

CITATION REPORT

List of articles citing

Consanguineous marriage in Iran

DOI: 10.1080/03014460310001652211

Annals of Human Biology, 2004, 31, 263-9.

Source: <https://exaly.com/paper-pdf/37455996/citation-report.pdf>

Version: 2024-04-26

This report has been generated based on the citations recorded by exaly.com for the above article. For the latest version of this publication list, visit the link given above.

The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

#	Paper	IF	Citations
245	ATM haplotypes and associated mutations in Iranian patients with ataxia-telangiectasia: recurring homozygosity without a founder haplotype. <i>Human Genetics</i> , 2005 , 117, 101-6	6.3	19
244	Epidemiology and mortality of hospitalized burn patients in Kohkiluyeh va Boyer-Ahmad province (Iran): 2002-2004. 2005 , 31, 306-9		80
243	Correlation between incidences of self-inflicted burns and means of inbreeding coefficients, an ecologic study. 2006 , 16, 708-11		26
242	Consanguinity in primary immunodeficiency disorders; the report from Iranian Primary Immunodeficiency Registry. 2006 , 56, 145-51		76
241	Consanguineous unions and child health in the State of Qatar. 2006 , 20, 372-8		80
240	Is consanguineous marriage religiously encouraged? Islamic and Iranian considerations. 2007 , 39, 313-6		48
239	Rising Cousins 2007 , 48, 756-764		13
238	Consanguineous marriages in Iranian folktales. 2007 , 10, 38-40		26
237	Consanguineous marriages and their effects on common adult diseases: studies from an endogamous population. 2007 , 16, 262-7		72
236	Ataxia-telangiectasia in Iran: clinical and laboratory features of 104 patients. 2007 , 37, 21-8		54
235	Distribution of primary immunodeficiency diseases in the Turk ethnic group, living in the northwestern Iran. 2007 , 27, 510-6		23
234	Genetic polymorphism of GSTT1 may be under natural selection in a population chronically exposed to natural sour gas. 2008 , 35, 673-6		8
233	Genetic counseling in southern Iran: consanguinity and reason for referral. 2008 , 17, 472-9		16
232	Common MEFV mutation analysis in Iranian Azeri Turkish patients with familial Mediterranean fever. 2008 , 37, 334-8		24
231	Prevalence and sociodemographic correlates of consanguineous marriages in Turkey. 2008 , 40, 137-48		62
230	Modernization or cultural maintenance: the practice of consanguineous marriage in Iran. 2008 , 40, 911-33		40
229	A community genetics perspective on consanguineous marriage. 2008 , 11, 324-30		37

228	Molecular analysis of factor IX gene in an Iranian female with severe hemophilia B. 2008 , 119, 151-3	4
227	Basic Concepts in Statistics and Epidemiology. 2008 , 62, 567-568	
226	Consanguinity and national IQ scores. 2008 , 62, 566-7	13
225	The Fertility Transition in Iran: Revolution and Reproduction. 2009 ,	81
224	Prevalence of consanguineous marriages in Syria. 2009 , 41, 685-92	55
223	Genetic counseling for thalassemia in the Islamic Republic of Iran. 2009 , 52, 364-76	14
222	Is there a significant trend in prevalence of consanguineous marriage in Tehran? A review of three generations. 2009 , 18, 82-6	25
221	Family Change and Continuity in Iran: Birth Control Use Before First Pregnancy. 2009 , 71, 1309-1324	30
220	Fragile X syndrome screening of families with consanguineous and non-consanguineous parents in the Iranian population. 2009 , 52, 170-3	11
219	Consanguinity and family clustering of male factor infertility in Lebanon. 2009 , 91, 1104-9	23
218	Prevalence of classical phenylketonuria in mentally retarded individuals in Iran. 2009 , 32 Suppl 1, S283-7	7
217	Common MEFV mutation analysis in 36 Iranian patients with familial Mediterranean fever: clinical and demographic significance. 2010 , 20, 566-572	8
216	Are radiologists and radiological technologists at greater risk of reproductive health problems? Data from seven provinces in Iran. 2010 , 7, 167	0
215	Correlation between consanguineous marriages and age-standardized mortality rate due to breast cancer, an ecologic study. 2010 , 121, 795-7	4
214	Effect of inbreeding on weight gain of offspring from birth to 12 months after birth: a study from Iran. 2010 , 42, 195-200	10
213	Bleeding disorders in the tribe: result of consanguineous in breeding. 2010 , 5, 23	34
212	Consanguinity, Genetic Drift, and Genetic Diseases in Populations with Reduced Numbers of Founders. 2010 , 507-528	15
211	Parental consanguinity and susceptibility to drug abuse among offspring, a case-control study. 2010 , 180, 57-9	8

