

# Anita de Breuk

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/991671/publications.pdf>

Version: 2024-02-01

16  
papers

260  
citations

1307594

7  
h-index

1125743

13  
g-index

16  
all docs

16  
docs citations

16  
times ranked

334  
citing authors

#	ARTICLE	IF	CITATIONS
1	Genetic Risk, Lifestyle, and Age-Related Macular Degeneration in Europe. <i>Ophthalmology</i> , 2021, 128, 1039-1049.	5.2	46
2	Integrating Metabolomics, Genomics, and Disease Pathways in Age-Related Macular Degeneration. <i>Ophthalmology</i> , 2020, 127, 1693-1709.	5.2	43
3	Implications of genetic variation in the complement system in age-related macular degeneration. <i>Progress in Retinal and Eye Research</i> , 2021, 84, 100952.	15.5	41
4	Development of a Genotype Assay for Age-Related Macular Degeneration. <i>Ophthalmology</i> , 2021, 128, 1604-1617.	5.2	38
5	Effect of rare coding variants in the CFI gene on Factor I expression levels. <i>Human Molecular Genetics</i> , 2020, 29, 2313-2324.	2.9	37
6	RETINAL HYPERREFLECTIVE FOCI IN TYPE 1 DIABETES MELLITUS. <i>Retina</i> , 2020, 40, 1565-1573.	1.7	14
7	Quantitative multiplex profiling of the complement system to diagnose complement-mediated diseases. <i>Clinical and Translational Immunology</i> , 2020, 9, e1225.	3.8	9
8	Functional Analysis of Variants in Complement Factor I Identified in Age-Related Macular Degeneration and Atypical Hemolytic Uremic Syndrome. <i>Frontiers in Immunology</i> , 2021, 12, 789897.	4.8	9
9	Systemic complement levels in patients with age-related macular degeneration carrying rare or low-frequency variants in the <i>CFH</i> gene. <i>Human Molecular Genetics</i> , 2022, 31, 455-470.	2.9	7
10	Evaluating the Occurrence of Rare Variants in the Complement Factor H Gene in Patients With Early-Onset Drusen Maculopathy. <i>JAMA Ophthalmology</i> , 2021, 139, 1218.	2.5	6
11	Genetic Risk in Families with Age-Related Macular Degeneration. <i>Ophthalmology Science</i> , 2021, 1, 100087.	2.5	5
12	Generation and characterization of human induced pluripotent stem cells (iPSCs) from three patients with age-related macular degeneration carrying rare variants in the CFH gene. <i>Stem Cell Research</i> , 2022, 60, 102669.	0.7	4
13	Generation and characterization of human induced pluripotent stem cells (iPSCs) from three individuals without age-related macular degeneration. <i>Stem Cell Research</i> , 2022, 60, 102670.	0.7	1
14	Genetic and environmental risk factors for reticular pseudodrusen in the EUGENDA study.. <i>Molecular Vision</i> , 2021, 27, 757-767.	1.1	0
15	Generation of an iPSC line (SCTCi015-A) and isogenic control line (SCTCi015-A-1) from an age-related macular degeneration patient carrying the variant c.355G>A in the CFI gene. <i>Stem Cell Research</i> , 2022, 62, 102796.	0.7	0
16	Generation of an iPSC line (SCTCi014-A) and isogenic control line (SCTCi014-A-1) from an age-related macular degeneration patient carrying the variant c.355G>A in the CFI gene. <i>Stem Cell Research</i> , 2022, 62, 102797.	0.7	0