Christos Yapijakis

List of Publications by Year in Descending Order

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

120
papers

1,843
citations

24
h-index

39
g-index

123
ext. papers

2,021
avg, IF

L-index

#	Paper	IF	Citations
120	The Impact of and Gene Polymorphisms in Pulmonary Diseases Including COVID-19 <i>In Vivo</i> , 2022 , 36, 13-29	2.3	2
119	The Role of MicroRNAs in Thrombosis Advances in Experimental Medicine and Biology, 2021, 1339, 409	9-43 <u>l.€</u>	
118	Prenatal Genetic Testing for X-Linked Hypohidrotic Ectodermal Dysplasia <i>Advances in Experimental Medicine and Biology</i> , 2021 , 1339, 337-340	3.6	O
117	Craniofacial and Neurological Phenotype in a Case of Oculodentodigital Syndrome <i>Advances in Experimental Medicine and Biology</i> , 2021 , 1339, 325-329	3.6	
116	Phenotype and Genotype Study in a Case of Frontometaphyseal Dysplasia 1 Advances in Experimental Medicine and Biology, 2021 , 1339, 319-323	3.6	
115	Clinical and Molecular Study of Common Thrombophilia Mutation Prothrombin G20210A <i>Advances in Experimental Medicine and Biology</i> , 2021 , 1339, 331-336	3.6	
114	Preimplantation Genetic Testing for Spastic Paraplegia Type 3 <i>Advances in Experimental Medicine and Biology</i> , 2021 , 1339, 341-345	3.6	O
113	Hot cross bun sign and prominent cerebellar peduncle involvement in a patient with oculodentodigital dysplasia. <i>Neurological Sciences</i> , 2021 , 42, 343-345	3.5	1
112	Association study indicates combined effect of interleukin-10 and angiotensin-converting enzyme in basal cell carcinoma development. <i>Archives of Dermatological Research</i> , 2021 , 313, 373-380	3.3	1
111	The Angiotensin-converting Enzyme Insertion/Deletion Polymorphism as a Common Risk Factor for Major Pregnancy Complications. <i>In Vivo</i> , 2021 , 35, 95-103	2.3	4
110	Increased Incidence of Stress-related Tic Habit Cough in Children During the Recent Greek Financial Crisis. <i>In Vivo</i> , 2021 , 35, 1811-1820	2.3	1
109	Lack of Association Between the G8790A Gene Variation and Risk for Basal Cell Carcinoma. <i>Anticancer Research</i> , 2021 , 41, 4021-4026	2.3	1
108	Craniofacial and Neurological Phenotype in a Patient with De Novo 18q Microdeletion and 18p Microduplication. <i>Advances in Experimental Medicine and Biology</i> , 2020 , 1195, 163-166	3.6	1
107	Genetic Counseling for Adult-Onset Spinal and Bulbar Muscular Atrophy (Kennedy Syndrome): Multiple Cases of Prenatal Testing in a Family. <i>Advances in Experimental Medicine and Biology</i> , 2020 , 1195, 199-204	3.6	3
106	Regulatory Role of MicroRNAs in Brain Development and Function. <i>Advances in Experimental Medicine and Biology</i> , 2020 , 1195, 237-247	3.6	8
105	History of Hygiene Focusing on the Crucial Role of Water in the Hellenic Asclepieia (i.e., Ancient Hospitals). <i>Water (Switzerland)</i> , 2020 , 12, 754	3	6
104	The Hamster Model of Sequential Oral Carcinogenesis: An Update. <i>In Vivo</i> , 2019 , 33, 1751-1755	2.3	6

(2015-2019)

