Keith Vaux

List of Publications by Year
in descending order

21 papers	2,583 citations	430874 18 h-index	752698 20 g-index
22 all docs	22 docs citations	22 times ranked	5341 citing authors

#	Article	IF	Citations
1	Neural responses to affective speech, including motherese, map onto clinical and social eye tracking profiles in toddlers with ASD. Nature Human Behaviour, 2022, 6, 443-454.	12.0	14
2	Birth outcomes in women who have taken adalimumab in pregnancy: A prospective cohort study. PLoS ONE, 2019, 14, e0223603.	2.5	40
3	Paternally inherited cis-regulatory structural variants are associated with autism. Science, 2018, 360, 327-331.	12.6	174
4	Prevalence of Fetal Alcohol Spectrum Disorders in 4 US Communities. JAMA - Journal of the American Medical Association, 2018, 319, 474.	7.4	562
5	Inside â€œOutsideâ€•Job: Unexpected Geometric Skin Ulcerations Overlying Orthopedic Hardware After Multimodal Laser Scar Revision. Dermatologic Surgery, 2018, 44, 1231-1233.	0.8	0
6	Biallelic Mutations in ADPRHL2, Encoding ADP-Ribosylhydrolase 3, Lead to a Degenerative Pediatric Stress-Induced Epileptic Ataxia Syndrome. American Journal of Human Genetics, 2018, 103, 431-439.	6.2	62
7	FOXP1-related intellectual disability syndrome: a recognisable entity. Journal of Medical Genetics, 2017, 54, 613-623.	3.2	48
8	Frequency and Complexity of De Novo Structural Mutation in Autism. American Journal of Human Genetics, 2016, 98, 667-679.	6.2	88
9	Functional genome-wide siRNA screen identifies KIAA0586 as mutated in Joubert syndrome. ELife, 2015, 4, e06602.	6.0	64
10	Inactivating mutations in MFSD2A, required for omega-3 fatty acid transport in brain, cause a lethal microcephaly syndrome. Nature Genetics, 2015, 47, 809-813.	21.4	180
11	Biallelic mutations in SNX14 cause a syndromic form of cerebellar atrophy and lysosome-autophagosome dysfunction. Nature Genetics, 2015, 47, 528-534.	21.4	111
12	The genetic landscape of autism spectrum disorders. Developmental Medicine and Child Neurology, 2014, 56, 12-18.	2.1	83
13	Exome Sequencing Links Corticospinal Motor Neuron Disease to Common Neurodegenerative Disorders. Science, 2014, 343, 506-511.	12.6	466
14	AMPD2 Regulates GTP Synthesis and Is Mutated in a Potentially Treatable Neurodegenerative Brainstem Disorder. Cell, 2013, 154, 505-517.	28.9	94
15	Placenta Accreta in a Separate Uterine Horn. Pediatric and Developmental Pathology, 2010, 13, 63-65.	1.0	4
16	Parkes Weber syndrome, vein of Galen aneurysmal malformation, and other fast-flow vascular anomalies are caused byRASA1 mutations. Human Mutation, 2008, 29, 959-965.	2.5	382
17	The role of molecular testing and enzyme analysis in the management of hypomorphic citrullinemia. American Journal of Medical Genetics, Part A, 2008, 146A, 2885-2890.	1.2	24
18	Cyclophosphamide, methotrexate, and cytarabine embropathy: Is apoptosis the common pathway?. Birth Defects Research Part A: Clinical and Molecular Teratology, 2003, 67, 403-408.	1.6	44

#	Article	IF	Citations
19	The Safe and Effective Use of Propofol Sedation in Children Undergoing Diagnostic and Therapeutic Procedures: Experience in a Pediatric ICU and a Review of the Literature. Pediatric Emergency Care, 2003, 19, 385-392.	0.9	61
20	Use of gabapentin in the treatment of childhood reflex sympathetic dystrophy. Pediatric Neurology, 2000, 22, 220-221.	2.1	62
21	Preventive Control of AIDS by the Dental Profession: A Survey of Practices in a Large Urban Area. Journal of Public Health Dentistry, 1990, 50, 7-12.	1.2	20