPCAM founder mutation c.499dup in Qatar. European Journal of Medical Genetics, 2020, 63, 104023.	1.3	3
137	Consanguinity Marriage Increases Risk of Newborn's Congenital Anomalies in Sulaimani City. , 2020, , .		3
138	rs1542705–67,992,843-1,050,239 represents a novel informative haplotype at the SMPD1 locus in the Iranian population. Meta Gene, 2020, 25, 100744.	0.6	0
139	Investigation of the most common clinical and imaging findings and the role of tubulin genes in the etiology of malformations of cortical development. Turkish Journal of Medical Sciences, 2020, 50, 1573-1579.	0.9	0
140	The structure of first-cousin marriages in Brazil. Scientific Reports, 2020, 10, 15573.	3.3	5
141	Etiology of intellectual disability in individuals from special education schools in the south of Brazil. BMC Pediatrics, 2020, 20, 506.	1.7	3
142	Expanded carrier screening in reproductive medicine. , 2020, , 223-239.		0
143	Global perspectives on primary immune deficiency diseases., 2020,, 1129-1142.		0
144	Profound intellectual disability caused by homozygous TRAPPC9 pathogenic variant in a man from Malta. Molecular Genetics & Enomic Medicine, 2020, 8, e1211.	1.2	20
145	When to think outside the autozygome: Best practices for exome sequencing in "consanguineous― families. Clinical Genetics, 2020, 97, 835-843.	2.0	11
147	Clinical Reasoning: Complex ataxia. Neurology, 2020, 95, 136-141.	1.1	1

#	Article	IF	Citations
148	Worldwide carrier frequency and genetic prevalence of autosomal recessive inherited retinal diseases. Proceedings of the National Academy of Sciences of the United States of America, 2020, 117, 2710-2716.	7.1	195
149	Two decades of experience in a combined adult/pediatric allogeneic hematopoietic stem cell transplantation center in Algiers, Algeria. Annals of Hematology, 2020, 99, 619-625.	1.8	3
150	Consanguineous marriage and its effect on reproductive behavior and uptake of prenatal screening. Journal of Genetic Counseling, 2020, 29, 849-856.	1.6	5
151	Is consanguinity an impediment to child development?. Population Studies, 2020, 74, 139-159.	2.1	7
152	Consanguinity and Autism. Current Psychiatry Reports, 2020, 22, 3.	4.5	11
153	Signals Exchanged between Potential Spouses in the Saudi Arabian Arranged Marriage Market. Society, 2020, 57, 185-194.	1.2	0
154	Diabetes management in Wolcott-Rallison syndrome: analysis from the German/Austrian DPV database. Orphanet Journal of Rare Diseases, 2020, 15, 100.	2.7	8
155	Triadin Knockout Syndrome Is Absent in a Multi-Center Molecular Autopsy Cohort of Sudden Infant Death Syndrome and Sudden Unexplained Death in the Young and Is Extremely Rare in the General Population. Circulation Genomic and Precision Medicine, 2020, 13, e002731.	3.6	4
156	Consanguinity rates among Syrian refugees in Lebanon: a study on genetic awareness. Journal of Biosocial Science, 2021, 53, 356-366.	1.2	3
157	The experience of Preimplantation Genetic Testing (PGT) among Muslim couples in Oman in the Middle East. Journal of Genetic Counseling, 2021, 30, 121-131.	1.6	2
158	A large-scale study exploring understanding of the national premarital screening program among Jordanians: Is an at-risk marriage a valid option for Jordanians?. Public Understanding of Science, 2021, 30, 319-330.	2.8	3
159	Infant mortality in Turkey: Causes and effects in a regional context. Papers in Regional Science, 2021, 100, 429-454.	1.9	1
160	Distribution and management of the pediatric refugee population with renal replacement: A German pediatric cohort. Pediatric Nephrology, 2021, 36, 271-277.	1.7	9
161	Prevalence of Intimate Partner Violence Against Arab Women in Consanguineous Marriages., 2021,, 2483-2501.		0
