Dawn Laney

List of Publications by Year
in descending order

Source: https://exaly.com/author-pdf/1602445/publications.pdf

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19	845	933447 10	888059 17
papers	citations	h-index	g-index

36	36	36	983
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Fabry disease revisited: Management and treatment recommendations for adult patients. Molecular Genetics and Metabolism, 2018, 123, 416-427.	1.1	391
2	The management and treatment of children with Fabry disease: A United States-based perspective. Molecular Genetics and Metabolism, 2016, 117, 104-113.	1.1	85
3	Fabry disease in infancy and early childhood: a systematic literature review. Genetics in Medicine, 2015, 17, 323-330.	2.4	82
4	Phenotypic characteristics of the p.Asn215Ser (p.N215S) G<scp>LA</scp> mutation in male and female patients with Fabry disease: A multicenter Fabry Registry study. Molecular Genetics & Genomic Medicine, 2018, 6, 492-503.	1.2	70
5	Fabry Disease Practice Guidelines: Recommendations of the National Society of Genetic Counselors. Journal of Genetic Counseling, 2013, 22, 555-564.	1.6	67
6	Antiproteinuric therapy and Fabry nephropathy: factors associated with preserved kidney function during agalsidase-beta therapy. Journal of Medical Genetics, 2015, 52, 860-866.	3.2	53
7	Machine learning based analytics of micro-MRI trabecular bone microarchitecture and texture in type 1 Gaucher disease. Journal of Biomechanics, 2016, 49, 1961-1968.	2.1	18
8	Treatment of Depression in Adults with Fabry Disease. JIMD Reports, 2017, 38, 13-21.	1.5	17
9	Patients' perspectives on newborn screening for later-onset lysosomal storage diseases. Molecular Genetics and Metabolism, 2016, 119, 109-114.	1.1	16
10	Climbing the Branches of a Family Tree: Diagnosis of Fragile X Syndrome. Journal of Pediatrics, 2014, 164, 1292-1295.	1.8	13
11	A Retrospective Survey Studying the Impact of Fabry Disease on Pregnancy. JIMD Reports, 2014, 21, 57-63.	1.5	10
12	The Psychosocial Impact of Carrying a Debated Variant in the GLA Gene. Journal of Genetic Counseling, 2018, 27, 217-224.	1.6	10
13	Barriers in communication and available resources to facilitate conversation about infertility with girls diagnosed with Turner syndrome. Journal of Pediatric Endocrinology and Metabolism, 2016, 29, 185-91.	0.9	7
14	Creating genetics-based infusion centers: a case study of two models. Genetics in Medicine, 2008, 10, 626-632.	2.4	2
15	Attention Deficits and ADHD Symptoms in Adults with Fabry Diseaseâ€"A Pilot Investigation. Journal of Clinical Medicine, 2021, 10, 3367.	2.4	1
16	Preliminary validation of telecounseling for depression in patients with Fabry disease. Molecular Genetics and Metabolism, 2017, 120, S19-S20.	1.1	0
17	Identification of a HEXB variant of unknown clinical significance in a family with Sandhoff disease. Molecular Genetics and Metabolism, 2018, 123, S131-S132.	1.1	0
18	ThinkGenetic: A pilot project to create an educational website/application providing increased access to information on accurate natural history, diagnosis, and treatment information with treatable genetic disorders to healthcare providers. Molecular Genetics and Metabolism, 2018, 123, S82.	1.1	0

#	Article	IF	Citations
19	Determining the disease-specific knowledge gaps in patients, family members, and caregivers living with lysosomal diseases. Molecular Genetics and Metabolism, 2019, 126, S95.	1.1	0