

# Daniela Barge-Schaapveld

## List of Publications by Year in descending order

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Version: 2024-02-01

31  
papers

958  
citations

516710

16  
h-index

454955

30  
g-index

33  
all docs

33  
docs citations

33  
times ranked

2190  
citing authors

#	ARTICLE	IF	CITATIONS
1	Early improvement in positive rather than negative emotion predicts remission from depression after pharmacotherapy. <i>European Neuropsychopharmacology</i> , 2011, 21, 241-247.	0.7	94
2	A mutation update for the <i>FLNC</i> gene in myopathies and cardiomyopathies. <i>Human Mutation</i> , 2020, 41, 1091-1111.	2.5	92
3	The Tatton-Brown-Rahman Syndrome: A clinical study of 55 individuals with de novo constitutive DNMT3A variants. <i>Wellcome Open Research</i> , 2018, 3, 46.	1.8	75
4	Effects of Antidepressant Treatment on the Quality of Daily Life. <i>Journal of Clinical Psychiatry</i> , 2002, 63, 477-485.	2.2	73
5	What Is New in Dilatation of the Ascending Aorta?. <i>Circulation</i> , 2011, 123, 924-928.	1.6	72
6	Isolated Subepicardial Right Ventricular Outflow Tract Scar in Athletes With Ventricular Tachycardia. <i>Journal of the American College of Cardiology</i> , 2017, 69, 497-507.	2.8	56
7	Aberrant Function of the C-Terminal Tail of HIST1H1E Accelerates Cellular Senescence and Causes Premature Aging. <i>American Journal of Human Genetics</i> , 2019, 105, 493-508.	6.2	48
8	The atypical 16p11.2 deletion: A not so atypical microdeletion syndrome?. <i>American Journal of Medical Genetics, Part A</i> , 2011, 155, 1066-1072.	1.2	46
9	Frameshift mutations at the C-terminus of HIST1H1E result in a specific DNA hypomethylation signature. <i>Clinical Epigenetics</i> , 2020, 12, 7.	4.1	40
10	Toward an effective exome-based genetic testing strategy in pediatric dilated cardiomyopathy. <i>Genetics in Medicine</i> , 2018, 20, 1374-1386.	2.4	36
11	Panel-Based Exome Sequencing for Neuromuscular Disorders as a Diagnostic Service. <i>Journal of Neuromuscular Diseases</i> , 2019, 6, 241-258.	2.6	32
12	Testing an mHealth Momentary Assessment Routine Outcome Monitoring Application: A Focus on Restoration of Daily Life Positive Mood States. <i>PLoS ONE</i> , 2014, 9, e115254.	2.5	31
13	Prospective risk of cancer and the influence of tobacco use in carriers of the p16-Leiden germline variant. <i>European Journal of Human Genetics</i> , 2015, 23, 711-714.	2.8	29
14	Expanding the clinical and genetic spectrum of ALPK3 variants: Phenotypes identified in pediatric cardiomyopathy patients and adults with heterozygous variants. <i>American Heart Journal</i> , 2020, 225, 108-119.	2.7	25
15	Identification of known and unknown genes associated with mitral valve prolapse using an exome slice methodology. <i>Journal of Medical Genetics</i> , 2020, 57, 843-850.	3.2	22
16	Characterization of Degenerative Mitral Valve Disease: Differences between Fibroelastic Deficiency and Barlow's Disease. <i>Journal of Cardiovascular Development and Disease</i> , 2021, 8, 23.	1.6	21
17	Clustered mutations in the GRIK2 kainate receptor subunit gene underlie diverse neurodevelopmental disorders. <i>American Journal of Human Genetics</i> , 2021, 108, 1692-1709.	6.2	18
18	Intellectual Disability and Hemizygous <i>GPD2</i> Mutation. <i>American Journal of Medical Genetics, Part A</i> , 2013, 161, 1044-1050.	1.2	16

#	ARTICLE	IF	CITATIONS
19	The first titin (c.59926 + 1G &gt; A) founder mutation associated with dilated cardiomyopathy. <i>European Journal of Heart Failure</i> , 2018, 20, 803-806.	7.1	16
20	<i>GATA6</i> mutations: Characterization of two novel patients and a comprehensive overview of the <i>GATA6</i> genotypic and phenotypic spectrum. <i>American Journal of Medical Genetics, Part A</i> , 2019, 179, 1836-1845.	1.2	16
21	Delineating the molecular and phenotypic spectrum of the <i>SETD1B</i> -related syndrome. <i>Genetics in Medicine</i> , 2021, 23, 2122-2137.	2.4	16
22	Complex genetics of radial ray deficiencies: screening of a cohort of 54 patients. <i>Genetics in Medicine</i> , 2013, 15, 195-202.	2.4	15
23	Putting genome-wide sequencing in neonates into perspective. <i>Genetics in Medicine</i> , 2019, 21, 1074-1082.	2.4	15
24	Beare-Stevenson Syndrome: Two Dutch Patients With Cerebral Abnormalities. <i>Pediatric Neurology</i> , 2011, 44, 303-307.	2.1	12
25	Consolidation of the clinical and genetic definition of a <i>SOX4</i> -related neurodevelopmental syndrome. <i>Journal of Medical Genetics</i> , 2022, 59, 1058-1068.	3.2	10
26	Autosomal dominant Marfan syndrome caused by a previously reported recessive <i>FBN1</i> variant. <i>Molecular Genetics &amp; Genomic Medicine</i> , 2019, 7, e00518.	1.2	6
27	Possible hints and pitfalls in diagnosing Peutz-Jeghers syndrome. <i>Journal of Pediatric Endocrinology and Metabolism</i> , 2018, 31, 1381-1386.	0.9	5
28	Mortality Risk Associated With Truncating Founder Mutations in Titin. <i>Circulation Genomic and Precision Medicine</i> , 2019, 12, e002436.	3.6	5
29	Distinct Metabolomic Signatures in Preclinical and Obstructive Hypertrophic Cardiomyopathy. <i>Cells</i> , 2021, 10, 2950.	4.1	5
30	Case series, Chemotherapy-induced cardiomyopathy: mind the family history!. <i>European Heart Journal - Case Reports</i> , 2021, 5, ytab333.	0.6	4
31	Pathogenic effect of a <i>TGFBR1</i> mutation in a family with Loey's "Dietz syndrome. <i>Molecular Genetics &amp; Genomic Medicine</i> , 2019, 7, e00943.	1.2	3