162	The Effect of Consanguineous Marriages in Solving DNA Cases. , 2021, , 1-13.		0
163	Pitfalls in the diagnosis of Gaucher disease in Iraq:ÂAÂdiagnostic experience from a developing country. Pakistan Journal of Medical Sciences, 2021, 37, 782-787.	0.6	0
164	Designing a Simulation Model to Examine The Genetic Impact of Inbreeding In a Sudanese Population. , 2021, , .		0
165	Variant Analysis of Alkaptonuria Families with Significant Founder Effect in Jordan. BioMed Research International, 2021, 2021, 1-8.	1.9	3

#	Article	IF	CITATIONS
166	Screening of Obese Offspring of First-Cousin Consanguineous Subjects for the Angiotensin-Converting Enzyme Gene with a 287-bp Alu Sequence. Journal of Obesity and Metabolic Syndrome, 2021, 30, 63-71.	3.6	35
167	Diagnostic exome-based preconception carrier testing in consanguineous couples: results from the first 100 couples in clinical practice. Genetics in Medicine, 2021, 23, 1125-1136.	2.4	20
168	Changing Trends of Consanguineous Marriages in South India. Journal of Asian and African Studies, 2022, 57, 209-225.	1.5	3
169	Molecular analysis of CFTR gene mutations among Iraqi cystic fibrosis patients. Egyptian Journal of Medical Human Genetics, 2021, 22, .	1.0	3
170	Consanguinity: A Form of Social Capital Among the Muslims of Char Areas in Barpeta District, Assam. Journal of the Anthropological Survey of India, 2021, 70, 103-125.	0.6	0
171	Dental Anomalies in Consanguineous Marriage: A Clinical-Radiological Study. International Dental Journal, 2021, 72, 133-133.	2.6	4
172	Consanguineous marriage and rare bleeding disorders. Expert Review of Hematology, 2021, 14, 467-472.	2,2	6
173	Genetic Characterization of Short Stature Patients With Overlapping Features of Growth Hormone Insensitivity Syndromes. Journal of Clinical Endocrinology and Metabolism, 2021, 106, e4716-e4733.	3.6	11
174	The Effect of Internal Migration on the General and Reproductive Health Status of Women Migrating to Aydın Province. Hacettepe Üniversitesi Hemşirelik Fakültesi Dergisi, 2021, 8, 213-222.	0.8	3
175	High prevalence of multilocus pathogenic variation in neurodevelopmental disorders in the Turkish population. American Journal of Human Genetics, 2021, 108, 1981-2005.	6.2	38
176	Health Literacy From a Pediatrician's Perspective. , 2022, , 251-273.		2
177	Targeted genome editing for the correction or alleviation of primary Immunodeficiencies. Progress in Molecular Biology and Translational Science, 2021, 182, 111-151.	1.7	3
178	Incest. , 2021, , 4033-4036.		0
179	Consanguinity, awareness, and genetic disorders among female university students in Riyadh, Saudi Arabia. Journal of Biochemical and Clinical Genetics, 0, , 27-34.	0.1	4
180	Clinical Management of NF1 and Indications for Surgery. , 2021, , 347-354.		0
183	Health Research, Regulations and Ethics in the United Arab Emirates. Research Ethics Forum, 2017, , 255-266.	0.1	3
184	What should be the focus of counseling in parental consanguinity: genetic disorders or underlying beliefs. Egyptian Journal of Medical Human Genetics, 2020, 21, .	1.0	4
185	Family history of cardiovascular disease and risk of premature coronary heart disease: A matched case-control study. Wellcome Open Research, 2020, 5, 70.	1.8	12

#	Article	IF	CITATIONS
186	A perinatal review of singleton stillbirths in an Australian metropolitan tertiary centre. PLoS ONE, 2017, 12, e0171829.	2.5	4
187	Adolescents' opinions and thoughts about consanguineous marriages: A Turkey sample. Journal of Human Sciences, 2017, 14, 105.	0.2	4
188	Genetic pattern of cystic fibrosis patients in Azeri Turkish population. Russian Open Medical Journal, 2017, 6, e0101.	0.3	3