210	Prevalence and causes of severe visual impairment and blindness among children in the Lorestan province of Iran, using the key informant method. 2010 , 17, 95-102	29
209	Parental consanguineous marriages and age at onset of schizophrenia. 2011 , 126, 298-9	11
208	Impact of consanguineous marriages in GJB2-related hearing loss in the Iranian population: a report of a novel variant. 2011 , 15, 489-93	26
207	Chromosome abnormality rate among Iranian patients with idiopathic mental retardation from consanguineous marriages. 2011 , 7, 321-5	5
206	Increased availability of family donors for hematopoietic stem cell transplantation in a population with increased incidence of consanguinity. 2011 , 25, 475-80	8
205	The demographics of primary immunodeficiency diseases across the unique ethnic groups in Iran, and approaches to diagnosis and treatment. 2011 , 1238, 24-32	14
204	Distribution and clinical features of primary immunodeficiency diseases in Chinese children (2004-2009). 2011 , 31, 297-308	20
203	Inheritance pattern and clinical aspects of 93 Iranian patients with chronic granulomatous disease. 2011 , 31, 792-801	73
202	Association between consanguineous marriages and risk of pre-eclampsia. 2011 , 283 Suppl 1, 5-7	9
201	Parental consanguinity and the risk of primary immunodeficiency disorders: report from the Kuwait National Primary Immunodeficiency Disorders Registry. 2011 , 154, 76-80	27
200	Prevalence of consanguineous marriages among Iranian Georgians. 2011 , 43, 47-50	13
199	IL-12R β 1 deficiency in two of fifty children with severe tuberculosis from Iran, Morocco, and Turkey. 2011 , 6, e18524	91
198	Association between healthy life expectancy at birth and consanguineous marriages in 63 countries. 2011 , 43, 475-80	11
197	Mutation analysis of the CYP21A2 gene in the Iranian population. 2012 , 16, 82-90	19
196	Consanguineous marriages in Afghanistan. 2012 , 44, 73-81	18
195	Influence of parental consanguineous marriages on age at onset of bipolar disorder. 2012 , 198, 327-8	7
194	Spectrum of GJB2 (Cx26) gene mutations in Iranian Azeri patients with nonsyndromic autosomal recessive hearing loss. 2012 , 76, 268-71	21
193	A comprehensive study to determine heterogeneity of autosomal recessive nonsyndromic hearing loss in Iran. 2012 , 158A, 2485-92	47

192	Prenatal genetic counseling and consanguinity. 2012 , 32, 1133-8		9
191	References. 232-314		
190	A transversion mutation in non-coding exon 3 of the TMC1 gene in two ethnically related Iranian deaf families from different geographical regions; evidence for founder effect. 2013 , 77, 821-6		14
189	Molecular characterization of WFS1 in an Iranian family with Wolfram syndrome reveals a novel frameshift mutation associated with early symptoms. 2013 , 528, 309-13		9
188	Consanguinity, endogamy, and genetic disorders in Tunisia. 2013 , 4, 273-84		57
187	Rare bleeding disorders: worldwide efforts for classification, diagnosis, and management. <i>Seminars in Thrombosis and Hemostasis</i> , 2013 , 39, 579-84	5-3	43
186	Prevalence of consanguineous marriages in west and south of Afghanistan. 2013 , 45, 799-805		15
185	Prevalence of consanguineous marriages among shi'a populations of Lebanon. 2013 , 45, 675-82		10
184	The prevalence of keratoconus in a young population in Mashhad, Iran. 2014 , 34, 519-27		58
183	Clinical and genetic study on 356 Brazilian patients with a distinct phenotype of cleft lip and palate without alveolar ridge involvement. 2014 , 42, 1952-7		4
182	Color vision deficiency in Zahedan, Iran: lower than expected. 2014 , 91, 1372-6		2
181	CIVIL UNREST AND THE CURRENT PROFILE OF CONSANGUINEOUS MARRIAGE IN KHYBER PAKHTUNKHWA PROVINCE, PAKISTAN. 2014 , 46, 698-701		11
180	Investigation of ATP6V1B1 and ATP6V0A4 genes causing hereditary hearing loss associated with distal renal tubular acidosis in Iranian families. 2014 , 128, 1056-9		3
179	Consanguineous marriage, reproductive behaviour and postnatal mortality in contemporary Iran. 2014 , 77, 16-25		24
178	Oral abnormalities in an Iranian newborn population. 2014 , 24, 8-11		
177	Parental consanguinity and nonsyndromic orofacial clefts in children: a systematic review and meta-analyses. 2014 , 51, 501-13		11
176	Genetics of consanguineous marriage: Impact and importance of counseling. 2012 , 1, 217-20		1
175	Rare bleeding disorders: diagnosis and treatment. 2015 , 125, 2052-61		180

