Ioannis A Georgiou

List of Publications by Year
in descending order

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145 papers	3,336 citations	201385 27 h-index	214527 47 g-index
148 all docs	148 docs citations	148 times ranked	4289 citing authors

#	Article	IF	Citations
1	Fragile X premutation is a significant risk factor for premature ovarian failure: The international collaborative POF in fragile X study?preliminary data. , 1999, 83, 322-325.		405
2	Arrayed Primer Extension: Solid-Phase Four-Color DNA Resequencing and Mutation Detection Technology. Genetic Testing and Molecular Biomarkers, 2000, 4, 1-7.	1.7	167
3	CpG methylation analysis of the MEG3 and SNRPN imprinted genes in acute myeloid leukemia and myelodysplastic syndromes. Leukemia Research, 2010, 34, 148-153.	0.4	150
4	Association of estrogen receptor gene polymorphisms with endometriosis. Fertility and Sterility, 1999, 72, 164-166.	0.5	148
5	Endothelial Function, But Not Carotid Intima-Media Thickness, Is Affected Early in Menopause and Is Associated with Severity of Hot Flushes. Journal of Clinical Endocrinology and Metabolism, 2010, 95, 1199-1206.	1.8	126
6	Association of the (TAAAA)n Repeat Polymorphism in the Sex Hormone-Binding Globulin (SHBG) Gene with Polycystic Ovary Syndrome and Relation to SHBG Serum Levels. Journal of Clinical Endocrinology and Metabolism, 2003, 88, 5976-5980.	1.8	117
7	A polymorphism in the 3â€™-UTR of interleukin-1 receptor-associated kinase (IRAK1), a target gene of miR-146a, is associated with rheumatoid arthritis susceptibility. Joint Bone Spine, 2010, 77, 411-413.	0.8	108
8	Genetic and epigenetic risks of intracytoplasmic sperm injection method. Asian Journal of Andrology, 2006, 8, 643-673.	0.8	89
9	Promoter Hypermethylation of the MEG3 (DLK1/MEG3) Imprinted Gene in Multiple Myeloma. Clinical Lymphoma and Myeloma, 2008, 8, 171-175.	1.4	88
10	Effect of Adiponectin Gene Polymorphisms on Circulating Adiponectin and Insulin Resistance Indexes in Women with Polycystic Ovary Syndrome. Clinical Chemistry, 2005, 51, 416-423.	1.5	82
11	miRNAs and related polymorphisms in rheumatoid arthritis susceptibility. Autoimmunity Reviews, 2012, 11, 636-641.	2.5	71
12	Interaction between the polymorphisms of the reninâ€“angiotensin system in preeclampsia. European Journal of Obstetrics, Gynecology and Reproductive Biology, 2003, 110, 8-11.	0.5	67
13	Retrotransposon RNA expression and evidence for retrotransposition events in human oocytes. Human Molecular Genetics, 2009, 18, 1221-1228.	1.4	65
14	CYP19 gene: a genetic modifier of polycystic ovary syndrome phenotype. Fertility and Sterility, 2010, 94, 250-254.	0.5	50
15	Association of the T45G polymorphism in exon 2 of the adiponectin gene with polycystic ovary syndrome: role of Â4-androstenedione. Human Reproduction, 2004, 19, 1728-1733.	0.4	45
16	Vitamin D receptor gene polymorphisms in multiple sclerosis patients in northwest Greece. Journal of Negative Results in BioMedicine, 2011, 10, 3.	1.4	44
17	Clinical application of whole-genome array CGH during prenatal diagnosis: Study of 25 selected pregnancies with abnormal ultrasound findings or apparently balanced structural aberrations. Molecular Cytogenetics, 2010, 3, 24.	0.4	43
18	The adiponectin-to-leptin ratio in women with polycystic ovary syndrome: relation to insulin resistance and proinflammatory markers. Metabolism: Clinical and Experimental, 2007, 56, 766-771.	1.5	41

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19	The synergistic effect of sex hormone-binding globulin and aromatase genes on polycystic ovary syndrome phenotype.. European Journal of Endocrinology, 2008, 158, 861-865.	1.9	39
