

# Dawn Laney

## List of Publications by Year in descending order

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

Version: 2024-02-01

19  
papers

845  
citations

933447

10  
h-index

888059

17  
g-index

36  
all docs

36  
docs citations

36  
times ranked

983  
citing authors

#	ARTICLE	IF	CITATIONS
1	Attention Deficits and ADHD Symptoms in Adults with Fabry Disease—A Pilot Investigation. <i>Journal of Clinical Medicine</i> , 2021, 10, 3367.	2.4	1
2	Determining the disease-specific knowledge gaps in patients, family members, and caregivers living with lysosomal diseases. <i>Molecular Genetics and Metabolism</i> , 2019, 126, S95.	1.1	0
3	Identification of a HEXB variant of unknown clinical significance in a family with Sandhoff disease. <i>Molecular Genetics and Metabolism</i> , 2018, 123, S131-S132.	1.1	0
4	Fabry disease revisited: Management and treatment recommendations for adult patients. <i>Molecular Genetics and Metabolism</i> , 2018, 123, 416-427.	1.1	391
5	Phenotypic characteristics of the p.Asn215Ser (p.N215S) mutation in male and female patients with Fabry disease: A multicenter Fabry Registry study. <i>Molecular Genetics &amp; Genomic Medicine</i> , 2018, 6, 492-503.	1.2	70
6	The Psychosocial Impact of Carrying a Debated Variant in the GLA Gene. <i>Journal of Genetic Counseling</i> , 2018, 27, 217-224.	1.6	10
7	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. <i>Molecular Genetics and Metabolism</i> , 2018, 123, S82.	1.1	0
8	Treatment of Depression in Adults with Fabry Disease. <i>JIMD Reports</i> , 2017, 38, 13-21.	1.5	17
9	Preliminary validation of telecounseling for depression in patients with Fabry disease. <i>Molecular Genetics and Metabolism</i> , 2017, 120, S19-S20.	1.1	0
10	Patients' perspectives on newborn screening for later-onset lysosomal storage diseases. <i>Molecular Genetics and Metabolism</i> , 2016, 119, 109-114.	1.1	16
11	Machine learning based analytics of micro-MRI trabecular bone microarchitecture and texture in type 1 Gaucher disease. <i>Journal of Biomechanics</i> , 2016, 49, 1961-1968.	2.1	18
12	The management and treatment of children with Fabry disease: A United States-based perspective. <i>Molecular Genetics and Metabolism</i> , 2016, 117, 104-113.	1.1	85
13	Barriers in communication and available resources to facilitate conversation about infertility with girls diagnosed with Turner syndrome. <i>Journal of Pediatric Endocrinology and Metabolism</i> , 2016, 29, 185-91.	0.9	7
14	Antiproteinuric therapy and Fabry nephropathy: factors associated with preserved kidney function during agalsidase-beta therapy. <i>Journal of Medical Genetics</i> , 2015, 52, 860-866.	3.2	53
15	Fabry disease in infancy and early childhood: a systematic literature review. <i>Genetics in Medicine</i> , 2015, 17, 323-330.	2.4	82
16	A Retrospective Survey Studying the Impact of Fabry Disease on Pregnancy. <i>JIMD Reports</i> , 2014, 21, 57-63.	1.5	10
17	Climbing the Branches of a Family Tree: Diagnosis of Fragile X Syndrome. <i>Journal of Pediatrics</i> , 2014, 164, 1292-1295.	1.8	13
18	Fabry Disease Practice Guidelines: Recommendations of the National Society of Genetic Counselors. <i>Journal of Genetic Counseling</i> , 2013, 22, 555-564.	1.6	67

#	ARTICLE	IF	CITATIONS
19	Creating genetics-based infusion centers: a case study of two models. <i>Genetics in Medicine</i> , 2008, 10, 626-632.	2.4	2