174	Family History and Relapse in Pediatric Acute Myeloid Leukemia. 2015 , 62, 2235-7	2
173	Consanguineous marriage, prepregnancy maternal characteristics and stillbirth risk: a population-based case-control study. 2015 , 94, 1095-101	15
172	Risk factors for keratoconus in Israel: a case-control study. 2015 , 35, 673-81	37
171	Age-standardized Incidence Rates for Leukemia Associated with Consanguineous Marriages in 68 Countries, an Ecological Study. 2015 , 7, e2015027	1
170	The genetic and environmental factors for keratoconus. <i>BioMed Research International</i> , 2015 , 2015, 795738	169
169	Incidence of primary breast cancer in Iran: Ten-year national cancer registry data report. 2015 , 39, 519-27	91
168	Association between consanguinity and survival of marriages. <i>Egyptian Journal of Medical Human Genetics</i> , 2015 , 16, 67-70	2 6
167	A study of consanguineous marriage as a risk factor for developing comitant strabismus. 2015 , 6, 177-80	5
166	Genotype and phenotype correlations in Iranian patients with hyperinsulinaemic hypoglycaemia. 2015 , 8, 350	6
165	Characterising the spectrum of autosomal recessive hereditary hearing loss in Iran. 2015 , 52, 823-9	72
164	Novel missense mutation in the GALNS gene in an affected patient with severe form of mucopolysaccharidosis type IVA. 2015 , 450, 121-4	2
163	Allelic heterogeneity among Iranian DFNB7/11 families: report of a new Iranian deaf family with TMC1 mutation identified by next-generation sequencing. 2015 , 135, 125-9	6
162	Health of children born through artificial oocyte activation: a pilot study. 2015 , 22, 322-8	32
161	The Effect of Consanguineous Marriage on Mental Health among the Students of the Shahrekord University of Medical Sciences. 2016 , 10, GC01-GC04	5
160	Consanguinity Among Parents of Iranian Deaf Children. 2016 , 18, e22038	6
159	Significant association of susceptibility to schizophrenia and type I bipolar disorder with parental consanguineous marriages. 2016 , 5, 37-40	
158	Associations between the codon 72 polymorphism of the TP53 gene and the risk of recurrent implantation failure. 2016 , 42, 184-9	3
157	DOCK8 deficiency in six Iranian patients. 2016 , 4, 593-600	3

156	Association between Family History and Keratoconus Severity. 2016 , 41, 1414-1418	17
155	A novel mutation in alpha sarcoglycan gene in an Iranian family with limb girdle muscular dystrophy 2D. 2016 , 38, 220-3	2
154	Consanguinity and multiple sclerosis susceptibility: A case control study. 2016 , 10, 179-180	6
153	Co-existence of phenylketonuria either with maple syrup urine disease or Sandhoff disease in two patients from Iran: emphasizing the role of consanguinity. 2016 , 29, 1215-1219	5
152	Mate Preferences in Young Iranian Women: Cultural and Individual Difference Correlates. 2016 , 2, 247-253	13
151	The genetic background of Southern Iranian couples before marriage. 2016 , 19, 71-74	1
150	Gene mutations responsible for primary immunodeficiency disorders: A report from the first primary immunodeficiency biobank in Iran. 2016 , 12, 62	7
149	A rare form of limb girdle muscular dystrophy (type 2E) seen in an Iranian family detected by autozygosity mapping. 2016 , 30, 1-4	2
148	Homozygosity mapping in albinism patients using a novel panel of 13 STR markers inside the nonsyndromic OCA genes: introducing 5 novel mutations. 2016 , 61, 373-9	5
147	Identification of six novel mutations in Iranian patients with maple syrup urine disease and their in silico analysis. 2016 , 786, 34-40	9
146	The relations between marital quality, social support, social acceptance and coping strategies among the infertile Iranian couples. 2016 , 200, 58-62	11
145	Clinical, genetic, and structural basis of congenital adrenal hyperplasia due to 11 β hydroxylase deficiency. 2017 , 114, E1933-E1940	70
144	Paecilomyces formosus Infection in an Adult Patient with Undiagnosed Chronic Granulomatous Disease. 2017 , 37, 342-346	8
143	A novel pathogenic variant in the FZD6 gene causes recessive nail dysplasia in a large Iranian kindred. 2017 , 88, 134-138	1
142	The association between parental consanguinity and primary immunodeficiency diseases: A systematic review and meta-analysis. 2017 , 28, 280-287	11
141	A novel pathogenic variant in an Iranian Ataxia telangiectasia family revealed by next-generation sequencing followed by in silico analysis. 2017 , 379, 212-216	9
140	An update of common autosomal recessive non-syndromic hearing loss genes in Iranian population. 2017 , 97, 113-126	18
139	Measurement and associative factors of adherence to epilepsy drug treatment among the elderly population in Tehran (Iran). 2017 , 04, 046-050	