20	Transcription regulatory polymorphism â^'43T>C in the 5â€²-flanking region of SLC19A1 gene could affect rheumatoid arthritis patient response to methotrexate therapy. Rheumatology International, 2007, 27, 1057-1061.	1.5	37
21	GENOTYPES OF N-ACETYLTRANSFERASE-2 AND RISK OF BLADDER CANCER: A CASE-CONTROL STUDY. Journal of Urology, 1999, 161, 1672-1675.	0.2	36
22	Estrogen Receptor Â and Â Polymorphisms Are Associated With Semen Quality. Journal of Andrology, 2010, 31, 291-298.	2.0	35
23	Assessment of sperm chromatin condensation and ploidy status using flow cytometry correlates to fertilization, embryo quality and pregnancy following in vitro fertilization. Journal of Assisted Reproduction and Genetics, 2011, 28, 885-891.	1.2	34
24	FoSTeS, MMBIR and NAHR at the human proximal Xp region and the mechanisms of human Xq isochromosome formation. Human Molecular Genetics, 2011, 20, 1925-1936.	1.4	34
25	HOX A10 and HOX A11 mutation scan in congenital malformations of the female genital tract. Reproductive BioMedicine Online, 2010, 21, 126-132.	1.1	33
26	Personal protective equipment and infection prevention and control: a national survey of UK medical students and interim foundation doctors during the COVID-19 pandemic. Journal of Public Health, 2021, 43, 67-75.	1.0	33
27	Fragile X premutations and (TA)n estrogen receptor polymorphism in women with ovarian dysfunction. , 1999, 84, 306-308.		30
28	Glutathione S–Transferase Null Genotypes in Transitional Cell Bladder Cancer. European Urology, 2000, 37, 660-664.	0.9	29
29	A polymorphism in the resistin gene promoter is associated with body mass index in women with polycystic ovary syndrome. Fertility and Sterility, 2004, 82, 1466-1467.	0.5	29
30	A Watershed Based Segmentation Method for Multispectral Chromosome Images Classification. , 2006, 2006, 3009-12.		29
31	Î´-Thalassemia in Cyprus. Hemoglobin, 2006, 30, 455-462.	0.4	28
32	Ultralong administration of gonadotropin-releasing hormone agonists before inÂvitro fertilization improves fertilization rate but not clinical pregnancy rate in women with mild endometriosis: a prospective, randomized, controlled trial. Fertility and Sterility, 2020, 113, 828-835.	0.5	28
33	Global profiling of EGFR gene mutation, amplification, regulation and tissue protein expression in unknown primary carcinomas: to target or not to target?. Clinical and Experimental Metastasis, 2007, 24, 79-86.	1.7	26
34	CYP19gene variants affect the assisted reproduction outcome of women with polycystic ovary syndrome. Gynecological Endocrinology, 2013, 29, 478-482.	0.7	26
35	FRAXA and FRAXE prevalence in patients with nonspecific mental retardation in the Hellenic population. Genetic Epidemiology, 1998, 15, 103-109.	0.6	25
36	How does closed system vitrification of human oocytes affect the clinical outcome? A prospective, observational, cohort, noninferiority trial in an oocyte donation program. Fertility and Sterility, 2016, 106, 1348-1355.	0.5	25

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37	Impact of renin-angiotensin-aldosterone system genes on the treatment response of patients with hypertension and metabolic syndrome. JRAAS - Journal of the Renin-Angiotensin-Aldosterone System, 2007, 8, 181-189.	1.0	23
38	A prospective randomized study comparing gonadotropin-releasing hormone agonists or gonadotropin-releasing hormone antagonists in couples with unexplained infertility and/or mild oligozoospermia. Fertility and Sterility, 2005, 83, 1354-1362.	0.5	21
