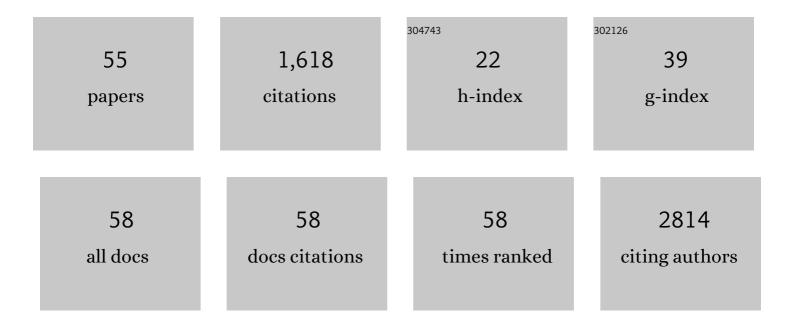
Nikolaos Drakoulis

List of Publications by Year in descending order

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Version: 2024-02-01



#	Article	lF	CITATIONS
1	VDR Gene Polymorphisms and Cluster Headache Susceptibility: Case–Control Study in a Southeastern European Caucasian Population. Journal of Molecular Neuroscience, 2022, 72, 382-392.	2.3	8
2	Deciphering the Role of the rs2651899, rs10166942, and rs11172113 Polymorphisms in Migraine: A Meta-Analysis. Medicina (Lithuania), 2022, 58, 491.	2.0	6
3	The Greek Collaborative Long COVID Study: Non-Hospitalized and Hospitalized Patients Share Similar Symptom Patterns. Journal of Personalized Medicine, 2022, 12, 987.	2.5	9
4	Back to basics in COVIDâ€19: Antigens and antibodies—Completing the puzzle. Journal of Cellular and Molecular Medicine, 2021, 25, 4523-4533.	3.6	35
5	Evidence That HFE H63D Variant Is a Potential Disease Modifier in Cluster Headache. Journal of Molecular Neuroscience, 2021, , 1.	2.3	1
6	Replication of chromosomal loci involved in Parkinson's disease: A quantitative synthesis of GWAS. Toxicology Reports, 2021, 8, 1762-1768.	3.3	0
7	Analysis of HCRTR2, GNB3, and ADH4 Gene Polymorphisms in a Southeastern European Caucasian Cluster Headache Population. Journal of Molecular Neuroscience, 2020, 70, 467-474.	2.3	12
8	Positron Emission Tomography in Merkel Cell Carcinoma. Cancers, 2020, 12, 2897.	3.7	9
9	A new threat from an old enemy: Reâ€'emergence of coronavirus (Review). International Journal of Molecular Medicine, 2020, 45, 1631-1643.	4.0	175
10	CYP1A2 rs762551 and ADORA2A rs5760423 Polymorphisms in Patients with Blepharospasm. Journal of Molecular Neuroscience, 2020, 70, 1370-1375.	2.3	12
11	Does the CD33 rs3865444 Polymorphism Confer Susceptibility to Alzheimer's Disease?. Journal of Molecular Neuroscience, 2020, 70, 851-860.	2.3	23
12	ANTIBIOTIC RESISTANCE IN COMMUNITY-ACQUIRED PNEUMONIA. A ROMANIAN PERSPECTIVE. Farmacia, 2020, 68, 512-520.	0.4	4
13	Comprehensive analysis of drugs to treat SARS‑CoV‑2 infection: Mechanistic insights into current COVID‑19 therapies (Review). International Journal of Molecular Medicine, 2020, 46, 467-488.	4.0	116
14	Unraveling the roles of vitaminÂD status and melanin during Covid‑19 (Review). International Journal of Molecular Medicine, 2020, 47, 92-100.	4.0	18
15	Clinical significance of serum melatonin in predicting the severity of oral squamous cell carcinoma. Oncology Letters, 2020, 19, 1537-1543.	1.8	18
16	Translational Application of Circulating DNA in Oncology: Review of the Last Decades Achievements. Cells, 2019, 8, 1251.	4.1	53
17	Targeted Metabolomic Analysis of Serum Fatty Acids for the Prediction of Autoimmune Diseases. Frontiers in Molecular Biosciences, 2019, 6, 120.	3.5	71
18	A Mechanistic and Pathophysiological Approach for Stroke Associated with Drugs of Abuse. Journal of Clinical Medicine, 2019, 8, 1295.	2.4	89