189	Epidemiological Study of Legal Abortion due to Fetal Defects in the Files Referred to Fars Province Forensic Medicine Centers from 2007 to 2013. Shiraz E Medical Journal, 2016, 17, .	0.3	5
190	Knowledge and Health Beliefs Regarding Sickle Cell Disease Among Omanis in a Primary Healthcare Setting: Cross-sectional study. Sultan Qaboos University Medical Journal, 2016, 16, e437-444.	1.0	10
191	Public Awareness of the Consanguinity and Genetic Disease Prevalence Connection in Saudi Arabia. Asian Journal of Pharmaceutical Research and Health Care, 2020, 12, 213-222.	0.1	2
192	Homozygosity Mapping and Targeted Sanger Sequencing Identifies Three Novel CRB1 (Cumbs) Tj ETQq0 0 0 rgBT 294-302.		10 Tf 50 50 7
193	Need for a universal thalassemia screening programme in India? A public health perspective. Journal of Family Medicine and Primary Care, 2019, 8, 1528.	0.9	9
194	Severe Disseminated Phaeohyphomycosis in a Patient with Inherited CARD9 Deficiency. Archives of Clinical Infectious Diseases, $2018,13,\ldots$	0.2	7
195	Association Between Outcomes and Demographic Factors in an Azeri Turkish Population With Cystic Fibrosis: A Cross-Sectional Study in Iran From 2001 Through 2014. Iranian Red Crescent Medical Journal, 2016, 18, e29615.	0.5	10
196	Knowledge, awareness, and attitude of medical students concerning genetics and premarital screening. Journal of Nature and Science of Medicine, 2021, 4, 356.	0.3	1
197	A case-control study on factor V Leiden: an independent, gender-dependent risk factor for venous thromboembolism. Thrombosis Journal, 2021, 19, 74.	2.1	2
198	Social determinants of health and primary immunodeficiency. Annals of Allergy, Asthma and Immunology, 2022, 128, 12-18.	1.0	10
200	Analysis of consanguineous and non-consanguineous pedigrees related to the mode of transmission of essential hypertension- study based on Serene Threonine Kinase-39 gene Single Nucleotide Polymorphism rs35929607 in a population of Sindh, Pakistan. International Journal of Medical Science and Public Health, 2014, 3, 955.	0.2	O
201	Reclaiming Oromo Indigenous Organizational Structures and Fostering Supportive Environments for Health. Archives of Business Research, 2014, 2, 23-45.	0.1	0
202	Consanguinity – Its Impact, Consequences and Management. , 2014, , .		O
203	Variations in Types of First-Cousin Marriages over a Two-Generation Period among Arabs in Israel. Advances in Anthropology, 2015, 05, 171-176.	0.2	2
205	CONSANGUINITY AND ITS EFFECT ON OFFSPRING IN AN URBAN COMMUNITY OF GUNTUR CITY OF ANDHRA PRADESH. Journal of Evidence Based Medicine and Healthcare, 2017, 4, 4221-4225.	0.0	O

#	Article	IF	CITATIONS
206	Premarital Health and Social Issues … Suffering in Silence?. Gynecology & Obstetrics (Sunnyvale, Calif) Tj ETQqC	0 9 9 rgBT	/Qverlock 10
207	Study of Chromosomal Abnormalities in Couples with Recurrent Spontaneous Abortions (RSA) in Ardabil Province. Journal of Ardabil University of Medical Sciences, 2018, 18, 91-107.	0.2	0
208	Consanguinité, Isonymie et Age Précoce au Mariage dans les deux Provinces de Tétouan et M'diq-fnide (Maroc). European Scientific Journal, 2018, 14, 63.	²⁹ 0.1	0
209	Incest. , 2020, , 1-4.		3
210	Family history of cardiovascular disease and risk of premature coronary heart disease: A matched case-control study. Wellcome Open Research, 2020, 5, 70.	1.8	5
211	Consanguineous marriage as socio-medical problem. I P Pavlov Russian Medical Biological Herald, 2020, 28, 249-258.	0.5	0
212	Interactions of Consanguinity and Number of Siblings with Childhood Acute Lymphoblastic Leukemia. BioMed Research International, 2020, 2020, 1-11.	1.9	2