138	The spectrum of Familial Mediterranean Fever gene (MEFV) mutations and genotypes in Iran, and report of a novel missense variant (R204H). 2017 , 60, 701-705	6
137	Multigene Next-Generation Sequencing Panel Identifies Pathogenic Variants in Patients with Unknown Subtype of Epidermolysis Bullosa: Subclassification with Prognostic Implications. 2017 , 137, 2649-2652	23
136	Autosomal recessive congenital ichthyosis: CERS3 mutations identified by a next generation sequencing panel targeting ichthyosis genes. 2017 , 25, 1282-1285	14
135	The Genetic Spectrum of Familial Hypercholesterolemia (FH) in the Iranian Population. 2017 , 7, 17087	13
134	Dystrophic Epidermolysis Bullosa: COL7A1 Mutation Landscape in a Multi-Ethnic Cohort of 152 Extended Families with High Degree of Customary Consanguineous Marriages. 2017 , 137, 660-669	34
133	Exome sequencing of Pakistani consanguineous families identifies 30 novel candidate genes for recessive intellectual disability. 2017 , 22, 1604-1614	69
132	In silico analysis of novel mutations in maple syrup urine disease patients from Iran. 2017 , 32, 105-113	12
131	Primary Immunodeficiency Diseases in Highly Consanguineous Populations from Middle East and North Africa: Epidemiology, Diagnosis, and Care. 2017 , 8, 678	29
130	A Novel Pathologic Variant in OTOF in an Iranian Family Segregating Hereditary Hearing Loss. 2018 , 158, 1084-1092	2
129	The First Report of Relative Incidence of Inherited White Matter Disorders in an Asian Country Based on an Iranian Bioregistry System. 2018 , 33, 255-259	9
128	Four novel mutations of the BCKDHA, BCKDHB and DBT genes in Iranian patients with maple syrup urine disease. 2018 , 31, 205-212	2
127	CONSANGUINEOUS MARRIAGES AMONG IRANIAN MANDAEANS LIVING IN SOUTH-WEST IRAN. 2018 , 50, 451-456	5
126	Clinical, immunologic, and genetic spectrum of 696 patients with combined immunodeficiency. 2018 , 141, 1450-1458	56
125	Management of rare coagulation disorders in 2018. 2018 , 57, 705-712	17
124	Molecular genetics of a cohort of 635 cases of phenylketonuria in a consanguineous population. 2018 , 41, 1159-1167	7
123	Molecular genetic study of Calpainopathy in Iran. 2018 , 677, 259-265	2
122	Autozygosity mapping of methylmalonic acidemia associated genes by short tandem repeat markers facilitates the identification of five novel mutations in an Iranian patient cohort. 2018 , 33, 1689-1697	5
121	Does Gold Price Matter for Divorce Rate in Iran?. 2018 , 39, 588-599	2