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#	Article	IF	CITATIONS
19	Hemochromatosis: Hereditary hemochromatosis and HFE gene. Vitamins and Hormones, 2019, 110, 201-222.	1.7	25
20	Contribution of MTHFR gene variants in lupus related subclinical atherosclerosis. Clinical Immunology, 2018, 193, 110-117.	3.2	25
21	Assessment of telomerase activity in leukocytes of typeÃ ⁻ ¿½2 diabetes mellitus patients having or not foot ulcer: Possible correlation with other clinical parameters. Experimental and Therapeutic Medicine, 2018, 15, 3420-3424.	1.8	12
22	Genetically driven antioxidant capacity in a Caucasian Southeastern European population. Mechanisms of Ageing and Development, 2018, 172, 1-5.	4.6	3
23	Beta 1, Beta 2 and Beta 3 Adrenergic Receptor Gene Polymorphisms in a Southeastern European Population. Frontiers in Genetics, 2018, 9, 560.	2.3	21
24	Population-Based Analysis of Cluster Headache-Associated Genetic Polymorphisms. Journal of Molecular Neuroscience, 2018, 65, 367-376.	2.3	11
25	Antimicrobial resistance development following surgical site infections. Molecular Medicine Reports, 2017, 15, 681-688.	2.4	49
26	CYP polymorphisms and pathological conditions related to chronic exposure to organochlorine pesticides. Toxicology Reports, 2017, 4, 335-341.	3.3	63
27	Pharmacological management of non-alcoholic fatty liver disease: Atorvastatin versus pentoxifylline. Experimental and Therapeutic Medicine, 2017, 13, 2375-2381.	1.8	34
28	Application of metabolomics: Focus on the quantification of organic acids in healthy adults. International Journal of Molecular Medicine, 2017, 40, 112-120.	4.0	32
29	Environmental exposure to organophosphorus nerve agents. Environmental Toxicology and Pharmacology, 2017, 56, 163-171.	4.0	43
30	Neuroendocrine factors: The missing link in non-melanoma skin cancer. Oncology Reports, 2017, 38, 1327-1340.	2.6	55
31	MTHFR gene variants and non-MALT lymphoma development in primary Sjogren's syndrome. Scientific Reports, 2017, 7, 7354.	3.3	28
32	Effect of singleâ€nucleotide polymorphisms in <i><scp>ADH</scp>1B</i> , <i><scp>ADH</scp>1C</i> , <i><scp>OPRM</scp>1</i> ,< and <i><scp>ALDH</scp>2</i> genes on alcohol dependence in a Caucasian population. Pharmacology Research and Perspectives, 2017, 5, e00326.	i> <scp>DR</scp>	RD2
33	Colon Cancer Cells Gene Expression Signature As Response to 5- Fluorouracil, Oxaliplatin, and Folinic Acid Treatment. Frontiers in Pharmacology, 2016, 7, 172.	3.5	33
34	Preliminary SEM Observations on the Surface of Elastomeric Impression Materials after Immersion or Ozone Disinfection. Journal of Clinical and Diagnostic Research JCDR, 2016, 10, ZC01-ZC05.	0.8	6
35	Population-based analysis of the frequency of HFE gene polymorphisms: Correlation with the susceptibility to develop hereditary hemochromatosis. Molecular Medicine Reports, 2016, 14, 630-636.	2.4	21
36	HPV strain distribution in patients with genital warts in a female population sample. Oncology Letters, 2016, 12, 1779-1782.	1.8	37

