Ioannis A Georgiou

List of Publications by Year in Descending Order

138 papers	2,748 citations	25 h-index	47 g-index
148 ext. papers	3,022 ext. citations	3 avg, IF	4.43 L-index

#	Paper	IF	Citations
138	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,	3.1	2
137	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	3	
136	The Factors That Influenced Medical Students' Decision to Work Within the NHS During the COVID-19 Pandemic-A National, Cross-sectional Study. *Journal of Occupational and Environmental Medicine*, **2021**, 63, 296-301	2	3
135	Insights into the Role of Telomeres in Human Embryological Parameters. Opinions Regarding IVF. *Journal of Developmental Biology*, **2021**, 9,	3.5	1
134	Presence of HPV 16 and HPV 18 in Spermatozoa and Embryos of Mice. *In Vivo*, **2021**, 35, 3203-3209	2.3	1
133	Biological and Clinical Significance of Mosaicism in Human Preimplantation Embryos. *Journal of Developmental Biology*, **2021**, 9,	3.5	2
132	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	5.7	0
131	A mutation in NOTCH2 gene first associated with Hajdu-Cheney syndrome in a Greek family: diversity in phenotype and response to treatment. *Endocrine*, **2021**, 71, 208-215	4	2
130	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	3.5	16
129	Expression of Retroelements in Mammalian Gametes and Embryos. *In Vivo*, **2021**, 35, 1921-1927	2.3	1
128	Ultralong administration of gonadotropin-releasing hormone agonists before in´vitro 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	4.8	11
127	T300A polymorphism is associated with Crohn's disease in a Northwest Greek cohort, but T130M and G290S polymorphisms are not associated with ulcerative colitis. *Annals of Gastroenterology*, **2020**, 33, 38-44	2.2	2
126	Pathways Involved in Premature Ovarian Failure: A Systematic Review of Experimental Studies. *Current Pharmaceutical Design*, **2020**, 26, 2087-2095	3.3	1
125	An open study of valproate in subfertile men with epilepsy. *Acta Neurologica Scandinavica*, **2020**, 142, 317-322	3.8	1
124	Reproductive health in patients with epilepsy. *Epilepsy and Behavior*, **2020**, 113, 107563	3.2	5
123	Follicle inhibition at the primordial stage without increasing apoptosis, with a combination of everolimus, verapamil. *Molecular Biology Reports*, **2020**, 47, 8711-8726	2.8	3
122	Reverse Transcriptase Affects Gametogenesis and Preimplantation Development in Mouse. *In Vivo*, **2020**, 34, 2269-2276	2.3	3

121	A Novel □Thalassemia Deletion Associated with Severe Anemia at Birth and a □Thalassemia Intermedia Phenotype Later in Life in Three Generations of a Greek Family. *Hemoglobin*, **2019**, 1-4	0.6	2
120	Rare Association of Hb D-Los Angeles (HBB: c.364G>C) with Hb H Disease: Diagnosis and Clinical Implications. *Hemoglobin*, **2018**, 42, 336-338	0.6	
119	TCF7L2 gene variants predispose to the development of type 2 diabetes mellitus among individuals with metabolic syndrome. *Hormones*, **2018**, 17, 359-365	3.1	11
118	Retrotransposon expression and incorporation of cloned human and mouse retroelements in human spermatozoa. *Fertility and Sterility*, **2017**, 107, 821-830	4.8	9
117	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	2.9	8
116	UGT1A6- and UGT2B7-related valproic acid pharmacogenomics according to age groups and total drug concentration levels. *Pharmacogenomics*, **2016**, 17, 827-35	2.6	10
115	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	2.4	3
114	The ovarian response to standard gonadotropin stimulation is influenced by AMHRII genotypes. *Gynecological Endocrinology*, **2016**, 32, 641-645	2.4	9
113	ROBO2 gene variants in children with primary nonsyndromic vesicoureteral reflux with or without renal hypoplasia/dysplasia. *Pediatric Research*, **2016**, 80, 72-6	3.2	1
112	Holliday Junctions Are Associated with Transposable Element Sequences in the Human Genome. *Journal of Molecular Biology*, **2016**, 428, 658-667	6.5	2
111	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	2.4	
110	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	4.8	10
109	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, 702-7	2.4	6
108	Mini- and Micro-Satellite Markers in Health, Disease and Evolution **2015**, 155-193		
107	Molecular and Clinical Investigation of Cystinuria in the Greek-Cypriot Population. *Genetic Testing and Molecular Biomarkers*, **2015**, 19, 641-5	1.6	1
106	The follicular outcome after standard gonadotropin stimulation is associated with ER□and ER□ genotypes. *Endocrine*, **2014**, 47, 930-5	4	4
105	Ovarian hyperstimulation syndrome inhibition by targeting VEGF, COX-2 and calcium pathways: a preclinical randomized study. *Gynecological Endocrinology*, **2014**, 30, 587-92	2.4	8
104	Association of the (TTTA)n repeat polymorphism of CYP19 gene with bone mineral density in Greek peri- and postmenopausal women. *Clinical Endocrinology*, **2014**, 81, 38-44	3.4	7