103	Association of Polymorphisms in the Genes of Angiotensinogen and Angiotensin Receptors With Risk for Basal Cell Carcinoma. <i>Anticancer Research</i> , 2019 , 39, 5525-5530	2.3	4
102	Gene Variant Causing High Blood Pressure May Be Associated With Medication-related Jaw Osteonecrosis. <i>In Vivo</i> , 2019 , 33, 559-562	2.3	2
101	An unusual phenocopy of the HANAC syndrome without genetic involvement of COL4A1/COL4A2. <i>Acta Neurologica Belgica</i> , 2018 , 118, 135-136	1.5	
100	Recognition of Cerebral Autosomal Dominant Arteriopathy with Subcortical Infarcts and Leukoencephalopathy (CADASIL) in Two Oligosymptomatic Sisters with Low CADASIL Scale Scores and a Venous Dysplasia: Report of a Novel Greek Family. <i>Journal of Stroke and Cerebrovascular</i>	2.8	Ο
99	Ancestral Concepts of Human Genetics and Molecular Medicine in Epicurean Philosophy 2017 , 41-57		1
98	History of sanitation and hygiene technologies in the Hellenic world. <i>Journal of Water Sanitation and Hygiene for Development</i> , 2017 , 7, 163-180	1.5	7
97	Huntington Disease: Genetics, Prevention, and Therapy Approaches. <i>Advances in Experimental Medicine and Biology</i> , 2017 , 987, 55-65	3.6	4
96	Neurofibromatosis-Noonan Syndrome: A Possible Paradigm of the Combination of Genetic and Epigenetic Factors. <i>Advances in Experimental Medicine and Biology</i> , 2017 , 987, 151-159	3.6	4
95	Cerebral Thrombosis: A Neurogenetic Approach. <i>Advances in Experimental Medicine and Biology</i> , 2017 , 987, 13-21	3.6	3
94	Novel PANK2 mutation in the first Greek compound heterozygote patient with pantothenate-kinase-associated neurodegeneration. <i>SAGE Open Medical Case Reports</i> , 2017 , 5, 205031.	3X777	20101
93	Cerebral autosomal dominant arteriopathy with subcortical infarcts and leucoancepahlopathy presenting with postpartum psychosis and late-onset stroke. <i>Future Neurology</i> , 2016 , 11, 207-213	1.5	1
92	Mutation Screening of Her-2, N-ras and Nf1 Genes in Brain Tumor Biopsies. <i>Anticancer Research</i> , 2016 , 36, 4607-11	2.3	6
91	Effect of Olmesartan on the Level of Oral Cancer Risk Factor PAI1. Anticancer Research, 2016, 36, 6093-	·6 <u>0</u> .96	2
90	Mutations of Two Major Coagulation Factors Are Not Associated with Male Infertility. <i>In Vivo</i> , 2016 , 30, 927-930	2.3	3
89	Is Neurofibromatosis Type 1-Noonan Syndrome a Phenotypic Result of Combined Genetic and Epigenetic Factors?. <i>In Vivo</i> , 2016 , 30, 315-20	2.3	4
88	High frequency of TTTY2-like gene-related deletions in patients with idiopathic oligozoospermia and azoospermia. <i>Andrologia</i> , 2015 , 47, 536-44	2.4	1
87	Common Gene Polymorphisms Associated with Thrombophilia 2015,		1
86	Cerebrovascular Aneurysms May Be Associated with Thrombophilia-predisposing Mutations in Patients with Familial Risk. <i>In Vivo</i> , 2015 , 29, 395-8	2.3	3

85	Juvenile myoclonic epilepsy is not associated with the DRPLA gene in a European population. <i>In Vivo</i> , 2014 , 28, 1193-6	2.3	1
84	Prevalence of human papillomavirus in saliva and cervix of sexually active women. <i>Gynecologic Oncology</i> , 2013 , 129, 395-400	4.9	21
83	Association of angiotensin-converting enzyme gene insertion/deletion polymorphism with decreased risk for basal cell carcinoma. <i>Archives of Dermatological Research</i> , 2013 , 305, 333-9	3.3	8
82	Effect of thrombosis-related gene polymorphisms upon oral cancer: a regression analysis. <i>Anticancer Research</i> , 2013 , 33, 4033-9	2.3	14
81	Potential prevention of thromboembolism by genetic counseling and testing for two common thrombophilia mutations. <i>In Vivo</i> , 2012 , 26, 165-72	2.3	5
80	The interplay between hemostasis and malignancy: the oral cancer paradigm. <i>Anticancer Research</i> , 2012 , 32, 1791-800	2.3	12
79	Oral carcinogenesis is not achieved in different carcinogen-treated PAI-1 transgenic and wild-type mouse models. <i>In Vivo</i> , 2012 , 26, 1001-5	2.3	
78	Prevalence of thrombosis-related DNA polymorphisms in a healthy Greek population. <i>In Vivo</i> , 2012 , 26, 1095-101	2.3	6
77	Homocysteinemia-associated anetoderma, in a young woman with anorexia nervosa history. <i>International Journal of Dermatology</i> , 2011 , 50, 343-5	1.7	3
76	Screening for Familial Mediterranean Fever M694V and V726A mutations in the Greek population. <i>Genetic Testing and Molecular Biomarkers</i> , 2009 , 13, 291-3	1.6	3
75	Gene polymorphisms related to angiogenesis, inflammation and thrombosis that influence risk for oral cancer. <i>Oral Oncology</i> , 2009 , 45, 247-53	4.4	46
74	Bisphosphonate-induced avascular osteonecrosis of the mandible associated with a common thrombophilic mutation in the prothrombin gene. <i>Journal of Oral and Maxillofacial Surgery</i> , 2009 , 67, 2009-12	1.8	12
73	Association of leptin -2548G/A and leptin receptor Q223R polymorphisms with increased risk for oral cancer. <i>Journal of Cancer Research and Clinical Oncology</i> , 2009 , 135, 603-12	4.9	31
72	Association of polymorphisms in Tumor Necrosis Factor Alpha and Beta genes with increased risk for oral cancer. <i>Anticancer Research</i> , 2009 , 29, 2379-86	2.3	28
71	Hippocrates of Kos, the father of clinical medicine, and Asclepiades of Bithynia, the father of molecular medicine. Review. <i>In Vivo</i> , 2009 , 23, 507-14	2.3	45
70	A common 9 bp deletion in the ataxia-telangiectasia-mutated gene is not associated with oral cancer. <i>Anticancer Research</i> , 2009 , 29, 3191-3	2.3	2
69	Comparison of cytology, colposcopy, HPV typing and biomarker analysis in cervical neoplasia. <i>Anticancer Research</i> , 2009 , 29, 3401-9	2.3	10
68	The hamster model of sequential oral oncogenesis. <i>Oral Oncology</i> , 2008 , 44, 315-24	4.4	77