39	The Pro12Ala polymorphism of the PPAR-Î³ gene is not associated with the polycystic ovary syndrome. Hormones, 2009, 8, 267-272.	0.9	21
40	The association of aromatase (CYP19) gene variants with sperm concentration and motility. Asian Journal of Andrology, 2011, 13, 292-297.	0.8	21
41	Preimplantation genetic diagnosis for spinal and bulbar muscular atrophy (SBMA). Human Genetics, 2001, 108, 494-498.	1.8	20
42	Functional impact and prevalence of polymorphisms involved in the hepatic glucuronidation of valproic acid. Pharmacogenomics, 2012, 13, 1055-1071.	0.6	20
43	The ovarian response to standard gonadotrophin stimulation depends on FSHR, SHBG and CYP19 gene synergism. Journal of Assisted Reproduction and Genetics, 2012, 29, 1185-1191.	1.2	19
44	Evidence for high-risk haplotypes and (CGG)n expansion in fragile X syndrome in the Hellenic population of Greece and Cyprus. , 1996, 64, 234-238.		18
45	Evidence for association of SLC7A9 gene haplotypes with cystinuria manifestation in SLC7A9 mutation carriers. Urological Research, 2006, 34, 299-303.	1.5	18
46	Chronic NF-ÎºB activation delays RasV12-induced premature senescence of human fibroblasts by suppressing the DNA damage checkpoint response. Mechanisms of Ageing and Development, 2009, 130, 409-419.	2.2	18
47	Association of Paraoxonase Gene Polymorphisms With Sperm Parameters. Journal of Andrology, 2011, 32, 394-401.	2.0	18
48	Gender association of the angiotensin-converting enzyme gene with ischaemic stroke. JRAAS - Journal of the Renin-Angiotensin-Aldosterone System, 2011, 12, 510-515.	1.0	17
49	Retrotransposon expression and incorporation of cloned human and mouse retroelements in human spermatozoa. Fertility and Sterility, 2017, 107, 821-830.	0.5	17
50	Association of serum and follicular fluid SHBG levels andSHBG(TAAAA)npolymorphism with follicle size in women undergoing ovarian stimulation. Gynecological Endocrinology, 2011, 27, 27-32.	0.7	16
51	Recombinant human erythropoietin treatment of postpartum anemia. European Journal of Obstetrics, Gynecology and Reproductive Biology, 1998, 81, 27-31.	0.5	15
52	Effect of recombinant human erythropoietin in preterm infants. European Journal of Haematology, 1999, 63, 71-76.	1.1	15
53	Association of TNFÂ, TNFR1, and TNFR2 Polymorphisms With Sperm Concentration and Motility. Journal of Andrology, 2012, 33, 74-80.	2.0	15
54	The importance of the (TAAAA)n alleles at the SHBG gene promoter for the severity of coronary artery disease in postmenopausal women. Menopause, 2008, 15, 461-468.	0.8	14

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55	Short-term hormone therapy improves sCD40L and endothelial function in early menopausal women: Potential role of estrogen receptor polymorphisms. Maturitas, 2012, 71, 389-395.	1.0	14
56	TCF7L2 gene variants predispose to the development of type 2 diabetes mellitus among individuals with metabolic syndrome. Hormones, 2018, 17, 359-365.	0.9	14
57	Reproductive health in patients with epilepsy. Epilepsy and Behavior, 2020, 113, 107563.	0.9	14
58	R831X mutation of the androgen receptor gene in an adolescent with Complete Androgen Insensitivity Syndrome and bilateral testicular hamartomata. Hormones, 2006, 5, 200-204.	0.9	14
59	Novel PKD1 deletions and missense variants in a cohort of Hellenic polycystic kidney disease families. European Journal of Human Genetics, 2001, 9, 677-684.	1.4	13
60	Identification of Novel Cystinuria Mutations and Polymorphisms in SLC3A1 and SLC7A9 Genes: Absence of SLC7A10 Gene Mutations in Cystinuric Patients. Genetic Testing and Molecular Biomarkers, 2005, 9, 175-184.	1.7	13