103	Association of the (TAAAA)n repeat polymorphism of SHBG gene with the age at menopause in Greek postmenopausal women. *Maturitas*, **2014**, 78, 113-6	5	2
102	Role of 9p21 and 2q36 variants and arterial stiffness in the prediction of coronary artery disease. *European Journal of Clinical Investigation*, **2014**, 44, 784-94	4.6	2
101	A novel nonsense mutation of the EPM2A gene in northwest Greece causing myoclonic epilepsy. *Seizure: the Journal of the British Epilepsy Association*, **2013**, 22, 315-7	3.2	2
100	Validation of the TAGAP rs212389 polymorphism in rheumatoid arthritis susceptibility. *Joint Bone Spine*, **2013**, 80, 543-4	2.9	8
99	Sperm flow cytometric parameters are associated with ICSI outcome. *Reproductive BioMedicine Online*, **2013**, 26, 611-8	4	10
98	CYP19 gene variants affect the assisted reproduction outcome of women with polycystic ovary syndrome. *Gynecological Endocrinology*, **2013**, 29, 478-82	2.4	20
97	miRNAs and related polymorphisms in rheumatoid arthritis susceptibility. *Autoimmunity Reviews*, **2012**, 11, 636-41	13.6	63
96	Aromatase (CYP19) gene variants influence ovarian response to standard gonadotrophin stimulation. *Journal of Assisted Reproduction and Genetics*, **2012**, 29, 203-9	3.4	10
95	The ovarian response to standard gonadotrophin stimulation depends on FSHR, SHBG and CYP19 gene synergism. *Journal of Assisted Reproduction and Genetics*, **2012**, 29, 1185-91	3.4	16
94	Short-term hormone therapy improves sCD40L and endothelial function in early menopausal women: potential role of estrogen receptor polymorphisms. *Maturitas*, **2012**, 71, 389-95	5	11
93	Association of TNF, TNFR1, and TNFR2 polymorphisms with sperm concentration and motility. *Journal of Andrology*, **2012**, 33, 74-80		12
92	Associations of ESR2 AluI (G/A) polymorphism with ischemic stroke in Caucasians. *Journal of the Neurological Sciences*, **2012**, 316, 126-30	3.2	9
91	Functional impact and prevalence of polymorphisms involved in the hepatic glucuronidation of valproic acid. *Pharmacogenomics*, **2012**, 13, 1055-71	2.6	16
90	Phosphatidylethanolamine N-methyltransferase and choline dehydrogenase gene polymorphisms are associated with human sperm concentration. *Asian Journal of Andrology*, **2012**, 14, 778-83	2.8	5
89	Association of paraoxonase gene polymorphisms with sperm parameters. *Journal of Andrology*, **2011**, 32, 394-401		14
88	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, 110-5	4.8	9
87	Association of the A1330V and V667M polymorphisms of LRP5 with bone mineral density in Greek peri- and postmenopausal women. *Maturitas*, **2011**, 70, 188-93	5	9
86	Association of the SHBG gene promoter polymorphism with early markers of atherosclerosis in apparently healthy women. *Atherosclerosis*, **2011**, 219, 205-10	3.1	9