213	Prevalence of Intimate Partner Violence Against Arab Women in Consanguineous Marriages. , 2020, , 1-19.		0
214	Genetic Health Care Before Conception. , 2020, , 35-52.		1
215	Health Literacy From a Pediatrician's Perspective. Advances in Media, Entertainment and the Arts, 2020, , 308-330.	0.1	0
216	IDENTIFICATION OF MUTATIONS IN THE PAH GENE IN PKU PATIENTS IN THE STATE OF MATO GROSSO. Revista Paulista De Pediatria, 2020, 38, e2018351.	1.0	3
217	Implantable and Wearable Neuroengineering Education: A Review of Postgraduate Programmes. IEEE Access, 2020, 8, 212396-212408.	4.2	4
218	Prevalence of Coagulation Factors Deficiency among Young Adults in Saudi Arabia: A National Survey. TH Open, 2020, 04, e457-e462.	1.4	5
219	La endogamia como causa de consanguinidad y su asociación con anomalÃas congénitas. Medicina Y Laboratorio, 2020, 25, 409-418.	0.1	1
220	Consanguineous marriages in the genetic counseling centers of Isfahan and the ethical issues of clinical consultations. Journal of Medical Ethics and History of Medicine, 2017, 10, 12.	0.6	5
221	Consanguineous Marriage as a Risk Factor for Developing Keratoconus. Medical Hypothesis, Discovery, and Innovation in Ophthalmology, 2018, 7, 17-21.	0.2	4
222	Targeted next generation sequencing reveals genetic defects underlying inherited retinal disease in Iranian families. Molecular Vision, 2019, 25, 106-117.	1.1	10
223	Induced Pluripotent Stem Cells in Pediatric Research and Clinical Translation. , 2021, , 203-216.		5

#	Article	IF	CITATIONS
224	Trends of Hospital Admissions Due to Congenital Anomalies in England and Wales between 1999 and 2019: An Ecological Study. International Journal of Environmental Research and Public Health, 2021, 18, 11808.	2.6	17
225	Quantitative dissection of multilocus pathogenic variation in an Egyptian infant with severe neurodevelopmental disorder resulting from multiple molecular diagnoses. American Journal of Medical Genetics, Part A, 2022, 188, 735-750.	1.2	14
226	Novel mutations in PDE6A and CDHR1 cause retinitis pigmentosa in Pakistani families. International Journal of Ophthalmology, 2021, 14, 1843-1851.	1.1	1
227	Custom Nextâ€Generation Sequencing Identifies Novel Mutations Expanding the Molecular and clinical spectrum of isolated Hearing Impairment or along with defects of the retina, the thyroid, and the kidneys. Molecular Genetics & Denomic Medicine, 2022, 10, e1868.	1.2	2
228	LEGAL REGULATION OF THE "BIOLOGICAL―PROTECTION OF MARRIAGE AND FAMILY: ISSUES OF BALANCING BETWEEN THE RIGHT TO MEDICAL SECRET AND THE RIGHT TO CONSCIOUS MARRIAGE. WiadomoÅ⋅ci Lekarskie, 2021, 74, 3009-3015.		0
229	Newborn Hearing Screening Benefits Children, but Global Disparities Persist. Journal of Clinical Medicine, 2022, 11, 271.	2.4	17
231	Fine-Scale Genetic Structure in the United Arab Emirates Reflects Endogamous and Consanguineous Culture, Population History, and Geography. Molecular Biology and Evolution, 2022, 39, .	8.9	3
232	Importance of multigene panel test in patients with consanguineous marriage and family history of breast cancer. Oncology Letters, 2022, 23, 118.	1.8	0
233	Consanguinity and Congenital Heart Disease Susceptibility: Insights into Rare Genetic Variations in Saudi Arabia. Genes, 2022, 13, 354.	2.4	7
234	Arginase 1 Deficiency: using genetic databases as a tool to establish global prevalence. Orphanet Journal of Rare Diseases, 2022, 17, 94.	2.7	7
235	Progress, challenges, and future perspectives in genetic researches of stuttering. Journal of Genetic Medicine, 2021, 18, 75-82.	0.2	0
