

# Fotis Tsetsos

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/249542/publications.pdf>

Version: 2024-02-01

25  
papers

1,883  
citations

758635

12  
h-index

580395

25  
g-index

33  
all docs

33  
docs citations

33  
times ranked

5255  
citing authors

#	ARTICLE	IF	CITATIONS
1	Analysis of shared heritability in common disorders of the brain. <i>Science</i> , 2018, 360, .	6.0	1,085
2	Interrogating the Genetic Determinants of Touretteâ€™s Syndrome and Other Tic Disorders Through Genome-Wide Association Studies. <i>American Journal of Psychiatry</i> , 2019, 176, 217-227.	4.0	242
3	Rare Copy Number Variants in NRXN1 and CNTN6 Increase Risk for Tourette Syndrome. <i>Neuron</i> , 2017, 94, 1101-1111.e7.	3.8	137
4	Maritime route of colonization of Europe. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2014, 111, 9211-9216.	3.3	71
5	Genetic association signal near <i>NTN4</i> in Tourette syndrome. <i>Annals of Neurology</i> , 2014, 76, 310-315.	2.8	53
6	Investigating Shared Genetic Basis Across Tourette Syndrome and Comorbid Neurodevelopmental Disorders Along the Impulsivity-Compulsivity Spectrum. <i>Biological Psychiatry</i> , 2021, 90, 317-327.	0.7	49
7	Synaptic processes and immune-related pathways implicated in Tourette syndrome. <i>Translational Psychiatry</i> , 2021, 11, 56.	2.4	31
8	Targeted Re-Sequencing Approach of Candidate Genes Implicates Rare Potentially Functional Variants in Tourette Syndrome Etiology. <i>Frontiers in Neuroscience</i> , 2016, 10, 428.	1.4	29
9	Meta-Analysis of Tourette Syndrome and Attention Deficit Hyperactivity Disorder Provides Support for a Shared Genetic Basis. <i>Frontiers in Neuroscience</i> , 2016, 10, 340.	1.4	26
10	Concordance of genetic variation that increases risk for Tourette Syndrome and that influences its underlying neurocircuitry. <i>Translational Psychiatry</i> , 2019, 9, 120.	2.4	24
11	Genetics of the peloponnesean populations and the theory of extinction of the medieval peloponnesean Greeks. <i>European Journal of Human Genetics</i> , 2017, 25, 637-645.	1.4	22
12	Association of ALOX12 gene polymorphism with all-cause and cardiovascular mortality in diabetic nephropathy. <i>International Urology and Nephrology</i> , 2018, 50, 321-329.	0.6	20
13	Oxidative Stress Genes in Diabetes Mellitus Type 2: Association with Diabetic Kidney Disease. <i>Oxidative Medicine and Cellular Longevity</i> , 2021, 2021, 1-10.	1.9	15
14	Association of rs11780592 Polymorphism in the Human Soluble Epoxide Hydrolase Gene (EPHX2) with Oxidized LDL and Mortality in Patients with Diabetic Chronic Kidney Disease. <i>Oxidative Medicine and Cellular Longevity</i> , 2021, 2021, 1-8.	1.9	13
15	Elevated common variant genetic risk for tourette syndrome in a densely-affected pedigree. <i>Molecular Psychiatry</i> , 2021, 26, 7522-7529.	4.1	8
16	The Genetics of Gilles de la Tourette Syndrome: a Common Aetiological Basis with Comorbid Disorders?. <i>Current Behavioral Neuroscience Reports</i> , 2016, 3, 218-231.	0.6	7
17	Assessment of association between lipoxigenase genes variants in elderly Greek population and type 2 diabetes mellitus. <i>Diabetes and Vascular Disease Research</i> , 2018, 15, 340-343.	0.9	7
18	Association between CUBN gene variants, type 2 diabetes and vitamin D concentrations in an elderly Greek population. <i>Journal of Steroid Biochemistry and Molecular Biology</i> , 2020, 198, 105549.	1.2	7

#	ARTICLE	IF	CITATIONS
19	Genetic variation in CARD8, a gene coding for an NLRP3 inflammasome-associated protein, alters the genetic risk for diabetic nephropathy in the context of type 2 diabetes mellitus. <i>Diabetes and Vascular Disease Research</i> , 2020, 17, 147916412097089.	0.9	6
20	Investigation of SNP rs2060546 Immediately Upstream to NTN4 in a Danish Gilles de la Tourette Syndrome Cohort. <i>Frontiers in Neuroscience</i> , 2016, 10, 531.	1.4	5
21	Myasthenia gravis genome-wide association study implicates AGRN as a risk locus. <i>Journal of Medical Genetics</i> , 2022, 59, 801-809.	1.5	5
22	Genetic history of the population of Crete. <i>Annals of Human Genetics</i> , 2019, 83, 373-388.	0.3	2
23	Variants in clock genes could be associated with lower risk of type 2 diabetes in an elderly Greek population. <i>Maturitas</i> , 2021, 152, 20-25.	1.0	1
24	Genetics and Population Analysis. , 2019, , 363-378.		0
25	Outcomes and issues of 12 chordomas treated in a single center. <i>Egyptian Journal of Neurology, Psychiatry and Neurosurgery</i> , 2021, 57, .	0.4	0