85	Association of TNF-857C>T, TNFRSF1A36A>G, and TNFRSF1B676T>G Polymorphisms with Ischemic Stroke in a Greek Population. *Stroke Research and Treatment*, **2011**, 2011, 920584	1.7	3
84	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, 885-91	3.4	26
83	Vitamin D receptor gene polymorphisms in multiple sclerosis patients in northwest Greece. *Journal of Negative Results in BioMedicine*, **2011**, 10, 3		40
82	Un polymorphisme de la r□gion 3?-UTR de la kinase associ□ au r□cepteur de l□interleukine-1 (IRAK´1), g□ne cible de miR-146a, est associ□□une susceptibilit□la polyarthrite rhumato□de. *Revue Du Rhumatisme (Edition Francaise)*, **2011**, 78, 242-244	0.1	
81	Association of serum and follicular fluid SHBG levels and SHBG (TAAAA)n polymorphism with follicle size in women undergoing ovarian stimulation. *Gynecological Endocrinology*, **2011**, 27, 27-32	2.4	14
80	Gender association of the angiotensin-converting enzyme gene with ischaemic stroke. *JRAAS - Journal of the Renin-Angiotensin-Aldosterone System*, **2011**, 12, 510-5	3	15
79	Association of endothelial nitric oxide synthase polymorphism G894T with functional outcome in acute stroke patients. *Neurological Research*, **2011**, 33, 835-40	2.7	5
78	FoSTeS, MMBIR and NAHR at the human proximal Xp region and the mechanisms of human Xq isochromosome formation. *Human Molecular Genetics*, **2011**, 20, 1925-36	5.6	29
77	Renin-angiotensin-aldosterone system genes and nonarteritic anterior ischemic optic neuropathy. *Molecular Vision*, **2011**, 17, 1254-60	2.3	7
76	The association of aromatase (CYP19) gene variants with sperm concentration and motility. *Asian Journal of Andrology*, **2011**, 13, 292-7	2.8	20
75	Endothelial nitric oxide synthase polymorphism (G894T) and nonarteritic anterior ischemic optic neuropathy. *Visual Neuroscience*, **2010**, 27, 183-5	1.7	3
74	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, 1199-206	5.6	112
73	Estrogen receptor alpha and beta polymorphisms are associated with semen quality. *Journal of Andrology*, **2010**, 31, 291-8		29
72	Polymerase chain reaction (PCR) and sequence specific oligonucleotide probes (SSOP) genotyping assay for detection of genes associated with rheumatoid arthritis and multiple sclerosis. *Annual International Conference of the IEEE Engineering in Medicine and Biology Society IEEE Engineering in Medicine and Biology Society Annual International Conference*, **2010**, 2010, 6202-5	0.9	
71	CYP19 gene: a genetic modifier of polycystic ovary syndrome phenotype. *Fertility and Sterility*, **2010**, 94, 250-4	4.8	42
70	HOX A10 and HOX A11 mutation scan in congenital malformations of the female genital tract. *Reproductive BioMedicine Online*, **2010**, 21, 126-32	4	19
69	A polymorphism in the 3'-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, 411-3	2.9	99
68	CpG methylation analysis of the MEG3 and SNRPN imprinted genes in acute myeloid leukemia and myelodysplastic syndromes. *Leukemia Research*, **2010**, 34, 148-53	2.7	120

67	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	2	37
66	The Pro12Ala polymorphism of the PPAR-gamma gene is not associated with the polycystic ovary syndrome. *Hormones*, **2009**, 8, 267-72	3.1	19
65	Retrotransposon RNA expression and evidence for retrotransposition events in human oocytes. *Human Molecular Genetics*, **2009**, 18, 1221-8	5.6	54
64	Chronic NF-kappaB activation delays RasV12-induced premature senescence of human fibroblasts by suppressing the DNA damage checkpoint response. *Mechanisms of Ageing and Development*, **2009**, 130, 409-19	5.6	15
63	Enhancement of multichannel chromosome classification using a region-based classifier and vector median filtering. *IEEE Transactions on Information Technology in Biomedicine*, **2009**, 13, 561-70		7
62	Gamma-chain heterogeneity in Greek (β0-thalassemia.. *European Journal of Haematology*, **2009**, 54, 101-105	3.8	2
61	Absence of mutations in the HOXA11 and HOXD11 genes in children with congenital renal malformations. *Pediatric Nephrology*, **2009**, 24, 1569-72	3.2	5
60	Post-fertilization effects of chronic renal failure in male rats. *Journal of Developmental and Physical Disabilities*, **2009**, 32, 675-86		2
59	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, 1272-5	2.7	10
58	Developing a genomic-based point-of-care diagnostic system for rheumatoid arthritis and multiple sclerosis. *Annual International Conference of the IEEE Engineering in Medicine and Biology Society IEEE Engineering in Medicine and Biology Society Annual International Conference*, **2009**, 2009, 827-30	0.9	4
57	Promoter hypermethylation of the MEG3 (DLK1/MEG3) imprinted gene in multiple myeloma. *Clinical Lymphoma and Myeloma*, **2008**, 8, 171-5		75
56	An overview of SLC3A1 and SLC7A9 mutations in Greek cystinuria patients. *Molecular Genetics and Metabolism*, **2008**, 95, 192-3	3.7	7
55	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	2.4	10
54	Estrogen receptor alpha gene polymorphisms and stroke. *Cerebrovascular Diseases*, **2008**, 26, 338	3.2	1
53	The synergistic effect of sex hormone-binding globulin and aromatase genes on polycystic ovary syndrome phenotype. *European Journal of Endocrinology*, **2008**, 158, 861-5	6.5	36
52	Estrogen receptor alpha gene haplotypes and diplotypes in the risk of stroke. *Stroke*, **2008**, 39, e172-3; author reply e174	6.7	7
51	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, 461-8	2.5	12
50	Distinct association of SLC19A1 polymorphism -43T>C with red cell folate levels and of MTHFR polymorphism 677C>T with plasma folate levels. *Clinical Biochemistry*, **2008**, 41, 174-6	3.5	9

