Shweta U Dhar

List of Publications by Year
in descending order

25 papers	589 citations	759055 12 h-index	677027 22 g-index
27 all docs	27 docs citations	27 times ranked	1684 citing authors

#	Article	IF	Citations
1	Molecular diagnostic experience of whole-exome sequencing in adult patients. Genetics in Medicine, 2016, 18, 678-685.	1.1	186
2	A recurrent p.Arg92Trp variant in steroidogenic factor-1 (NR5A1) can act as a molecular switch in human sex development. Human Molecular Genetics, 2016, 25, 3446-3453.	1.4	90
3	Enhancing exposure to genetics and genomics through an innovative medical school curriculum. Genetics in Medicine, 2012, 14, 163-167.	1.1	48
4	Disruptive mutations in TANC2 define a neurodevelopmental syndrome associated with psychiatric disorders. Nature Communications, 2019, 10, 4679.	5.8	43
5	Outcomes of Integrating Genetics in Management of Patients With Retinoblastoma. JAMA Ophthalmology, 2011, 129, 1428.	2.6	29
6	De novo missense variant in the GTPase effector domain (GED) of DNM1L leads to static encephalopathy and seizures. Journal of Physical Education and Sports Management, 2019, 5, a003673.	0.5	24
7	Tetrasomy 13q mosaicism associated with phylloid hypomelanosis and precocious puberty. American Journal of Medical Genetics, Part A, 2009, 149A, 993-996.	0.7	22
8	Autism genetic testing information needs among parents of affected children: A qualitative study. Patient Education and Counseling, 2016, 99, 1011-1016.	1.0	21
9	Genetic Testing Experiences Among Parents of Children with Autism Spectrum Disorder in the United States. Journal of Autism and Developmental Disorders, 2019, 49, 4821-4833.	1.7	18
10	Financial barriers in a county genetics clinic: Problems and solutions. Journal of Genetic Counseling, 2020, 29, 678-688.	0.9	17
11	Development and evaluation of a genomics training program for community health workers in Texas. Genetics in Medicine, 2018, 20, 1030-1037.	1.1	15
12	Needs assessment in genetic testing education: A survey of parents of children with autism spectrum disorder in the united states. Autism Research, 2019, 12, 1162-1170.	2.1	15
13	The practice of adult genetics: A 7â€year experience from a single center. American Journal of Medical Genetics, Part A, 2013, 161, 89-93.	0.7	10
14	Pursuing genetic testing for children with autism spectrum disorders: What do parents think?. Journal of Genetic Counseling, 2021, 30, 370-382.	0.9	9
15	Cranioâ€metaâ€diaphyseal dysplasia: 25 year followâ€up and review of literature. American Journal of Medical Genetics, Part A, 2010, 152A, 2335-2338.	0.7	7
16	Atypical Alexander disease with dystonia, retinopathy, and a brain mass mimicking astrocytoma. Neurology: Genetics, 2018, 4, e248.	0.9	7
17	Family Health Historyâ€“Based Cancer Prevention Training for Community Health Workers. American Journal of Preventive Medicine, 2021, 60, e159-e167.	1.6	4
18	Quality improvement of clinic flow for complex genetic conditions: Using Ehlersâ€“Danlos syndrome as a model. Molecular Genetics & Genomic Medicine, 2018, 6, 993-1000.	0.6	3

#	Article	IF	Citations
19	An innovative medical school curriculum to enhance exposure to genetics and genomics: Updates and outcomes. Genetics in Medicine, 2022, 24, 722-728.	1.1	3
20	Psychometric Properties of the POAGTS: A Tool for Understanding Parentsâ€™ Perceptions Regarding Autism Spectrum Disorder Genetic Testing. International Journal of Environmental Research and Public Health, 2021, 18, 3323.	1.2	2
21	Implementation of a Medical School Elective Course Incorporating Case-Based Learning: a Pilot Study. Medical Science Educator, 2020, 30, 339-344.	0.7	1
22	Texas health educatorsâ€™Âpractice in basic genomics education and services. Personalized Medicine, 2021, 18, 55-66.	0.8	1
23	Genetic Evaluation for Common, Chronic Disorders of Adulthood. , 2019, , 265-282.		0
24	Cancer syndromes. , 2020, , 85-108.		0
25	Introduction to the practice of adult genetics. , 2020, , 3-7.		0