236	Consanguinity in Risk Assessment of Retinoblastoma Using Machine Learning. Lecture Notes in Electrical Engineering, 2022, , 579-585.	0.4	2
237	Association of maternal age and presence of non-communicable diseases in consanguineous marriage with congenital abnormalities in infants. , 0, , 24-27.		0
238	The Effect of Consanguineous Marriages in Solving DNA Cases. , 2022, , 545-557.		O
239	Les mariages consanguins et leurs effets sur les maladies non transmissibles dans la population Marocaine: étude transversale. Pan African Medical Journal, 0, 41, .	0.8	1
240	Knowledge and attitude of pregnant women in the Kingdom of Saudi Arabia toward Noninvasive prenatal testing: A single center study. Molecular Genetics & Samp; Genomic Medicine, 2022, , e1960.	1.2	1
241	Primary hyperoxaluria and genetic linkages: an insight into the disease burden from Pakistan. Urolithiasis, 2022, 50, 439-445.	2.0	3
242	Redislocation After Posteromedial Open Reduction in Developmental Dysplasia of the Hip: Analyzing the Risk Factors and Determining the Optimal Treatment Method. Journal of Pediatric Orthopaedics, 0, Publish Ahead of Print, .	1.2	2

#	Article	IF	Citations
243	Models of Love as Social Connections. , 2022, , 249-304.		0
244	Effect of Women's Status on Consanguinity in the Arab Society of Israel. Advances in Anthropology, 2022, 12, 137-148.	0.2	1
245	Case report: First case report of an Emirati child with a novel gene variant causing aromatic L-amino acid decarboxylase deficiency. Frontiers in Pediatrics, $0,10,10$	1.9	2
246	Knowledge regarding consanguineous marriage and it effects on pregnancy outcome among the adolescents of village in Bagalakote district. Indian Journal of Forensic and Community Medicine, 2022, 9, 117-123.	0.3	0
247	Whole-exome sequencing analysis in a case of primary congenital glaucoma due to the partial uniparental isodisomy. Genomics and Informatics, 2022, 20, e28.	0.8	2
248	Heterozygosity reduction in children whose parents are closely related. Forensic Science International: Genetics Supplement Series, 2022, 8, 71-73.	0.3	1
249	Education and Consanguineous Marriage. Journal of Human Capital, 2023, 17, 114-171.	1.3	3
250	Investigating the genetic profile of familial atypical cystic fibrosis patients (DeltaF508-CFTR) with neonatal biliary atresia. Journal of Applied Genetics, 2023, 64, 71-80.	1.9	2
251	Cleft lip and palate based on birth order and family history at Mitra Sejati General Hospital, Indonesia. Dental Journal: Majalah Kedokteran Gigi, 2022, 55, 221-225.	0.2	1
252	Parental occupational and environmental risk factors for childhood bone cancer in Mansoura oncology center: a case control study. International Journal of Environmental Health Research, 2024, 34, 248-256.	2.7	0
254	An Updated Review on MSMD Research Globally and A Literature Review on the Molecular Findings, Clinical Manifestations, and Treatment Approaches in China. Frontiers in Immunology, 0, 13, .	4.8	7
255	CASE REPORT: MUCOPOLYSACCHRIDOSIS TYPE IV - MORQUIO SYNDROME. , 2022, , 62-63.		0
256	Hereditary thrombotic thrombocytopenic purpura (TTP) with co-occurring autosomal dominant polycystic kidney disease (ADPKD). BMJ Case Reports, 2022, 15, e250378.	0.5	0
257	Multidisciplinary management of Papillon-Lefevre syndrome as a result of consanguineous marriage. BMJ Case Reports, 2022, 15, e252992.	0.5	0
258	Consanguinity and childhood acute lymphoblastic leukaemia: a case-control study. The Gazette of the Egyptian Paediatric Association, 2022, 70, .	0.4	0
259	Characterization of Fertility Clinic Attendees in the Abu Dhabi Emirate, United Arab Emirates: A Cross-Sectional Study. International Journal of Environmental Research and Public Health, 2023, 20, 1692.	2.6	4