(2007-2008)

67	Genetic association of cytokine DNA polymorphisms with head and neck cancer. <i>Oral Oncology</i> , 2008 , 44, 1093-9	4.4	46
66	Diabetes enhances cell proliferation but not Bax/Bcl-2-mediated apoptosis during oral oncogenesis. <i>International Journal of Oral and Maxillofacial Surgery</i> , 2008 , 37, 60-5	2.9	7
65	A metalloproteinase-9 polymorphism which affects its expression is associated with increased risk for oral squamous cell carcinoma. <i>European Journal of Surgical Oncology</i> , 2008 , 34, 450-5	3.6	42
64	Enhancement of erbB2 and erbB3 expression during oral oncogenesis in diabetic rats. <i>Journal of Cancer Research and Clinical Oncology</i> , 2008 , 134, 337-44	4.9	4
63	Gene expression polymorphisms of interleukins-1 beta, -4, -6, -8, -10, and tumor necrosis factors-alpha, -beta: regression analysis of their effect upon oral squamous cell carcinoma. <i>Journal of Cancer Research and Clinical Oncology</i> , 2008 , 134, 821-32	4.9	57
62	Typhoid Fever Epidemic in Ancient Athens 2008 , 161-173		5
61	A DNA polymorphism of stromal-derived factor-1 is associated with advanced stages of oral cancer. <i>Anticancer Research</i> , 2008 , 28, 271-5	2.3	24
60	The interleukin-10 (-1082A/G) polymorphism is strongly associated with increased risk for oral squamous cell carcinoma. <i>Anticancer Research</i> , 2008 , 28, 309-14	2.3	37
59	Angiotensinogen polymorphism is associated with risk for malignancy but not for oral cancer. <i>Anticancer Research</i> , 2008 , 28, 1675-9	2.3	7
58	Prevalence of HPV types in a cohort of Greeks with clinical indication of infection. <i>Anticancer Research</i> , 2008 , 28, 2233-7	2.3	5
57	HPV detection rate in saliva may depend on the immune system efficiency. <i>In Vivo</i> , 2008 , 22, 599-602	2.3	19
56	H-ras and c-fos exhibit similar expression patterns during most stages of oral oncogenesis. <i>In Vivo</i> , 2008 , 22, 621-8	2.3	9
55	Increased risk of oral cancer in diabetic animals is not associated with c-jun activation pathway. <i>Journal of Cranio-Maxillo-Facial Surgery</i> , 2007 , 35, 382-7	3.6	1
54	The 1040C/T polymorphism influencing thermal stability and activity of thrombin activatable fibrinolysis inhibitor is associated with risk for oral cancer. <i>American Journal of Hematology</i> , 2007 , 82, 1010-2	7.1	9
53	Ancient typhoid epidemic reveals possible ancestral strain of Salmonella enterica serovar Typhi. <i>Infection, Genetics and Evolution</i> , 2007 , 7, 126-7	4.5	14
52	Diabetes may increase risk for oral cancer through the insulin receptor substrate-1 and focal adhesion kinase pathway. <i>Oral Oncology</i> , 2007 , 43, 165-73	4.4	23
51	Expression of ets-1 is not affected by N-ras or H-ras during oral oncogenesis. <i>Journal of Cancer Research and Clinical Oncology</i> , 2007 , 133, 227-33	4.9	9
50	The low VEGF production allele of the +936C/T polymorphism is strongly associated with increased risk for oral cancer. <i>Journal of Cancer Research and Clinical Oncology</i> , 2007 , 133, 787-91	4.9	45

