

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

393982

19
h-index

552369

26
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. <i>Translational Psychiatry</i> , 2022, 12, 149.	2.4	5
2	De Novo KAT5 Variants Cause a Syndrome with Recognizable Facial Dysmorphisms, Cerebellar Atrophy, Sleep Disturbance, and Epilepsy. <i>American Journal of Human Genetics</i> , 2020, 107, 564-574.	2.6	14
3	Twenty-four-hour motor activity and body temperature patterns suggest altered central circadian timekeeping in Smith-Magenis syndrome, a neurodevelopmental disorder. <i>American Journal of Medical Genetics, Part A</i> , 2019, 179, 224-236.	0.7	20
4	Auditory Phenotype of Smith-Magenis Syndrome. <i>Journal of Speech, Language, and Hearing Research</i> , 2017, 60, 1076-1087.	0.7	13
5	Delayed diagnosis in a house of correction: Smith-Magenis syndrome due to a de novo nonsense <i>RAI1</i> variant. <i>American Journal of Medical Genetics, Part A</i> , 2016, 170, 2383-2388.	0.7	4
6	Analysis of the Sensory Profile in Children with Smith-Magenis Syndrome. <i>Physical and Occupational Therapy in Pediatrics</i> , 2012, 32, 48-65.	0.8	16
7	Molecular Analysis of the Retinoic Acid Induced 1 Gene (<i>RAI1</i>) in Patients with Suspected Smith-Magenis Syndrome without the 17p11.2 Deletion. <i>PLoS ONE</i> , 2011, 6, e22861.	1.1	38
8	Diagnostic utility of daytime salivary melatonin levels in Smith-Magenis syndrome. <i>American Journal of Medical Genetics, Part A</i> , 2010, 152A, 96-101.	0.7	12
9	Autism spectrum features in Smith-Magenis syndrome. <i>American Journal of Medical Genetics, Part C: Seminars in Medical Genetics</i> , 2010, 154C, 456-462.	0.7	92
10	Pharmacological treatment of disruptive behavior in Smith-Magenis syndrome. <i>American Journal of Medical Genetics, Part C: Seminars in Medical Genetics</i> , 2010, 154C, 463-468.	0.7	24
11	Review of disrupted sleep patterns in Smith-Magenis syndrome and normal melatonin secretion in a patient with an atypical interstitial 17p11.2 deletion. <i>American Journal of Medical Genetics, Part A</i> , 2009, 149A, 1382-1391.	0.7	43
12	Neurodevelopment of Children Under 3 Years of Age With Smith-Magenis Syndrome. <i>Pediatric Neurology</i> , 2009, 41, 250-258.	1.0	39
13	New developments in Smith-Magenis syndrome (del 17p11.2). <i>Current Opinion in Neurology</i> , 2007, 20, 125-134.	1.8	58
14	Neurologic and Developmental Features of the Smith-Magenis Syndrome (del 17p11.2). <i>Pediatric Neurology</i> , 2006, 34, 337-350.	1.0	130
15	Adaptive and Maladaptive Behavior in Children with Smith-Magenis Syndrome. <i>Journal of Autism and Developmental Disorders</i> , 2006, 36, 541-552.	1.7	55
16	Craniofacial and dental phenotype of Smith-Magenis syndrome. <i>American Journal of Medical Genetics, Part A</i> , 2006, 140A, 2556-2561.	0.7	41
17	Diagnostic FISH probes for del(17)(p11.2p11.2) associated with Smith-Magenis syndrome should contain the <i>RAI1</i> gene. , 2005, 132A, 278-282.		33
18	Overview of Smith-Magenis syndrome. <i>Journal of the Association of Genetic Technologists</i> , 2005, 31, 163-7.	0.1	21

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19	Hypercholesterolemia in children with Smith-Magenis syndrome: del (17)(p11.2p11.2). Genetics in Medicine, 2002, 4, 118-125.	1.1	59
20	To Use or Not to Use: The Prenatal Genetic Technology/Worry Conundrum. Journal of Genetic Counseling, 2000, 9, 203-217.	0.9	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.	0.9	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). , 1998, 81, 179-185.		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.	1.4	176
26	PRENATAL DIAGNOSIS OF UNIPARENTAL DISOMY 15 FOLLOWING TRISOMY 15 MOSAICISM. Prenatal Diagnosis, 1996, 16, 323-332.	1.1	73
27	Genetic counseling for the next 25 years: Models for the future. Journal of Genetic Counseling, 1995, 4, 115-124.	0.9	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