61	Non-Invasive Prenatal Detection of Paternal Origin Hb Lepore in a Male Fetus at the 7th Week of Gestation. Fetal Diagnosis and Therapy, 2006, 21, 506-509.	0.6	13
62	Methylation status of the SNRPN and HUMARA genes in testicular biopsy samples. Fertility and Sterility, 2007, 87, 805-809.	0.5	13
63	UGT1A6- and UGT2B7-related valproic acid pharmacogenomics according to age groups and total drug concentration levels. Pharmacogenomics, 2016, 17, 827-835.	0.6	13
64	Insights into the Role of Telomeres in Human Embryological Parameters. Opinions Regarding IVF. Journal of Developmental Biology, 2021, 9, 49.	0.9	13
65	Prenatal Paternity Testing Using DNA Extracted from Coelomic Cells. Fetal Diagnosis and Therapy, 2004, 19, 75-77.	0.6	12
66	Region Based Segmentation and Classification of Multispectral Chromosome Images. Proceedings of the IEEE Symposium on Computer-Based Medical Systems, 2007, , .	0.0	12
67	Absence of methylation-dependent transcriptional silencing in TP73 irrespective of the methylation status of the CDKN2A CpG island in plasma cell neoplasia. Leukemia Research, 2009, 33, 1272-1275.	0.4	12
68	Association of the SHBG gene promoter polymorphism with early markers of atherosclerosis in apparently healthy women. Atherosclerosis, 2011, 219, 205-210.	0.4	12
69	The ovarian response to standard gonadotropin stimulation is influenced by AMHRII genotypes. Gynecological Endocrinology, 2016, 32, 641-645.	0.7	12
70	High HbF in pregnancy is associated with the Xmn I polymorphism at the â^'158bp of the Gî³-globin gene. European Journal of Obstetrics, Gynecology and Reproductive Biology, 1995, 60, 153-156.	0.5	11
71	Non-invasive first-trimester detection of paternal beta-globin gene mutations and polymorphisms as predictors of thalassemia risk at chorionic villous sampling. European Journal of Obstetrics, Gynecology and Reproductive Biology, 2008, 140, 17-20.	0.5	11
72	Enhancement of Multichannel Chromosome Classification Using a Region-Based Classifier and Vector Median Filtering. IEEE Transactions on Information Technology in Biomedicine, 2009, 13, 561-570.	3.6	11

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73	Aromatase (CYP19) gene variants influence ovarian response to standard gonadotrophin stimulation. Journal of Assisted Reproduction and Genetics, 2012, 29, 203-209.	1.2	11
74	Expanded Cytological Referral Criteria for Colposcopy in Cervical Screening: Comparison with Human Papillomavirus Testing. Gynecologic Oncology, 2001, 82, 355-359.	0.6	10
75	Distinct association of SLC19A1 polymorphism â˜'43T>C with red cell folate levels and of MTHFR polymorphism 677C>T with plasma folate levels. Clinical Biochemistry, 2008, 41, 174-176.	0.8	10
76	Using semen flow cytometry to evaluate association of ploidy status and chromatin condensation of spermatozoa with conventional semen parameters: Clinical application in intrauterine insemination. Fertility and Sterility, 2011, 95, 110-115.	0.5	10
77	Sperm flow cytometric parameters are associated with ICSI outcome. Reproductive BioMedicine Online, 2013, 26, 611-618.	1.1	10
78	Ovarian hyperstimulation syndrome inhibition by targeting VEGF, COX-2 and Calcium pathways: a preclinical randomized study. Gynecological Endocrinology, 2014, 30, 587-592.	0.7	10
79	Effects of Different Drug Therapies and COVID-19 mRNA Vaccination on Semen Quality in a Man with Ankylosing Spondylitis: A Case Report. Medicina (Lithuania), 2022, 58, 173.	0.8	10
80	An overview of SLC3A1 and SLC7A9 mutations in Greek cystinuria patients. Molecular Genetics and Metabolism, 2008, 95, 192-193.	0.5	9
