Kyriacos Markianos

List of Publications by Year in descending order

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430442 433756 2,092 33 18 31 citations h-index g-index papers 36 36 36 3304 docs citations times ranked citing authors all docs

#	Article	IF	CITATIONS
1	Congenital Xâ€linked Neutropenia with Myelodysplasia and Somatic Tetraploidy due to a Germline Mutation in SEPT6. American Journal of Hematology, 2021, , .	2.0	1
2	Homozygous deletions implicate non-coding epigenetic marks in Autism spectrum disorder. Scientific Reports, 2020, 10, 14045.	1.6	12
3	Normalizing hepcidin predicts TMPRSS6 mutation status in patients with chronic iron deficiency. Blood, 2018, 132, 448-452.	0.6	16
4	Congenital macrothrombocytopenia with focal myelofibrosis due to mutations in human G6b-B is rescued in humanized mice. Blood, 2018, 132, 1399-1412.	0.6	37
5	Congenital X-Linked Myelodysplasia with Tetraploidy Is Associated with De Novo Germline C-Terminal Mutation of SEPT6, a Septin Filament Protein. Blood, 2018, 132, 644-644.	0.6	О
6	Biallelic mutations in human DCC cause developmental split-brain syndrome. Nature Genetics, 2017, 49, 606-612.	9.4	62
7	Ringed sideroblasts in βâ€thalassemia. Pediatric Blood and Cancer, 2017, 64, e26324.	0.8	4
8	Evolution of <scp>GOUNDRY</scp> , a cryptic subgroup of <i>AnophelesÂgambiaeÂs.l</i> , and its impact on susceptibility to <i>Plasmodium</i> infection. Molecular Ecology, 2016, 25, 1494-1510.	2.0	18
9	Hyperammonemia as a Presenting Feature in Two Siblings with FBXL4 Variants. JIMD Reports, 2016, 35, 7-15.	0.7	9
10	Mutations in the substrate binding glycine-rich loop of the mitochondrial processing peptidase- $\hat{l}\pm$ protein (PMPCA) cause a severe mitochondrial disease. Journal of Physical Education and Sports Management, 2016, 2, a000786.	0.5	33
11	Genetic Structure of a Local Population of the Anopheles gambiae Complex in Burkina Faso. PLoS ONE, 2016, 11, e0145308.	1.1	8
12	The kdr-bearing haplotype and susceptibility to Plasmodium falciparum in Anopheles gambiae: genetic correlation and functional testing. Malaria Journal, 2015, 14, 391.	0.8	35
13	Congenital sideroblastic anemia due to mutations in the mitochondrial HSP70 homologue HSPA9. Blood, 2015, 126, 2734-2738.	0.6	78
14	Association mapping by pooled sequencing identifies TOLL 11 as a protective factor against Plasmodium falciparum in Anopheles gambiae. BMC Genomics, 2015, 16, 779.	1.2	19
15	Mutations in PYCR2, Encoding Pyrroline-5-Carboxylate Reductase 2, Cause Microcephaly and Hypomyelination. American Journal of Human Genetics, 2015, 96, 709-719.	2.6	60
16	LARS2 Variants Associated with Hydrops, Lactic Acidosis, Sideroblastic Anemia, and Multisystem Failure. JIMD Reports, 2015, 28, 49-57.	0.7	48
17	Expanding the Phenotype Associated With the <i>NEFL</i> Mutation. JAMA Neurology, 2014, 71, 1413.	4.5	30
18	Mutations in TRNT1 cause congenital sideroblastic anemia with immunodeficiency, fevers, and developmental delay (SIFD). Blood, 2014, 124, 2867-2871.	0.6	162

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19	GATA2 Mutations In Pediatric Myelodysplastic Syndromes and Bone Marrow Failure. Blood, 2013, 122, 1520-1520.	0.6	3
20	Responseâ€"New Mosquito Subgroup Breeds Questions. Science, 2011, 332, 420-421.	6.0	0
21	A Cryptic Subgroup of <i>Anopheles gambiae</i> Is Highly Susceptible to Human Malaria Parasites. Science, 2011, 331, 596-598.	6.0	129
22	Lack of Association of the Serotonin Transporter Polymorphism with the Sudden Infant Death Syndrome in the San Diego Dataset. Pediatric Research, 2010, 68, 1.	1.1	33
23	Serotonin-Related FEV Gene Variant in the Sudden Infant Death Syndrome Is a Common Polymorphism in the African-American Population. Pediatric Research, 2009, 66, 631-635.	1.1	16
24	Mutations in TMPRSS6 cause iron-refractory iron deficiency anemia (IRIDA). Nature Genetics, 2008, 40, 569-571.	9.4	586
25	A major genetic locus controlling natural Plasmodium falciparum infection is shared by East and West African Anopheles gambiae. Malaria Journal, 2007, 6, 87.	0.8	23
26	Natural Malaria Infection in Anopheles gambiae Is Regulated by a Single Genomic Control Region. Science, 2006, 312, 577-579.	6.0	261
27	FINDING PROSTATE CANCER SUSCEPTIBILITY GENES. Annual Review of Genomics and Human Genetics, 2004, 5, 151-175.	2.5	36
28	No Bias in Linkage Analysis. American Journal of Human Genetics, 2004, 75, 722-723.	2.6	4
29	Sequence-Based Linkage Analysis. American Journal of Human Genetics, 2004, 75, 647-653.	2.6	4
30	A 3.9-Centimorgan-Resolution Human Single-Nucleotide Polymorphism Linkage Map and Screening Set. American Journal of Human Genetics, 2003, 73, 271-284.	2.6	112
31	Genetic Loci Affecting Resistance to Human Malaria Parasites in a West African Mosquito Vector Population. Science, 2002, 298, 213-216.	6.0	121
32	Efficient Multipoint Linkage Analysis through Reduction of Inheritance Space. American Journal of Human Genetics, 2001, 68, 963-977.	2.6	130
33	A Joint Analysis of Asthma Affection Status and IgE Levels in Multiple Data Sets Collected for Asthma. Genetic Epidemiology, 2001, 21, S148-53.	0.6	2