49	Association of angiotensin-converting enzyme gene insertion/deletion polymorphism with increased risk for oral cancer. <i>Acta Oncolgica</i> , 2007 , 46, 1097-102	3.2	24
48	Increased risk for oral cancer is associated with coagulation factor XIII but not with factor XII. Oncology Reports, 2007 , 18, 1537	3.5	5
47	Strong association of the tissue inhibitor of metalloproteinase-2 polymorphism with an increased risk of oral squamous cell carcinoma in Europeans. <i>Oncology Reports</i> , 2007 ,	3.5	1
46	Strong association of interleukin-4 (-590 C/T) polymorphism with increased risk for oral squamous cell carcinoma in Europeans. <i>Oral Surgery Oral Medicine Oral Pathology Oral Radiology and Endodontics</i> , 2007 , 104, 796-802		21
45	The interleukin-8 (-251A/T) polymorphism is associated with increased risk for oral squamous cell carcinoma. <i>European Journal of Surgical Oncology</i> , 2007 , 33, 504-7	3.6	54
44	Strong association of the tissue inhibitor of metalloproteinase-2 polymorphism with an increased risk of oral squamous cell carcinoma in Europeans. <i>Oncology Reports</i> , 2007 , 17, 963-8	3.5	30
43	Association of matrix metalloproteinase-1 (-1607 1G/2G) polymorphism with increased risk for oral squamous cell carcinoma. <i>Anticancer Research</i> , 2007 , 27, 459-64	2.3	27
42	Loss of tumour suppressor p16 expression in initial stages of oral oncogenesis. <i>Anticancer Research</i> , 2007 , 27, 979-84	2.3	19
41	Diabetes alters expression of p53 and c-myc in different stages of oral oncogenesis. <i>Anticancer Research</i> , 2007 , 27, 1465-73	2.3	8
40	Coagulation-related factors, thrombomodulin and protein Z, are not associated with risk for oral cancer. <i>Anticancer Research</i> , 2007 , 27, 2449-51	2.3	1
39	High gene expression of matrix metalloproteinase-7 is associated with early stages of oral cancer. <i>Anticancer Research</i> , 2007 , 27, 2493-8	2.3	29
38	Diabetes increases both N-ras and ets-1 expression during rat oral oncogenesis resulting in enhanced cell proliferation and metastatic potential. <i>In Vivo</i> , 2007 , 21, 615-21	2.3	5
37	Diabetes does not influence oral oncogenesis through fibroblast growth factor receptors. <i>In Vivo</i> , 2007 , 21, 623-8	2.3	
36	Increased risk for oral cancer is associated with coagulation factor XIII but not with factor XII. <i>Oncology Reports</i> , 2007 , 18, 1537-43	3.5	14
35	Expression of cell cycle regulator p16 is not affected by diabetes during oral oncogenesis. <i>In Vivo</i> , 2007 , 21, 745-50	2.3	1
34	EGFR and c-Jun exhibit the same pattern of expression and increase gradually during the progress of oral oncogenesis. <i>In Vivo</i> , 2007 , 21, 791-6	2.3	3
33	Evaluation of apoptosis in nasal and buccal cells of septic patients. <i>In Vivo</i> , 2007 , 21, 901-4	2.3	1
32	The interleukin-1 beta gene polymorphism +3953 C/T is not associated with risk for oral cancer. Anticancer Research, 2007, 27, 3981-6	2.3	7

(2005-2007)