81	Association of the A1330V and V667M polymorphisms of LRP5 with bone mineral density in Greek peri- and postmenopausal women. Maturitas, 2011, 70, 188-193.	1.0	9
82	Associations of ESR2 AluI (G/A) polymorphism with ischemic stroke in Caucasians. Journal of the Neurological Sciences, 2012, 316, 126-130.	0.3	9
83	Validation of the TAGAP rs212389 polymorphism in rheumatoid arthritis susceptibility. Joint Bone Spine, 2013, 80, 543-544.	0.8	9
84	Copy-number variation analysis in familial nonsyndromic congenital anomalies of the kidney and urinary tract: Evidence for the causative role of a transposable element-associated genomic rearrangement. Molecular Medicine Reports, 2017, 15, 3631-3636.	1.1	9
85	Unusual N Gene Dropout and Ct Value Shift in Commercial Multiplex PCR Assays Caused by Mutated SARS-CoV-2 Strain. Diagnostics, 2022, 12, 973.	1.3	9
86	Mosaic Trisomy r(14) Associated With Epilepsy and Mental Retardation. Journal of Child Neurology, 2007, 22, 869-873.	0.7	8
87	Everolimus, an mTOR pathway inhibitor, is highly successful on ovarian hyperstimulation syndrome by reducing ovarian weight and progesterone levels: a preclinical experimental randomized controlled study. Gynecological Endocrinology, 2015, 31, 702-707.	0.7	8
88	Infertility and Multiple Urogenital Abnormalities in a Male with Mosaic 46,XY/45,XO/47,XXY Karyotype and Mixed Phenotype. Urologia Internationalis, 1998, 61, 111-114.	0.6	7
89	Angiotensin II Type 2 Receptor Gene Polymorphism in Caucasian Children With a Wide Spectrum of Congenital Anomalies of the Kidney and Urinary Tract. Pediatric Research, 2007, 62, 83-87.	1.1	7
90	Estrogen Receptor Î± Gene Haplotypes and Diplotypes in the Risk of Stroke. Stroke, 2008, 39, e172-3; author reply e174.	1.0	7

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91	Association of the (<scp>TTTA</scp>)n repeat polymorphism of <scp>CYP</scp>19 gene with bone mineral density in <scp>G</scp>reek periâ€•and postmenopausal women. Clinical Endocrinology, 2014, 81, 38-44.	1.2	7
92	Holliday Junctions Are Associated with Transposable Element Sequences in the Human Genome. Journal of Molecular Biology, 2016, 428, 658-667.	2.0	7
93	Biological and Clinical Significance of Mosaicism in Human Preimplantation Embryos. Journal of Developmental Biology, 2021, 9, 18.	0.9	7
94	Interplay Between mTOR and Hippo Signaling in the Ovary: Clinical Choice Guidance Between Different Gonadotropin Preparations for Better IVF. Frontiers in Endocrinology, 2021, 12, 702446.	1.5	7
95	Phosphatidylethanolamine N-methyltransferase and choline dehydrogenase gene polymorphisms are associated with human sperm concentration. Asian Journal of Andrology, 2012, 14, 778-783.	0.8	7
96	Renin-angiotensin-aldosterone system genes and nonarteritic anterior ischemic optic neuropathy. Molecular Vision, 2011, 17, 1254-60.	1.1	7
97	Absence of mutations in the HOXA11 and HOXD11 genes in children with congenital renal malformations. Pediatric Nephrology, 2009, 24, 1569-1572.	0.9	6
98	Association of endothelial nitric oxide synthase polymorphism G894T with functional outcome in acute stroke patients. Neurological Research, 2011, 33, 835-840.	0.6	6
99	The follicular outcome after standard gonadotropin stimulation is associated with ERÎ± and ERÎ² genotypes. Endocrine, 2014, 47, 930-935.	1.1	6
100	Glutathione sulfur transferase M1 and T1 genotypes in chronic lymphoblastic leukemia. The Hematology Journal, 2004, 5, 500-504.	2.0	5
101	Absence of mutations of the EPO-receptor gene in Greek patients with familiar polycythemia. European Journal of Haematology, 2006, 76, 537-538.	1.1	5