120	Frequency of Inborn Errors of Metabolism in a Northeastern Iranian Sample with High Consanguinity Rates. 2018 , 83, 71-78	5
119	High Incidence of Severe Combined Immunodeficiency Disease in Saudi Arabia Detected Through Combined T Cell Receptor Excision Circle and Next Generation Sequencing of Newborn Dried Blood Spots. 2018 , 9, 782	34
118	Long-term exposure to ambient air pollution and autism spectrum disorder in children: A case-control study in Tehran, Iran. 2018 , 643, 1216-1222	37
117	Clinical subtypes and molecular basis of epidermolysis bullosa in Kuwait. 2018 , 57, 1058-1067	2
116	Outcomes Following Allogeneic Stem Cell Transplantation Using Non-sibling Family Donors. 2019 , 35, 43-49	2
115	Clinical and molecular assessment of 13 Iranian families with Wolfram syndrome. 2019 , 66, 185-191	1
114	Iranome: A catalog of genomic variations in the Iranian population. 2019 , 40, 1968-1984	63
113	Type 1 diabetes genetic risk score discriminates between monogenic and Type 1 diabetes in children diagnosed at the age of . 2019 , 36, 1694-1702	8
112	Distinct genetic variation and heterogeneity of the Iranian population. 2019 , 15, e1008385	18
111	An update of spectrum and frequency of GJB2 mutations causing hearing loss in the south of Iran: A literature review. 2019 , 119, 136-140	16
110	Maple syrup urine disease mutation spectrum in a cohort of 40 consanguineous patients and insilico analysis of novel mutations. 2019 , 34, 1145-1156	6
109	A pathogenic variant in is associated with Pendred syndrome in a consanguineous Iranian family. 2019 , 58, 628-634	0
108	Molecular genetic diagnosis of Glanzmann syndrome in Iranian population; reporting novel and recurrent mutations. 2019 , 14, 87	4
107	Genetics and genomic medicine in Iran. 2019 , 7, e00606	3
106	Parental consanguinity in patients with psychogenic nonepileptic seizures. 2019 , 94, 167-168	1
105	Autosomal recessive congenital ichthyosis: Genomic landscape and phenotypic spectrum in a cohort of 125 consanguineous families. 2019 , 40, 288-298	21
104	A novel association of rs13334070 in the RPGRIP1L gene with adiposity factors discovered by joint linkage and linkage disequilibrium analysis in Iranian pedigrees: Tehran Cardiometabolic Genetic Study (TCGS). 2019 , 43, 342-351	4
103	Congenital neutropenia and primary immunodeficiency diseases. 2019 , 133, 149-162	15

102	Genome-wide single nucleotide polymorphism-based autozygosity mapping facilitates identification of mutations in consanguineous families with epidermolysis bullosa. 2019 , 28, 1118-1121	13
101	Genetics of intellectual disability in consanguineous families. 2019 , 24, 1027-1039	79
100	Role of and in Iranian Nonsyndromic Hearing Impairment: From Molecular Analysis to Literature Reviews. 2020 , 39, 1-12	12
99	Clinical and laboratory diagnosis of rare coagulation disorders (RCDs). 2020 , 196, 603-608	1
98	Homozygosity mapping and direct sequencing identify a novel pathogenic variant in the CISD2 gene in an Iranian Wolfram syndrome family. 2020 , 57, 81-87	6
97	c.1227_1228dupGG (p.Glu410Glyfs), a frequent variant in Tunisian patients with MUTYH associated polyposis. 2020 , 240, 45-53	0
96	Consanguineous marriages among Andalusian : a genealogical analysis (1925-2006). 2020 , 52, 809-831	2
95	Assessment of the correlation between various risk factors and orofacial cleft disorder spectrum: a retrospective case-control study. 2020 , 42, 26	3
94	Corneal dystrophies. 2020 , 6, 46	7
93	Pre-implantation genetic diagnosis in an Iranian family with a novel mutation in MUT gene. 2020 , 21, 22	3
92	Copy number variations in miscarriage products and their relationship with consanguinity and recurrent miscarriage in individuals with normal karyotypes. 2020 , 51, 101526	
91	Genomics-based treatment in a patient with two overlapping heritable skin disorders: Epidermolysis bullosa and acrodermatitis enteropathica. 2020 , 41, 906-912	7
90	Genotype-phenotype correlation and description of two novel mutations in Iranian patients with glycogen storage disease 1b (GSD1b). 2020 , 15, 35	1
89	A novel pathogenic variant of gene causes rigidity and multifocal seizure syndrome, lethal neonatal. 2021 , 131, 875-878	3
88	Next-generation sequencing reveals a novel pathogenic variant in the gene. 2021 , 1-5	1
87	Comprehensive Mutation Analysis and Report of 12 Novel Mutations in a Cohort of Patients with Spinal Muscular Atrophy in Iran. 2021 , 71, 2281-2298	3
86	Development of a risk prediction model for early discrimination between permanent and transient congenital hypothyroidism. 2021 , 73, 374-383	0
85	Novel SCN5A variants identified in a group of Iranian Brugada syndrome patients. 2021 , 21, 331-340	