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37	The Role of Glutamine Supplementation in Thoracic and Upper Aerodigestive Malignancies. Nutrition and Cancer, 2015, 67, 231-237.	2.0	11
38	Genes and personality characteristics: Possible association of the genetic background with intelligence and decision making in 830 Caucasian Greek subjects. Meta Gene, 2014, 2, 844-853.	0.6	3
39	Resistin in mid-trimester amniotic fluid in trisomy 21. Journal of Maternal-Fetal and Neonatal Medicine, 2013, 26, 1576-1580.	1.5	3
40	Fetuin A Concentration in the Second Trimester Amniotic Fluid of Fetuses with Trisomy 21 Appears to Be Lower: Phenotypic Considerations. Mediators of Inflammation, 2012, 2012, 1-4.	3.0	8
41	Frequency Distribution of COMT Polymorphisms in Greek Patients with Schizophrenia and Controls: A Study of SNPs rs737865, rs4680, and rs165599. , 2012, 2012, 1-6.		7
42	The multi-cancer marker, rs6983267, located at region 3 of chromosome 8q24, is associated with prostate cancer in Greek patients but does not contribute to the aggressiveness of the disease. Clinical Chemistry and Laboratory Medicine, 2012, 50, 379-85.	2.3	13
43	Safety and Pharmacokinetics of Oseltamivir for Prophylaxis of Neonates Exposed to Influenza H1N1. Pediatric Infectious Disease Journal, 2012, 31, 527-529.	2.0	20
44	Association of the eNOS E298D polymorphism and the risk of myocardial infarction in the Greek population. BMC Medical Genetics, 2010, 11, 133.	2.1	13
45	Spontaneous preterm birth in women with chronic hepatitis B virus infection. International Journal of Gynecology and Obstetrics, 2010, 110, 241-244.	2.3	29
46	Role of the H1 haplotype of microtubule-associated protein tau (MAPT) gene in Greek patients with Parkinson's disease. BMC Neurology, 2009, 9, 26.	1.8	15
47	Kinetics of progenitor hemopoetic stem cells in sepsis: Correlation with patients survival?. BMC Infectious Diseases, 2006, 6, 142.	2.9	18
48	Herpes simplex virus type 2: a cause of acute retinal necrosis syndrome. Ocular Immunology and Inflammation, 2001, 9, 103-109.	1.8	17
49	Polymorphisms in the Human Cytochrome P-450 1A1 Gene <i>(CYP1A1) </i> as a Factor for Developing Acne. Dermatology, 1998, 196, 171-175.	2.1	55
50	Retinal Perivasculitis in an Immunocompetent Patient With Systemic Herpes Simplex Infection. American Journal of Ophthalmology, 1997, 123, 699-702.	3.3	12
51	Polymorphisms in the human CYP1A1 gene as susceptibility factors for lung cancer: exon-7 mutation (4889 A to G), and a T to C mutation in the 3?-flanking region. The Clinical Investigator, 1994, 72, 240-8.	0.6	92
52	Polymorphic arylamine N-acetyltransferase (NAT2) genes in children with insulin-dependent diabetes mellitus. Clinical Pharmacology and Therapeutics, 1994, 56, 626-634.	4.7	23
53	Correlation between trans-stilbene oxide-glutathione conjugation activity and the deletion mutation in the glutathione S-transferase class Mu gene detected by polymerase chain reaction. Biochemical Pharmacology, 1992, 43, 647-650.	4.4	88
54	Mutant genes of cytochrome P-450IID6, glutathione S-transferase class Mu, and arylamine N-acetyltransferase in lung cancer patients. The Clinical Investigator, 1992, 70-70, 307-19.	0.6	16

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55	Debrisoquine hydroxylation polymorphism: comparison of debrisoquine- and dextromethorphan-test and restriction fragment analysis in healthy volunteers. European Journal of Pharmacology, 1990, 183, 2399.	3.5	0