49	The adiponectin-to-leptin ratio in women with polycystic ovary syndrome: relation to insulin resistance and proinflammatory markers. *Metabolism: Clinical and Experimental*, **2007**, 56, 766-71	12.7	35
48	Transcription regulatory polymorphism -43T>C in the 5'-flanking region of SLC19A1 gene could affect rheumatoid arthritis patient response to methotrexate therapy. *Rheumatology International*, **2007**, 27, 1057-61	3.6	35
47	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	4.7	24
46	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, 83-7	3.2	7
45	Mosaic trisomy r(14) associated with epilepsy and mental retardation. *Journal of Child Neurology*, **2007**, 22, 869-73	2.5	8
44	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, 181-9	3	22
43	Methylation status of the SNRPN and HUMARA genes in testicular biopsy samples. *Fertility and Sterility*, **2007**, 87, 805-9	4.8	12
42	Region Based Segmentation and Classification of Multispectral Chromosome Images. *Proceedings of the IEEE Symposium on Computer-Based Medical Systems*, **2007**,		6
41	Evidence for association of SLC7A9 gene haplotypes with cystinuria manifestation in SLC7A9 mutation carriers. *Urological Research*, **2006**, 34, 299-303		16
40	Absence of mutations of the EPO-receptor gene in Greek patients with familial polycythemia. *European Journal of Haematology*, **2006**, 76, 537-8	3.8	4
39	Identification of the four most common beta-globin gene mutations in Greek beta-thalassemic patients and carriers by PCR-SSCP: advantages and limitations of the method. *Journal of Clinical Laboratory Analysis*, **2006**, 20, 1-7	3	5
38	A watershed based segmentation method for multispectral chromosome images classification. *Annual International Conference of the IEEE Engineering in Medicine and Biology Society*, **2006**, 2006, 3009-12		18
37	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, 506-9	2.4	13
36	Delta-thalassemia in Cyprus. *Hemoglobin*, **2006**, 30, 455-62	0.6	23
35	Genetic and epigenetic risks of intracytoplasmic sperm injection method. *Asian Journal of Andrology*, **2006**, 8, 643-73	2.8	75
34	R831X mutation of the androgen receptor gene in an adolescent with complete androgen insensitivity syndrome and bilateral testicular hamartomata. *Hormones*, **2006**, 5, 200-4	3.1	9
33	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, 1354-62	4.8	20
32	Real-time PCR analysis of trinucleotide repeat allele expansions in the androgen receptor gene. *Molecular Diagnosis and Therapy*, **2005**, 9, 217-9		1