84	Clinical and genetic spectrum of glycogen storage disease in Iranian population using targeted gene sequencing. 2021 , 11, 7040	2
83	Identifying the causes of recurrent pregnancy loss in consanguineous couples using whole exome sequencing on the products of miscarriage with no chromosomal abnormalities. 2021 , 11, 6952	3
82	The spectrum of ATM gene mutations in Iranian patients with ataxia-telangiectasia. 2021 , 32, 1316-1326	5
81	Identification of a novel MICU1 nonsense variant causes myopathy with extrapyramidal signs in an Iranian consanguineous family. 2021 , 8, 6	1
80	Consanguineous marriage and rare bleeding disorders. 2021 , 14, 467-472	0
79	Consanguineous Marriage and the Risk of Parents Having Children with Cleft Lip and Palate. 2021 , In Press,	1
78	Detection of Aneuploidies in Products of Conception and Neonatal Deaths in Iranian Patients Using the Multiplex Ligation-Dependent Probe Amplification (MLPA). 2021 , 13, 143-148	
77	Exome sequencing reveals novel rare variants in Iranian familial multiple sclerosis: The importance of POLD2 in the disease pathogenesis. 2021 , 113, 2645-2655	1
76	Phenotype of ST3GAL3 deficient patients: A case and review of the literature. 2021 , 64, 104250	3
75	CFTR gene mutation spectrum among 735 Iranian patients with cystic fibrosis: A comprehensive systematic review. 2021 , 56, 3644-3656	1
74	GJB2 mutations in Iranian Azeri population with autosomal recessive nonsyndromic hearing loss (ARNSHL): First report of c.238 C>A mutation in Iran. 2021 , 35, e24024	1
73	AutoMap is a high performance homozygosity mapping tool using next-generation sequencing data. 2021 , 12, 518	17
72	Genetics of Recessive Cognitive Disorders. 1-21	2
71	Common MEFV mutation analysis in 36 Iranian patients with familial Mediterranean fever: clinical and demographic significance. 2010 , 20, 566-72	11
70	Spectrum of PAH gene mutations in 1547 phenylketonuria patients from Iran: a comprehensive systematic review. 2021 , 36, 767-780	3
69	Male Infertility and Consanguinity in Lebanon: The Power of Ethnographic Epidemiology. 2009 , 165-195	3
68	Association between inbreeding coefficient and susceptibility to HIV-1 infection, a case-control study. 2013 , 3, 122-5	4
67	Prevalence and Socio-Demographic Characteristics Related to Stress, Anxiety, and Depression among Patients with Major Thalassemia in the Kermanshah County. 2015 , 4,	3

66	Epidemiological Study of Legal Abortion due to Fetal Defects in the Files Referred to Fars Province Forensic Medicine Centers from 2007 to 2013. 2016 , 17,	2
65	Knowledge and attitude toward genetic diseases and genetic tests among pre-marriage individuals: A cross-sectional study in northern Iran. 2019 , 17, 543-550	5
64	Applying Two Different Bioinformatic Approaches to Discover Novel Genes Associated with Hereditary Hearing Loss via Whole-Exome Sequencing: ENDEAVOUR and HomozygosityMapper. 2018 , 7, 141	2
63	Prevalence of Keratoconus among Young Arab students in Israel. 2014 , 3, 9-14	23
62	Consanguinity and isolated atrial septal defect in North East of Iran. 2014 , 34, 147-52	2
61	Causes of Chronic Kidney Disease in Iranian Children: A Meta-Analysis and Systematic Review. 2019 , 85,	2
60	Genetics of Hearing Loss in North Iran Population: An Update of Spectrum and Frequency of GJB2 Mutations. 2019 , 23, 175-180	6
59	Genetic Investigation of 261 Cases of Turner Syndrome Patients Referred to the Genetic Clinic.. <i>Iranian Journal of Public Health</i> , 2021 , 50, 2065-2075	0.7
58	X-linked SCID with a rare mutation. 2021 , 17, 107	
57	A Cohort Perspective on Changes in Family, Fertility Behaviour and Attitudes. 2009 , 135-161	
56	Cognitive Predictors of Cousin Marriage Among Couples Visiting Counseling Centers in Kohgiluyeh-Boyer Ahmad Province. 2014 , 1, 47-53	0
55	Variations in Types of First-Cousin Marriages over a Two-Generation Period among Arabs in Israel. 2015 , 05, 171-176	1
54	An Overview of Hereditary Diseases in Khuzestan Province, Southwest Iran. 2015 , 2,	
53	Murder and Motivation: A Qualitative Study. 2015 , 2,	
52	Murder and Motivation: A Qualitative Study. 2015 , 2,	1
51	Domestic Violence Subjected to Different Patterns of Cultural Marriage. 2016 , 04, 87-98	1
50	Which outcomes do women expect to achieve after undergoing induced abortion. 2017 , 9, 3741-3750	0
49	Potential Founder Effect of Tyrosinase Gene Mutations in Oculocutaneous Albinism Families from West of Iran. 2017 , 1,	