260	Relative frequency of inherited retinal disorders in Khuzestan province, southern Iran. Journal of Advanced Pharmacy Education and Research, 2023, 13, 42-45.	1.1	0
262	Comparison of energy and nutrient intakes of individuals with disabilities according to reference values: A cross-sectional study. Ankara Sağlık Bilimleri Dergisi, 2022, 11, 1-13.	0.3	0

#	Article	IF	CITATIONS
263	Health inequity in genomic personalized medicine in underrepresented populations: a look at the current evidence. Functional and Integrative Genomics, 2023, 23, .	3.5	2
264	Analysis of perinatal outcomes of pregnancies from consanguineous marriages in a tertiary hospital in Bursa, Turkey. Journal of Health Sciences and Medicine, 2023, 6, 359-363.	0.1	0
266	Delineating the Spectrum of Genetic Variants Associated with Bardet-Biedl Syndrome in Consanguineous Pakistani Pedigrees. Genes, 2023, 14, 404.	2.4	5
267	Prevalence of Consanguineous Marriage among Saudi Citizens of Albaha, a Cross-Sectional Study. International Journal of Environmental Research and Public Health, 2023, 20, 3767.	2.6	6
268	The added value of brain MR spectroscopy in children with Crigler-Najjar syndrome type-I: correlation with demographic, neurodevelopmental, and laboratory findings. British Journal of Radiology, 2023, 96, .	2.2	0
269	Rare manifestations of alobar holoprosencephaly and the potential causes: a report of two cases. Annals of Medicine and Surgery, 2023, 85, 252-256.	1.1	0
270	Consanguineous marriage as a key indicator of isolated congenital dental anomaly among South Indian population – A cross-sectional study. Journal of Oral and Maxillofacial Pathology, 2023, 27, 60.	0.6	0
271	Incidence of Inborn Errors of Metabolism in Newborn Infants: Five Years' Single-Center Experience, Jeddah, Saudi Arabia. Clinical Pediatrics, 0, , 000992282311635.	0.8	0
272	Construction and validation of ShaMaq developmental screening tool. Child: Care, Health and Development, 2024, 50, .	1.7	0
273	Features of chromosomal abnormalities in relation to consanguinity: analysis of 10,556 blastocysts from IVF/ICSI cycles with PGT-A from consanguineous and non-consanguineous couples. Scientific Reports, 2023, 13, .	3.3	2
275	Consanguineous couples $\hat{\mathbf{e}}^{\text{TM}}$ experiences and views regarding expanded carrier screening: Barriers and facilitators in the decision-making process. European Journal of Human Genetics, 0, , .	2.8	1
276	Consanguinity., 2023, , 55-63.		0
277	Familial Cataracts: Profile of Patients and Their Families at a Child Eye Care Tertiary Facility in a Developing Country. Korean Journal of Ophthalmology: KJO, 0, , .	1.1	0
278	Predictors of Consanguinity Marriage Decision in Saudi Arabia: A Pilot Study. Healthcare (Switzerland), 2023, 11, 1925.	2.0	1
279	Living Apart Together (LAT): A New Family Form in Urban India. Sociological Bulletin, 0, , .	0.4	0
280	Thalassemia, biobanking infrastructures, and personalized stem cell therapies in Chennai. Frontiers in Sociology, 0, 8, .	2.0	0
281	Perception of consanguineous marriage among the qatari population. Frontiers in Public Health, 0, 11 , .	2.7	1
282	Homogentisate 1,2-dioxygenase (HGD) gene variants in young Egyptian patients with alkaptonuria. Scientific Reports, 2023, 13, .	3.3	0

#	Article	IF	CITATIONS
283	Influence of autozygosity on common disease risk across the phenotypic spectrum. Cell, 2023, 186, 4514-4527.e14.	28.9	2
284	Consanguinity and willingness to perform premarital genetic screening in Sudan. European Journal of Human Genetics, 0, , .	2.8	1