31	Effect of adiponectin gene polymorphisms on circulating adiponectin and insulin resistance indexes in women with polycystic ovary syndrome. *Clinical Chemistry*, **2005**, 51, 416-23	5.5	75
30	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, 175-84		12
29	Prenatal paternity testing using DNA extracted from coelomic cells. *Fetal Diagnosis and Therapy*, **2004**, 19, 75-7	2.4	10
28	Association of the T45G polymorphism in exon 2 of the adiponectin gene with polycystic ovary syndrome: role of Delta4-androstenedione. *Human Reproduction*, **2004**, 19, 1728-33	5.7	40
27	Glutathione sulfur transferase M1 and T1 genotypes in chronic lymphoblastic leukemia. *The Hematology Journal*, **2004**, 5, 500-4		4
26	A polymorphism in the resistin gene promoter is associated with body mass index in women with polycystic ovary syndrome. *Fertility and Sterility*, **2004**, 82, 1466-7	4.8	25
25	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
24	Interaction between the polymorphisms of the renin-angiotensin system in preeclampsia. *European Journal of Obstetrics, Gynecology and Reproductive Biology*, **2003**, 110, 8-11	2.4	57
23	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, 5976-80	5.6	102
22	Distribution of Two X-Linked Trinucleotide Polymorphisms in Greece. *Public Health Genomics*, **2001**, 4, 125-128		1
21	Preimplantation genetic diagnosis for spinal and bulbar muscular atrophy (SBMA). *Human Genetics*, **2001**, 108, 494-8	6.3	17
20	Expanded cytological referral criteria for colposcopy in cervical screening: comparison with human papillomavirus testing. *Gynecologic Oncology*, **2001**, 82, 355-9	4.9	10
19	Novel PKD1 deletions and missense variants in a cohort of Hellenic polycystic kidney disease families. *European Journal of Human Genetics*, **2001**, 9, 677-84	5.3	10
18	Congenital obstructive azoospermia in a man with Marfan syndrome. *Fertility and Sterility*, **2001**, 76, 1256-7	4.8	1
17	Glutathione S-transferase null genotypes in transitional cell bladder cancer: a case-control study. *European Urology*, **2000**, 37, 660-4	10.2	24
16	Arrayed primer extension: solid-phase four-color DNA resequencing and mutation detection technology. *Genetic Testing and Molecular Biomarkers*, **2000**, 4, 1-7		150
15	Effect of recombinant human erythropoietin in preterm infants. *European Journal of Haematology*, **1999**, 63, 71-6	3.8	13
14	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		326

13	Fragile X premutations and (TA)n estrogen receptor polymorphism in women with ovarian dysfunction **1999**, 84, 306-308		24
12	GENOTYPES OF N-ACETYLTRANSFERASE-2 AND RISK OF BLADDER CANCER: A CASE-CONTROL STUDY. *Journal of Urology*, **1999**, 161, 1672-1675	2.5	32
11	Association of estrogen receptor gene polymorphisms with endometriosis. *Fertility and Sterility*, **1999**, 72, 164-6	4.8	139
10	Fragile X premutations and (TA)n estrogen receptor polymorphism in women with ovarian dysfunction **1999**, 84, 306		1
9	FRAXA and FRAXE prevalence in patients with nonspecific mental retardation in the Hellenic population. *Genetic Epidemiology*, **1998**, 15, 103-9	2.6	23
8	The role of deoxyribonuclease I in amniotic fluid during pregnancy and labour. *European Journal of Obstetrics, Gynecology and Reproductive Biology*, **1998**, 77, 177-80	2.4	
7	Recombinant human erythropoietin treatment of postpartum anemia. Preliminary results. *European Journal of Obstetrics, Gynecology and Reproductive Biology*, **1998**, 81, 27-31	2.4	12
6	Infertility and multiple urogenital abnormalities in a male with mosaic 46,XY/45,XO/47,XXY karyotype and mixed phenotype. *Urologia Internationalis*, **1998**, 61, 111-4	1.9	7
5	Fetal Heart Rate Following Coelocentesis. *Journal of Maternal-Fetal and Neonatal Medicine*, **1997**, 6, 314-316	2	
4	Increased nuchal translucency thickness in a fetus at risk for beta-thalassaemia. *The Journal of Maternal-fetal Medicine*, **1997**, 6, 301-2		
3	Evidence for high-risk haplotypes and (CGG)n expansion in fragile X syndrome in the Hellenic population of Greece and Cyprus. *American Journal of Medical Genetics Part A*, **1996**, 64, 234-8		17
2	High HbF in pregnancy is associated with the Xmn I polymorphism at the -158bp of the G gamma-globin gene. *European Journal of Obstetrics, Gynecology and Reproductive Biology*, **1995**, 60, 153-6	2.4	11
1	A Watershed Based Segmentation Method for Multispectral Chromosome Images Classification		2