Charlotte Brasch-Andersen

List of Publications by Year
in descending order

28 papers	593 citations	949033 11 h-index	721071 23 g-index
29 all docs	29 docs citations	29 times ranked	1423 citing authors

#	Article	IF	Citations
1	Monoâ€allelic loss of <scp>YTHDF3</scp> and neurodevelopmental disorder: clinical features of four individuals with 8q12.3 deletions. Clinical Genetics, 2022, 101, 208-213.	1.0	2
2	Variants in ADD1 cause intellectual disability, corpus callosum dysgenesis, and ventriculomegaly in humans. Genetics in Medicine, 2022, 24, 319-331.	1.1	6
3	Genotype-Phenotype Comparison in POGZ-Related Neurodevelopmental Disorders by Using Clinical Scoring. Genes, 2022, 13, 154.	1.0	6
4	Variants in PHF8 cause a spectrum of X-linked neurodevelopmental disorders and facial dysmorphology. Human Genetics and Genomics Advances, 2022, 3, 100102.	1.0	5
5	Loss-of-function variants in SRRM2 cause a neurodevelopmental disorder. Genetics in Medicine, 2022, 24, 1774-1780.	1.1	16
6	National data on the early clinical use of nonâ€invasive prenatal testing in public and private healthcare in Denmark 2013â€“2017. Acta Obstetricia Et Gynecologica Scandinavica, 2021, 100, 884-892.	1.3	11
7	Total number of reads affects the accuracy of fetal fraction estimates in NIPT. Molecular Genetics & Genomic Medicine, 2021, 9, e1653.	0.6	8
8	Rare deleterious mutations of HNRNP genes result in shared neurodevelopmental disorders. Genome Medicine, 2021, 13, 63.	3.6	50
9	Impaired glucose-1,6-biphosphate production due to bi-allelic PGM2L1 mutations is associated with a neurodevelopmental disorder. American Journal of Human Genetics, 2021, 108, 1151-1160.	2.6	9
10	<scp>ZMYND11</scp> variants are a novel cause of centrotemporal and generalised epilepsies with neurodevelopmental disorder. Clinical Genetics, 2021, 100, 412-429.	1.0	5
11	Phenotypic heterogeneity and mosaicism in Xia-Gibbs syndrome: Five Danish patients with novel variants in AHDC1. European Journal of Medical Genetics, 2021, 64, 104280.	0.7	3
12	Association of serum surfactant protein D and SFTPD gene variants with asthma in Danish children, adolescents, and young adults. Immunity, Inflammation and Disease, 2021, , .	1.3	2
13	NCKAP1 Disruptive Variants Lead to a Neurodevelopmental Disorder with Core Features of Autism. American Journal of Human Genetics, 2020, 107, 963-976.	2.6	18
14	A new 1p36.13â€1p36.12 microdeletion syndrome characterized by learning disability, behavioral abnormalities, and ptosis. Clinical Genetics, 2020, 97, 927-932.	1.0	6
15	Is MED13L-related intellectual disability a recognizable syndrome?. European Journal of Medical Genetics, 2019, 62, 129-136.	0.7	21
16	Enriched power of disease-concordant twin-case-only design in detecting interactions in genome-wide association studies. European Journal of Human Genetics, 2019, 27, 631-636.	1.4	4
17	Estimating the effect size of the 15Q11.2 BP1â€“BP2 deletion and its contribution to neurodevelopmental symptoms: recommendations for practice. Journal of Medical Genetics, 2019, 56, 701-710.	1.5	43
18	CD18 is redundant for the response to multiple vaccines: A case study. Pediatric Allergy and Immunology, 2019, 30, 136-139.	1.1	0

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19	Prostaglandin E₂â€<scp>EP</scp>₃ receptor subtype gene deletion in mother and son impairs platelet aggregation. British Journal of Haematology, 2019, 184, 851-853.	1.2	0
20	Deletion of T-type calcium channels Cav3.1 or Cav3.2 attenuates endothelial dysfunction in aging mice. Pflugers Archiv European Journal of Physiology, 2018, 470, 355-365.	1.3	12
21	A case-only genome-wide association study on gene-sex interaction in allergic rhinitis. Annals of Allergy, Asthma and Immunology, 2018, 121, 366-367.e2.	0.5	3
22	Genomic Analyses of Breast Cancer Progression Reveal Distinct Routes of Metastasis Emergence. Scientific Reports, 2017, 7, 43813.	1.6	24
23	YY1 Haploinsufficiency Causes an Intellectual Disability Syndrome Featuring Transcriptional and Chromatin Dysfunction. American Journal of Human Genetics, 2017, 100, 907-925.	2.6	125
24	The <scp>ABCB</scp>1, rs9282564, <scp>AG</scp> and <scp>TT</scp> Genotypes and the <scp>COMT</scp>, rs4680, <scp>AA</scp> Genotype are Less Frequent in Deceased Patients with Opioid Addiction than in Living Patients with Opioid Addiction. Basic and Clinical Pharmacology and Toxicology, 2016, 119, 381-388.	1.2	14
25	17q12 deletion and duplication syndrome in Denmarkâ€”A clinical cohort of 38 patients and review of the literature. American Journal of Medical Genetics, Part A, 2016, 170, 2934-2942.	0.7	53
26	<scp>S</scp>100A14 is a novel independent prognostic biomarker in the tripleâ€negative breast cancer subtype. International Journal of Cancer, 2015, 137, 2093-2103.	2.3	19
27	A candidate gene study of serotonergic pathway genes and pain relief during treatment with escitalopram in patients with neuropathic pain shows significant association to serotonin receptor2C (HTR2C). European Journal of Clinical Pharmacology, 2011, 67, 1131-1137.	0.8	34
28	Possible gene dosage effect of glutathione-S-transferases on atopic asthma: Using real-time PCR for quantification of GSTM1 and GSTT1 gene copy numbers. Human Mutation, 2004, 24, 208-214.	1.1	94