285	Reporting a novel growth hormone receptor gene variant in an Iranian consanguineous pedigree with Laron syndrome: a case report. BMC Endocrine Disorders, 2023, 23, .	2.2	1
286	Induzierte pluripotente Stammzellen in der pÄ d iatrischen Forschung und klinischen Umsetzung. , 2023, , 225-240.		0
288	Chromosomal aberration detection in Iraqi children with autism., 2023, 38, 201229.		0
289	Clinical and Genetic Characterization of Patients with Primary Ciliary Dyskinesia in Southwest Saudi Arabia: A Cross Sectional Study. Children, 2023, 10, 1684.	1.5	0
290	Genetic Testing for Parkinson's Disease and Movement Disorders in Less Privileged Areas: Barriers and Opportunities. Movement Disorders Clinical Practice, 2024, 11, 14-20.	1.5	0
291	South Asia: The Missing Diverse in Diversity. Behavior Genetics, 0, , .	2.1	0
292	Genetic literacy and experiential knowledge on sickle cell disease among Canadian- and foreign-born male and female Anglophone and Francophone youth in Canada. International Journal of Adolescent Medicine and Health, 2023, 35, 443-455.	1.3	0
293	Assessment of Perceptions and Predictors Towards Consanguinity: A Cross-Sectional Study from Palestine. Journal of Multidisciplinary Healthcare, 0, Volume 16, 3443-3453.	2.7	0
294	Determinants of childbearing intentions among pregnant women with a suspected fetal congenital heart disease. Midwifery, 2024, 128, 103875.	2.3	0
295	The Impact of Increased Homozygosity on Human Fertility: A Comprehensive Review. Cureus, 2023, , .	0.5	0
296	Direct and indirect predictors of postpartum depression symptoms among indigenous Bedouin mothers in Israel. Research in Nursing and Health, 0, , .	1.6	1
298	Î'lpha-thalassemia: A practical overview. Blood Reviews, 2024, 64, 101165.	5.7	1
299	Title-molecular diagnostics of dystrophinopathies in Sri Lanka towards phenotype predictions: an insight from a South Asian resource limited setting. European Journal of Medical Research, 2024, 29, .	2.2	0
300	Search for chromosomal instability aiding variants reveal naturally occurring kinetochore gene variants that perturb chromosome segregation. IScience, 2024, 27, 109007.	4.1	0
301	Primary sex ratio in euploid embryos of consanguine couples after IVF/ICSI. Journal of Assisted Reproduction and Genetics, 2024, 41, 957-965.	2.5	0
302	Mechanical Hatching as a Therapeutic Intervention for Improving Implantation Rate in a 32-Year-Old Female With Recurrent Implantation Failures: A Case Report. Cureus, 2024, , .	0.5	0

#	Article	IF	CITATIONS
303	Genetic, immunologic, and clinical features of 830 patients with Mendelian susceptibility to mycobacterial diseases (MSMD): AÂsystematic review. Journal of Allergy and Clinical Immunology, 2024, 153, 1432-1444.	2.9	0
304	Consanguineous Marriage and Its Association With Genetic Disorders in Saudi Arabia: A Review. Cureus, 2024, , .	0.5	0
305	Investigating the genetic makeup of the major histocompatibility complex (MHC) in the United Arab Emirates population through next-generation sequencing. Scientific Reports, 2024, 14, .	3.3	0
306	A Rare Presentation of Laurence-Moon-Bardet-Biedl Syndrome: Atypical Retinitis Punctata Albescens and Non-alcoholic Fatty Liver Disease. Cureus, 2024, , .	0.5	0
307	Global carrier frequency and predicted genetic prevalence of patients with pathogenic sequence variants in autosomal recessive genetic neuromuscular diseases. Scientific Reports, 2024, 14, .	3.3	0
308	Reproductive Carrier Screening., 2023,, 235-250.		0
309	Assessing Students' Knowledge and Attitudes Regarding the Risks and Prevention of Consanguineous Marriage: A Cross-Sectional Online Survey. Journal of Multidisciplinary Healthcare, 0, Volume 17, 1251-1263.	2.7	0