102	Identification of the four most common Î²â€globin gene mutations in Greek Î²â€thalassemic patients and carriers by PCRâ€SSCP: advantages and limitations of the method. Journal of Clinical Laboratory Analysis, 2006, 20, 1-7.	0.9	5
103	Developing a genomic-based point-of-care diagnostic system for rheumatoid arthritis and multiple sclerosis. , 2009, 2009, 827-30.		5
104	Follicle inhibition at the primordial stage without increasing apoptosis, with a combination of everolimus, verapamil. Molecular Biology Reports, 2020, 47, 8711-8726.	1.0	5
105	Presence of HPV 16 and HPV 18 in Spermatozoa and Embryos of Mice. In Vivo, 2021, 35, 3203-3209.	0.6	5
106	Postâ€fertilization effects of chronic renal failure in male rats. Journal of Developmental and Physical Disabilities, 2009, 32, 675-686.	3.6	4
107	Endothelial nitric oxide synthase polymorphism (G894T) and nonarteritic anterior ischemic optic neuropathy. Visual Neuroscience, 2010, 27, 183-185.	0.5	4
108	Role of 9p21 and 2q36 variants and arterial stiffness in the prediction of coronary artery disease. European Journal of Clinical Investigation, 2014, 44, 784-794.	1.7	4

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109	Expression of Retroelements in Mammalian Gametes and Embryos. In Vivo, 2021, 35, 1921-1927.	0.6	4
110	ATG16L1 T300A polymorphism is associated with Crohnâ€™s disease in a Northwest Greek cohort, but ECM1 T130M and G290S polymorphisms are not associated with ulcerative colitis. Annals of Gastroenterology, 2019, 33, 38-44.	0.4	4
111	The Factors That Influenced Medical Studentsâ€™ Decision to Work Within the NHS During the COVID-19 Pandemicâ€"A National, Cross-sectional Study. Journal of Occupational and Environmental Medicine, 2021, 63, 296-301.	0.9	4
112	A gene for nonsyndromic X-linked mental retardation (MRX77) maps to Xq12-Xq21.33. American Journal of Medical Genetics Part A, 2003, 122A, 46-50.	2.4	3
113	Association ofTNF-857C>T,TNFRSF1A36A>G, andTNFRSF1B676T>GPolymorphisms with Ischemic Stroke in a Greek Population. Stroke Research and Treatment, 2011, 2011, 1-5.	0.5	3
114	The combination of Everolimus with Verapamil reduces ovarian weight and vascular permeability on ovarian hyperstimulation syndrome: a preclinical experimental randomized controlled study. Gynecological Endocrinology, 2016, 32, 886-890.	0.7	3
115	A Novel Îµ[3]Î[2]-Thalassemia Deletion Associated with Severe Anemia at Birth and a Î²-Thalassemia Intermedia Phenotype Later in Life in Three Generations of a Greek Family. Hemoglobin, 2021, 45, 351-354.	0.4	3
116	Reverse Transcriptase Affects Gametogenesis and Preimplantation Development in Mouse. In Vivo, 2020, 34, 2269-2276.	0.6	3
117	A mutation in NOTCH2 gene first associated with Hajduâ€"Cheney syndrome in a Greek family: diversity in phenotype and response to treatment. Endocrine, 2021, 71, 208-215.	1.1	3
118	Development and Validation of a Targeted â€˜Liquidâ€™ NGS Panel for Treatment Customization in Patients with Metastatic Colorectal Cancer. Diagnostics, 2021, 11, 2375.	1.3	3
119	Gammaâ€chain heterogeneity in Greek (Î[2])^Oâ€thalassemia.. European Journal of Haematology, 1995, 54, 101-105.	1.1	2
120	A novel nonsense mutation of the EPM2A gene in Northwest Greece causing myoclonic epilepsy. Seizure: the Journal of the British Epilepsy Association, 2013, 22, 315-317.	0.9	2
121	Association of the (TAAAA)n repeat polymorphism of SHBG gene with the age at menopause in Greek postmenopausal women. Maturitas, 2014, 78, 113-116.	1.0	2
122	ROBO2 gene variants in children with primary nonsyndromic vesicoureteral reflux with or without renal hypoplasia/dysplasia. Pediatric Research, 2016, 80, 72-76.	1.1	2
