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	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
2	Texas health educatorsâ€™Âpractice in basic genomics education and services. Personalized Medicine, 2021, 18, 55-66.	0.8	1
3	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
4	Family Health Historyâ€“Based Cancer Prevention Training for Community Health Workers. American Journal of Preventive Medicine, 2021, 60, e159-e167.	1.6	4
5	Pursuing genetic testing for children with autism spectrum disorders: What do parents think?. Journal of Genetic Counseling, 2021, 30, 370-382.	0.9	9
6	Implementation of a Medical School Elective Course Incorporating Case-Based Learning: a Pilot Study. Medical Science Educator, 2020, 30, 339-344.	0.7	1
7	Financial barriers in a county genetics clinic: Problems and solutions. Journal of Genetic Counseling, 2020, 29, 678-688.	0.9	17
8	Cancer syndromes. , 2020, , 85-108.		0
9	Introduction to the practice of adult genetics. , 2020, , 3-7.		0
10	Disruptive mutations in TANC2 define a neurodevelopmental syndrome associated with psychiatric disorders. Nature Communications, 2019, 10, 4679.	5.8	43
11	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
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	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
14	Genetic Evaluation for Common, Chronic Disorders of Adulthood. , 2019, , 265-282.		0
15	Development and evaluation of a genomics training program for community health workers in Texas. Genetics in Medicine, 2018, 20, 1030-1037.	1.1	15
16	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
17	Atypical Alexander disease with dystonia, retinopathy, and a brain mass mimicking astrocytoma. Neurology: Genetics, 2018, 4, e248.	0.9	7
18	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

#	Article	IF	Citations
19	Autism genetic testing information needs among parents of affected children: A qualitative study. Patient Education and Counseling, 2016, 99, 1011-1016.	1.0	21
20	Molecular diagnostic experience of whole-exome sequencing in adult patients. Genetics in Medicine, 2016, 18, 678-685.	1.1	186
21	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
22	Enhancing exposure to genetics and genomics through an innovative medical school curriculum. Genetics in Medicine, 2012, 14, 163-167.	1.1	48
23	Outcomes of Integrating Genetics in Management of Patients With Retinoblastoma. JAMA Ophthalmology, 2011, 129, 1428.	2.6	29
24	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
25	Tetrasomy 13q mosaicism associated with phylloid hypomelanosis and precocious puberty. American Journal of Medical Genetics, Part A, 2009, 149A, 993-996.	0.7	22