31	The interleukin-18 -607A/C polymorphism is not associated with risk for oral cancer. <i>Anticancer Research</i> , 2007 , 27, 4011-4	2.3	17
30	A metalloproteinase-13 polymorphism affecting its gene expression is associated with advanced stages of oral cancer. <i>Anticancer Research</i> , 2007 , 27, 4027-30	2.3	10
29	Association of -1171 promoter polymorphism of matrix metalloproteinase-3 with increased risk for oral cancer. <i>Anticancer Research</i> , 2007 , 27, 4095-100	2.3	12
28	The platelet glycoprotein Ibalpha VNTR polymorphism is associated with risk for oral cancer. <i>Anticancer Research</i> , 2007 , 27, 4121-5	2.3	2
27	Diabetes and oral oncogenesis. <i>Anticancer Research</i> , 2007 , 27, 4185-93	2.3	24
26	Cell proliferation and apoptosis culminate in early stages of oral oncogenesis. <i>Oral Oncology</i> , 2006 , 42, 540-50	4.4	25
25	Plasminogen activator inhibitor-1 polymorphism is associated with increased risk for oral cancer. <i>Oral Oncology</i> , 2006 , 42, 888-92	4.4	29
24	DNA examination of ancient dental pulp incriminates typhoid fever as a probable cause of the Plague of Athens. <i>International Journal of Infectious Diseases</i> , 2006 , 10, 206-14	10.5	122
23	Insufficient phylogenetic analysis may not exclude candidacy of typhoid fever as a probable cause of the Plague of Athens (reply to Shapiro et al.). <i>International Journal of Infectious Diseases</i> , 2006 , 10, 335-336	10.5	8
22	Association of platelet glycoprotein Ia polymorphism with minor increase of risk for oral cancer. <i>European Journal of Surgical Oncology</i> , 2006 , 32, 455-7	3.6	18
21	Immunological and molecular detection of human immunodeficiency virus in saliva, and comparison with blood testing. <i>European Journal of Oral Sciences</i> , 2006 , 114, 175-9	2.3	7
20	Methylenetetrahydrofolate reductase polymorphism and minor increase of risk for oral cancer. Journal of Cancer Research and Clinical Oncology, 2006 , 132, 219-22	4.9	41
19	Strong association of interleukin-6 -174 G>C promoter polymorphism with increased risk of oral cancer. <i>International Journal of Biological Markers</i> , 2006 , 21, 246-50	2.8	21
18	The co-expression of c-myc and p53 increases and reaches a plateau early in oral oncogenesis. <i>Anticancer Research</i> , 2006 , 26, 2957-62	2.3	14
17	FGFR-2 and -3 play an important role in initial stages of oral oncogenesis. <i>Anticancer Research</i> , 2006 , 26, 4217-21	2.3	14
16	Real-time PCR analysis of trinucleotide repeat allele expansions in the androgen receptor gene. <i>Molecular Diagnosis and Therapy</i> , 2005 , 9, 217-9		1
15	Salivary gland manifestations of sarcoidosis: report of three cases. <i>Journal of Oral and Maxillofacial Surgery</i> , 2005 , 63, 1016-21	1.8	14
14	Are factor V and prothrombin mutations associated with increased risk of oral cancer?. <i>Anticancer Research</i> , 2005 , 25, 2561-5	2.3	22

13	Use of truncated pyramid representation methodology in three-dimensional reconstruction: an example. <i>Journal of Microscopy</i> , 2004 , 214, 70-5	1.9	8
12	Thyroid gland neurofibroma in a NF1 patient. Acta Neurologica Scandinavica, 2002, 106, 58-61	3.8	6
11	Association of oestrogen receptor alpha polymorphisms and androgen receptor CAG trinucleotide repeats with male infertility: a study in 109 Greek infertile men. <i>Journal of Developmental and Physical Disabilities</i> , 2002 , 25, 149-52		66
10	A simple and effective approach for detecting maternal cell contamination in molecular prenatal diagnosis. <i>Prenatal Diagnosis</i> , 2002 , 22, 425-9	3.2	24
9	Prenatal testing for Huntington@disease: a European collaborative study. <i>European Journal of Human Genetics</i> , 2002 , 10, 689-93	5.3	40
8	Predictive DNA-testing for Huntington@disease and reproductive decision making: a European collaborative study. <i>European Journal of Human Genetics</i> , 2002 , 10, 167-76	5.3	68
7	Clinical and genetic heterogeneity in benign hereditary chorea. <i>Neurology</i> , 2002 , 59, 579-84	6.5	89
6	Distribution of Two X-Linked Trinucleotide Polymorphisms in Greece. <i>Public Health Genomics</i> , 2001 , 4, 125-128		1
5	Prenatal diagnosis of X-linked spinal and bulbar muscular atrophy in a Greek family. <i>Prenatal Diagnosis</i> , 1996 , 16, 262-5	3.2	12
4	Hereditary neuropathy with liability to pressure palsies: the same molecular defect can result in diverse clinical presentation. <i>Journal of Neurology</i> , 1996 , 243, 225-30	5.5	11
3	Exclusion mapping of the benign hereditary chorea gene from the Huntington@ disease locus: report of a family. <i>Clinical Genetics</i> , 1995 , 47, 133-8	4	3
2	Linkage disequilibrium between the expanded (CAG)n repeat and an allele of the adjacent (CCG)n repeat in Huntington@ disease patients of Greek origin. <i>European Journal of Human Genetics</i> , 1995 , 3, 228-34	5.3	14
1	Isolation of temperature-sensitive mutants of 16 S rRNA in Escherichia coli. <i>Journal of Molecular</i>	6.5	80