48	Assisted Reproductive Technologies and Making and Unmaking of Kin in Iran: Transformation or Variation on a Theme?. 2018 , 83-93		1
47	SLC26A4 pathogenic variants as a third cause of hearing loss: Role of three exons in DFNB4 deafness in Iran. 2019 , 25, 146		
46	Prevalence of Hemoglobin Mutations and Hemoglobinopathies in Masjed Soleiman County, Southeastern Iran. 2019 , 13, 48-54		
45	Once in a Blue Moon, a Very Rare Coexistence of Glutaric Acidemia Type I and Mucopolysaccharidosis Type IIIB in a Patient. <i>Iranian Biomedical Journal</i> , 2020 , 24, 201-5	2	0
44	Interactions of Consanguinity and Number of Siblings with Childhood Acute Lymphoblastic Leukemia. <i>BioMed Research International</i> , 2020 , 2020, 7919310	3	
43	Rare Coagulation Factor Deficiencies. 2020 , 51-60		
42	Biliary Atresia in Northwest Iran: Epidemiologic Features and Long-Term Outcome. <i>Iranian Journal of Pediatrics</i> , 2020 , 30,	1	0
41	RELATIONSHIP BETWEEN LEAD AND CADMIUM LEVELS IN BLOOD AND REFRACTORY CHRONIC CONSTIPATION AMONG IRANIAN CHILDREN. <i>Arquivos De Gastroenterologia</i> , 2021 , 58, 329-336	1.3	0
40	Non-modifiable Factors of Coronary Artery Stenosis in Late Onset Patients with Coronary Artery Disease in Southern Iranian Population. <i>Journal of Cardiovascular and Thoracic Research</i> , 2014 , 6, 51-5	1.3	12
39	The prevalence and clinical study of galactosemia disease in a pilot screening program of neonates, southern iran. <i>Iranian Journal of Public Health</i> , 2011 , 40, 99-104	0.7	5
38	Genetic Linkage Analysis of 15 DFNB Loci in a Group of Iranian Families with Autosomal Recessive Hearing Loss. <i>Iranian Journal of Public Health</i> , 2011 , 40, 34-48	0.7	28
37	Chronic Kidney Disease Stages 3-5 in Iranian Children: Need for a School-based Screening Strategy: The CASPIAN-III Study. <i>International Journal of Preventive Medicine</i> , 2013 , 4, 95-101	1.6	14
36	Clinical and genetic features in patients with cystic fibrosis in southwestern iran. <i>Iranian Journal of Pediatrics</i> , 2013 , 23, 212-5	1	10
35	Down syndrome and consanguinity. <i>Journal of Research in Medical Sciences</i> , 2013 , 18, 995-7	1.6	2
34	Clinical and Para clinical Manifestations of Tuberous Sclerosis: A Cross Sectional Study on 81 Pediatric Patients. <i>Iranian Journal of Child Neurology</i> , 2012 , 6, 25-31	0.6	3
33	Consanguinity and neonatal death: a nested case-control study. <i>Journal of Family & Reproductive Health</i> , 2014 , 8, 189-93	0.6	2
32	Sequence analysis of tyrosinase gene in ocular and oculocutaneous albinism patients: introducing three novel mutations. <i>Molecular Vision</i> , 2015 , 21, 730-5	2.3	8
31	Heterogeneity of Hereditary Hearing Loss in Iran: a Comprehensive Review. <i>Archives of Iranian Medicine</i> , 2016 , 19, 720-728	2.4	16

