Ann C M Smith

List of Publications by Year
in descending order

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Version: 2024-02-01

29 papers	1,757 citations	394421 19 h-index	526287 27 g-index
33 all docs	33 docs citations	33 times ranked	1203 citing authors

#	Article	IF	Citations
1	Variegation of autism related traits across seven neurogenetic disorders. Translational Psychiatry, 2022, 12, 149.	4.8	5
2	De Novo KAT5 Variants Cause a Syndrome with Recognizable Facial Dysmorphisms, Cerebellar Atrophy, Sleep Disturbance, and Epilepsy. American Journal of Human Genetics, 2020, 107, 564-574.	6.2	14
3	Twentyâ€fourâ€hour motor activity and body temperature patterns suggest altered central circadian timekeeping in Smithâ€“Magenis syndrome, a neurodevelopmental disorder. American Journal of Medical Genetics, Part A, 2019, 179, 224-236.	1.2	20
4	Auditory Phenotype of Smithâ€“Magenis Syndrome. Journal of Speech, Language, and Hearing Research, 2017, 60, 1076-1087.	1.6	13
5	Delayed diagnosis in a house of correction: Smithâ€“Magenis syndrome due to a de novo nonsense RAI1 variant. American Journal of Medical Genetics, Part A, 2016, 170, 2383-2388.	1.2	4
6	Analysis of the Sensory Profile in Children with Smithâ€“Magenis Syndrome. Physical and Occupational Therapy in Pediatrics, 2012, 32, 48-65.	1.3	16
7	Molecular Analysis of the Retinoic Acid Induced 1 Gene (RAI1) in Patients with Suspected Smith-Magenis Syndrome without the 17p11.2 Deletion. PLoS ONE, 2011, 6, e22861.	2.5	38
8	Diagnostic utility of daytime salivary melatonin levels in Smithâ€“Magenis syndrome. American Journal of Medical Genetics, Part A, 2010, 152A, 96-101.	1.2	12
9	Autism spectrum features in Smithâ€“Magenis syndrome. American Journal of Medical Genetics, Part C: Seminars in Medical Genetics, 2010, 154C, 456-462.	1.6	92
10	Pharmacological treatment of disruptive behavior in Smithâ€“Magenis syndrome. American Journal of Medical Genetics, Part C: Seminars in Medical Genetics, 2010, 154C, 463-468.	1.6	24
11	Review of disrupted sleep patterns in Smithâ€“Magenis syndrome and normal melatonin secretion in a patient with an atypical interstitial 17p11.2 deletion. American Journal of Medical Genetics, Part A, 2009, 149A, 1382-1391.	1.2	43
12	Neurodevelopment of Children Under 3 Years of Age With Smith-Magenis Syndrome. Pediatric Neurology, 2009, 41, 250-258.	2.1	39
13	New developments in Smith-Magenis syndrome (del 17p11.2). Current Opinion in Neurology, 2007, 20, 125-134.	3.6	58
14	Neurologic and Developmental Features of the Smith-Magenis Syndrome (del 17p11.2). Pediatric Neurology, 2006, 34, 337-350.	2.1	130
15	Adaptive and Maladaptive Behavior in Children with Smith-Magenis Syndrome. Journal of Autism and Developmental Disorders, 2006, 36, 541-552.	2.7	55
16	Craniofacial and dental phenotype of Smithâ€“Magenis syndrome. American Journal of Medical Genetics, Part A, 2006, 140A, 2556-2561.	1.2	41
17	Diagnostic FISH probes for del(17)(p11.2p11.2) associated with Smith-Magenis syndrome should contain theRAI1gene. , 2005, 132A, 278-282.		33
18	Overview of Smith-Magenis syndrome. Journal of the Association of Genetic Technologists, 2005, 31, 163-7.	0.1	21

#	Article	IF	Citations
19	Hypercholesterolemia in children with Smith-Magenis syndrome: del (17)(p11.2p11.2). Genetics in Medicine, 2002, 4, 118-125.	2.4	59
20	To Use or Not to Use: The Prenatal Genetic Technology/Worry Conundrum. Journal of Genetic Counseling, 2000, 9, 203-217.	1.6	14
21	To Use or Not to Use: Male Partners' Perspectives on Decision Making About Prenatal Diagnosis. Journal of Genetic Counseling, 2000, 9, 33-45.	1.6	26
22	Risk of abnormal pregnancy outcome in carriers of balanced reciprocal translocations involving the Miller-Dieker syndrome (MDS) critical region in chromosome 17p13.3. , 1999, 85, 369-375.		13
23	Behavioral phenotype of smith-magenis syndrome (del 17p11.2). American Journal of Medical Genetics Part A, 1998, 81, 179-185.	2.4	139
24	Sleep disturbance in smith-magenis syndrome (del 17 p11.2). American Journal of Medical Genetics Part A, 1998, 81, 186-191.	2.4	132
25	A Revision of the Lissencephaly and Miller-Dieker Syndrome Critical Regions in Chromosome 17p13.3. Human Molecular Genetics, 1997, 6, 147-155.	2.9	176
26	PRENATAL DIAGNOSIS OF UNIPARENTAL DISOMY 15 FOLLOWING TRISOMY 15 MOSAICISM. Prenatal Diagnosis, 1996, 16, 323-332.	2.3	73
27	Genetic counseling for the next 25 years: Models for the future. Journal of Genetic Counseling, 1995, 4, 115-124.	1.6	26
28	Interstitial deletion of (17)(p11.2p11.2) in nine patients. American Journal of Medical Genetics Part A, 1986, 24, 393-414.	2.4	364
29	Consideration of connective tissue dysfunction in the fragile X syndrome. American Journal of Medical Genetics Part A, 1984, 17, 111-121.	2.4	59