Wesam Kurdi

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/1417423/publications.pdf

Version: 2024-02-01

25	1,614	516710	552781
papers	citations	16 h-index	26 g-index

27	27	27	3959
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Accelerating Novel Candidate Gene Discovery in Neurogenetic Disorders via Whole-Exome Sequencing of Prescreened Multiplex Consanguineous Families. Cell Reports, 2015, 10, 148-161.	6.4	375
2	The landscape of genetic diseases in Saudi Arabia based on the first 1000 diagnostic panels and exomes. Human Genetics, 2017, 136, 921-939.	3.8	209
3	Lessons Learned from Large-Scale, First-Tier Clinical Exome Sequencing in a Highly Consanguineous Population. American Journal of Human Genetics, 2019, 104, 1182-1201.	6.2	184
4	Expanding the genetic heterogeneity of intellectual disability. Human Genetics, 2017, 136, 1419-1429.	3.8	122
5	The genetic landscape of familial congenital hydrocephalus. Annals of Neurology, 2017, 81, 890-897.	5.3	108
6	Identification of embryonic lethal genes in humans by autozygosity mapping and exome sequencing in consanguineous families. Genome Biology, 2015, 16, 116.	8.8	91
7	Molecular autopsy in maternalâ€“fetal medicine. Genetics in Medicine, 2018, 20, 420-427.	2.4	84
8	Autozygome and high throughput confirmation of disease genes candidacy. Genetics in Medicine, 2019, 21, 736-742.	2.4	81
9	Autozygosity reveals recessive mutations and novel mechanisms in dominant genes: implications in variant interpretation. Genetics in Medicine, 2017, 19, 1144-1150.	2.4	77
10	KIAA1109 Variants Are Associated with a Severe Disorder of Brain Development and Arthrogryposis. American Journal of Human Genetics, 2018, 102, 116-132.	6.2	46
11	Identification of a novel MKS locus defined byTMEM107mutation. Human Molecular Genetics, 2015, 24, 5211-5218.	2.9	42
12	Absence of GP130 cytokine receptor signaling causes extended StÃ¼ve-Wiedemann syndrome. Journal of Experimental Medicine, 2020, 217, .	8.5	41
13	Genetic spectrum of Saudi Arabian patients with antenatal cystic kidney disease and ciliopathy phenotypes using a targeted renal gene panel. Journal of Medical Genetics, 2016, 53, 338-347.	3.2	28
14	Delayed normalisation of uterine artery doppler waveforms is not a benign phenomenon. European Journal of Obstetrics, Gynecology and Reproductive Biology, 2004, 117, 20-23.	1.1	23
15	Bialleleic PKD1 mutations underlie early-onset autosomal dominant polycystic kidney disease in Saudi Arabian families. Pediatric Nephrology, 2019, 34, 1615-1623.	1.7	21
16	Lethal variants in humans: lessons learned from a large molecular autopsy cohort. Genome Medicine, 2021, 13, 161.	8.2	13
17	Novel loss of function variants in FRAS1 AND FREM2 underlie renal agenesis in consanguineous families. Journal of Nephrology, 2021, 34, 893-900.	2.0	11
18	Bleeding and thrombotic risk in pregnant women with Fontan physiology. Heart, 2021, 107, 1390-1397.	2.9	9

#	Article	IF	Citations
19	Prenatal exome sequencing and chromosomal microarray analysis in fetal structural anomalies in a highly consanguineous population reveals a propensity of ciliopathy genes causing multisystem phenotypes. Human Genetics, 2022, 141, 101-126.	3.8	6
20	Fetal Anomalies Associated with Novel Pathogenic Variants in TMEM94. Genes, 2020, 11, 967.	2.4	4
21	The effect of hyoscine-N-butylbromide on pain perception during and after hysterosalpingography in infertile women: a systematic review and meta-analysis of randomised controlled trials. Human Fertility, 2020, , 1-8.	1.7	2
22	Robot-assisted surgical staging for ovarian cancer in pregnant women. Journal of Robotic Surgery, 2012, 6, 163-166.	1.8	1
23	Efficacy of hyoscine in pain management during hysteroscopy: a systematic review and meta-analysis. Journal of the Turkish German Gynecology Association, 2022, 23, 51-57.	0.6	1
24	Efficacy of hyoscine in pain management during hysteroscopy: a systematic review and meta-analysis. Journal of the Turkish German Gynecology Association, 2021, .	0.6	1
25	The Evolution of Fetal Procedures. Innovations in Surgery and Interventional Medicine, 2022, 2, 15-16.	0.1	0