30	Consanguineous marriages in the genetic counseling centers of Isfahan and the ethical issues of clinical consultations. <i>Journal of Medical Ethics and History of Medicine</i> , 2017 , 10, 112	0.6	4
29	Consanguineous Marriage as a Risk Factor for Developing Keratoconus. <i>Medical Hypothesis, Discovery, and Innovation in Ophthalmology</i> , 2018 , 7, 17-21	1.4	4
28	Inherited Bleeding Disorders in Iraq and Consanguineous Marriage. <i>International Journal of Hematology-Oncology and Stem Cell Research</i> , 2018 , 12, 273-281	0.5	2
27	Impact of group assertiveness-based sexual training on the quality of marital relationships among female university students. <i>Journal of Education and Health Promotion</i> , 2019 , 8, 111	1.4	1
26	CRB1-Related Leber Congenital Amaurosis: Reporting Novel Pathogenic Variants and a Brief Review on Mutations Spectrum. <i>Iranian Biomedical Journal</i> , 2019 , 23, 362-8	2	
25	Brief Report of Variants Detected in Hereditary Hearing Loss Cases in Iran over a 3-Year Period. <i>Iranian Journal of Public Health</i> , 2019 , 48, 1910-1915	0.7	
24	Characterization of Niemann-Pick diseases genes mutation spectrum in Iran and identification of a novel mutation in gene. <i>Medical Journal of the Islamic Republic of Iran</i> , 2019 , 33, 126	1.1	
23	Gene Mutations in Non-Syndromic Hearing Loss of Bloch, Kurd, and Turkmen Ethnicities in Iran. <i>Iranian Journal of Public Health</i> , 2020 , 49, 2128-2135	0.7	
22	Letter to the Editor Regarding "Nationwide Prevalence of Diabetes and Prediabetes and Associated Risk Factors Among Iranian Adults: Analysis of Data from PERSIAN Cohort Study". <i>Diabetes Therapy</i> , 2021 , 13, 217	3.6	1
21	Gjb3 Gene Mutations in Non-Syndromic Hearing Loss of Bloch, Kurd, and Turkmen Ethnicities in Iran. <i>Iranian Journal of Public Health</i> , 2020 , 49, 2128-2135	0.7	
20	Gastrointestinal Bleeding in Congenital Bleeding Disorders.. <i>Seminars in Thrombosis and Hemostasis</i> , 2022 ,	5.3	0
19	Morbidity and mortality of COVID-19 negatively associated with the frequency of consanguineous marriages, an ecologic study. <i>Egyptian Journal of Medical Human Genetics</i> , 2022 , 23,	2	
18	Genetic etiology of hearing loss in Iran.. <i>Human Genetics</i> , 2022 , 1	6.3	1
17	A novel missense variant in ESRRB gene causing autosomal recessive non-syndromic hearing loss: in silico analysis of a case.. <i>BMC Medical Genomics</i> , 2022 , 15, 18	3.7	
16	Epidemiology of familial multiple sclerosis in Iran: a national registry-based study.. <i>BMC Neurology</i> , 2022 , 22, 76	3.1	0
15	Comparative Analyses of Turkish Variome and Widely Used Genomic Variation Databases for the Evaluation of Rare Sequence Variants in Turkish Individuals: Idiopathic Hypogonadotropic Hypogonadism as a Disease Model.. <i>JCRPE Journal of Clinical Research in Pediatric Endocrinology</i> , 2022 ,	1.9	0
14	table_1.PDF. 2018 ,		
13	table_2.PDF. 2018 ,		

12	Clinical, biological, and genetic features in an afibrinogenemia patient series in Algeria.. <i>Haemophilia</i> , 2022 ,	3.3	○
11	Evaluation of the Correlation between Socioeconomic Factors and Pediatric Cleft Lip and Palate. <i>Modern Medical Laboratory Journal</i> , 2021 , 4, 40-49	0.5	○
10	Analysis of early neonatal case fatality rate among newborns with congenital hydrocephalus, a 2000-2014 multi-country registry-based study. <i>Birth Defects Research</i> ,	2.9	○
9	Relationship between risk factors of hearing loss and the results of otoacoustic emission in newborns. <i>Hearing, Balance and Communication</i> , 1-6	0.7	○
8	Determining the Changes in the Prevalence of Consanguineous Marriages and Resistant Groups in Turkey: What the 2018 Turkey Demographic and Health Survey Indicates?.		1
7	The psychosocial experiences of girls with visual impairment about the ideal spouse and marriage. 0264619622211244		○
6	Epidemiology of familial multiple sclerosis and its comparison to sporadic form in Markazi Province, Iran. 2022 , 68, 104231		○
5	Chloride Channel Mutations Leading to Congenital Myotonia. 2022 ,		○
4	Relative frequency of inherited retinal disorders in Khuzestan province, southern Iran. 2023 , 13, 42-45		○
3	The First Report of Iranian Registry of Patients with Spinal Muscular Atrophy. 2023 , 10, 211-225		○
2	Historical Cohort of Severe Congenital Neutropenia in Iran: Clinical Course, Laboratory Evaluation, Treatment, and Survival. Publish Ahead of Print,		○
1	Epidemiological and Spatiotemporal Descriptive Analysis of Patients with Nonsyndromic Cleft Lip and/or Palate: A 12-Year Retrospective Study in Southern Iran. 2023 , 2023, 1-19		○