123	An open study of valproate in subfertile men with epilepsy. Acta Neurologica Scandinavica, 2020, 142, 317-322.	1.0	2
124	A Watershed Based Segmentation Method for Multispectral Chromosome Images Classification. Annual International Conference of the IEEE Engineering in Medicine and Biology Society, 2006, , .	0.5	2
125	Pathways Involved in Premature Ovarian Failure: A Systematic Review of Experimental Studies. Current Pharmaceutical Design, 2020, 26, 2087-2095.	0.9	2
126	Is Hysteroscopy Prior to IVF Associated with an Increased Probability of Live Births in Patients with Normal Transvaginal Scan Findings after Their First Failed IVF Trial?. Journal of Clinical Medicine, 2022, 11, 1217.	1.0	2

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127	Congenital obstructive azoospermia in a man with Marfan syndrome. Fertility and Sterility, 2001, 76, 1256-1257.	0.5	1
128	Distribution of Two X-Linked Trinucleotide Polymorphisms in Greece. Public Health Genomics, 2001, 4, 125-128.	1.0	1
129	Real-Time PCR Analysis of Trinucleotide Repeat Allele Expansions in the Androgen Receptor Gene. Molecular Diagnosis and Therapy, 2005, 9, 217-219.	1.3	1
130	Estrogen Receptor Alpha Gene Polymorphisms and Stroke. Cerebrovascular Diseases, 2008, 26, 338-338.	0.8	1
131	Molecular and Clinical Investigation of Cystinuria in the Greek-Cypriot Population. Genetic Testing and Molecular Biomarkers, 2015, 19, 641-645.	0.3	1
132	Fragile X premutations and (TA)n estrogen receptor polymorphism in women with ovarian dysfunction. , 1999, 84, 306.		1
133	An intelligent web-based system for the detection and visualization of biomarkers in Microdeletion and Microduplication Syndromes. , 2020, , .		1
134	Fetal Heart Rate Following Coelocentesis. Journal of Maternal-Fetal and Neonatal Medicine, 1997, 6, 314-316.	0.7	0
135	Increased nuchal translucency thickness in a fetus at risk for ?-thalassaemia. , 1997, 6, 301-302.		0
136	The role of deoxyribonuclease I in amniotic fluid during pregnancy and labour. European Journal of Obstetrics, Gynecology and Reproductive Biology, 1998, 77, 177-180.	0.5	0
137	Point-of-care monitoring and diagnostics for autoimmune diseases. , 2008, , .		0
138	Data analysis of Genome-Wide Association studies (GWAS) concerning rheumatoid arthritis and multiple sclerosis. , 2010, , .		0
139	Polymerase chain reaction (PCR) and sequence specific oligonucleotide probes (SSOP) genotyping assay for detection of genes associated with rheumatoid arthritis and multiple sclerosis. , 2010, 2010, 6202-5.		0
140	Mini- and Micro-Satellite Markers in Health, Disease and Evolution. , 2015, , 155-193.		0
141	Rare Association of Hb D-Los Angeles (HBB: c.364G>C) with Hb H Disease: Diagnosis and Clinical Implications. Hemoglobin, 2018, 42, 336-338.	0.4	0
142	A genetic platform for studying the creation of structural abnormalities of chromosomes that cause micro-deletion and micro-duplication (MMS) syndromes. , 2021, , .		0
143	A CLUSTERING â€“ BASED METHOD FOR GENE EXPRESSION COMPARISON IN EMBRYONIC STEM CELL DERIVED HEAMNGIOBLASTS AND ADULT BONE MARROW STROMAL CELLS. , 2010, , .		0
144	Pharmacological preconditioning for short-term ex vivo expansion of human umbilical cord blood hematopoietic stem cells by filgrastim. American Journal of Stem Cells, 2016, 5, 29-38.	0.4	0

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145	Social and psychological impact of the COVID-19 pandemic on UK medical and nursing students: protocol for a national medical and nursing student survey. BMJ Open, 2022, 12, e057467.	0.8	0