Pawel Gawlinski

List of Publications by Year
in descending order

15 papers	290 citations	1040056 9 h-index	1058476 14 g-index
15 all docs	15 docs citations	15 times ranked	806 citing authors

#	Article	IF	Citations
1	Loss of SMPD4 Causes a Developmental Disorder Characterized by Microcephaly and Congenital Arthrogryposis. American Journal of Human Genetics, 2019, 105, 689-705.	6.2	48
2	Why Functional Pre-Erythrocytic and Bloodstage Malaria Vaccines Fail: A Meta-Analysis of Fully Protective Immunizations and Novel Immunological Model. PLoS ONE, 2010, 5, e10685.	2.5	37
3	Malaria's deadly secret: a skin stage. Trends in Parasitology, 2012, 28, 142-150.	3.3	37
4	Comprehensive genomic analysis of patients with disorders of cerebral cortical development. European Journal of Human Genetics, 2018, 26, 1121-1131.	2.8	35
5	PEHO Syndrome May Represent Phenotypic Expansion at the Severe End of the Early-Onset Encephalopathies. Pediatric Neurology, 2016, 60, 83-87.	2.1	25
6	Bi-allelic Pathogenic Variants in TUBGCP2 Cause Microcephaly and Lissencephaly Spectrum Disorders. American Journal of Human Genetics, 2019, 105, 1005-1015.	6.2	24
7	The Drosophila mitotic inhibitor FrÃ¼hstart specifically binds to the hydrophobic patch of cyclins. EMBO Reports, 2007, 8, 490-496.	4.5	23
8	Posterior Neocortex-Specific Regulation of Neuronal Migration by CEP85L Identifies Maternal Centriole-Dependent Activation of CDK5. Neuron, 2020, 106, 246-255.e6.	8.1	19
9	Phenotype expansion and development in Kosaki overgrowth syndrome. Clinical Genetics, 2018, 93, 919-924.	2.0	17
10	Exome Sequencing Reveals Novel Variants and Expands the Genetic Landscape for Congenital Microcephaly. Genes, 2021, 12, 2014.	2.4	8
11	A Patient with Berardinelli-Seip Syndrome, Novel <i>AGPAT2</i> Splicesite Mutation and Concomitant Development of Non-diabetic Polyneuropathy. JCRPE Journal of Clinical Research in Pediatric Endocrinology, 2019, 11, 319-326.	0.9	6
12	Further Delineation of Phenotype and Genotype of Primary Microcephaly Syndrome with Cortical Malformations Associated with Mutations in the WDR62 Gene. Genes, 2021, 12, 594.	2.4	5
13	Clinical and molecular characterization of craniofrontonasal syndrome: new symptoms and novel pathogenic variants in the EFNB1 gene. Orphanet Journal of Rare Diseases, 2021, 16, 286.	2.7	3
14	De Novo ACTG1 Variant Expands the Phenotype and Genotype of Partial Deafness and Baraitserâ€“Winter Syndrome. International Journal of Molecular Sciences, 2022, 23, 692.	4.1	3
15	Kosaki overgrowth syndrome due to a novel de novo <scp>PDGFRB</scp> variant. Clinical Genetics, 2022, 101, 144-145.	2.0	0