

# Bernhard H F Weber

## List of Publications by Year in descending order

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Version: 2024-02-01

76  
papers

4,886  
citations

147566

31  
h-index

114278

63  
g-index

76  
all docs

76  
docs citations

76  
times ranked

7853  
citing authors

#	ARTICLE	IF	CITATIONS
1	Incidence, progression and risk factors of age-related macular degeneration in 35â€“95-year-old individuals from three jointly designed German cohort studies. <i>BMJ Open Ophthalmology</i> , 2022, 7, e000912.	0.8	7
2	Vitronectin and Its Interaction with PAI-1 Suggests a Functional Link to Vascular Changes in AMD Pathobiology. <i>Cells</i> , 2022, 11, 1766.	1.8	2
3	Epistatic interactions of genetic loci associated with age-related macular degeneration. <i>Scientific Reports</i> , 2021, 11, 13114.	1.6	4
4	Assigning Co-Regulated Human Genes and Regulatory Gene Clusters. <i>Cells</i> , 2021, 10, 2395.	1.8	4
5	OUP accepted manuscript. <i>Human Molecular Genetics</i> , 2021, , .	1.4	1
6	Performance of Breast Cancer Polygenic Risk Scores in 760 Female <i>CHEK2</i> Germline Mutation Carriers. <i>Journal of the National Cancer Institute</i> , 2021, 113, 893-899.	3.0	21
7	Insights into the loss of the Y chromosome with age in control individuals and in patients with age-related macular degeneration using genotyping microarray data. <i>Human Genetics</i> , 2020, 139, 401-407.	1.8	18
8	Complement C3 Inhibitor Pegcetacoplan for Geographic Atrophy Secondary to Age-Related Macular Degeneration. <i>Ophthalmology</i> , 2020, 127, 186-195.	2.5	306
9	Altered Protein Function Caused by AMD-associated Variant rs704 Links Vitronectin to Disease Pathology. , 2020, 61, 2.		14
10	Pleiotropic Locus 15q24.1 Reveals a Gender-Specific Association with Neovascular but Not Atrophic Age-Related Macular Degeneration (AMD). <i>Cells</i> , 2020, 9, 2257.	1.8	5
11	A mega-analysis of expression quantitative trait loci in retinal tissue. <i>PLoS Genetics</i> , 2020, 16, e1008934.	1.5	22
12	Genome-wide association meta-analysis for early age-related macular degeneration highlights novel loci and insights for advanced disease. <i>BMC Medical Genomics</i> , 2020, 13, 120.	0.7	56
13	Clinical Heterogeneity in Autosomal Recessive Bestrophinopathy with Biallelic Mutations in the BEST1 Gene. <i>International Journal of Molecular Sciences</i> , 2020, 21, 9353.	1.8	8
14	Learning from Fifteen Years of Genome-Wide Association Studies in Age-Related Macular Degeneration. <i>Cells</i> , 2020, 9, 2267.	1.8	18
15	Resolving the dark matter of ABCA4 for 1054 Stargardt disease probands through integrated genomics and transcriptomics. <i>Genetics in Medicine</i> , 2020, 22, 1235-1246.	1.1	92
16	Mutation-Dependent Pathomechanisms Determine the Phenotype in the Bestrophinopathies. <i>International Journal of Molecular Sciences</i> , 2020, 21, 1597.	1.8	20
17	Correlating Adaptive Optics Images to Clinical Findings in Juvenile Macular Dystrophy with Hypotrichosis in Siblings with Homozygous <i>CDH3</i> Pathogenic Variation. <i>Ophthalmic Research</i> , 2020, 63, 141-151.	1.0	1
18	A transcriptome-wide association study based on 27 tissues identifies 106 genes potentially relevant for disease pathology in age-related macular degeneration. <i>Scientific Reports</i> , 2020, 10, 1584.	1.6	39

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19	A Circulating MicroRNA Profile in a Laser-Induced Mouse Model of Choroidal Neovascularization. <i>International Journal of Molecular Sciences</i> , 2020, 21, 2689.	1.8	8
20	Investigating the effects of additional truncating variants in DNA-repair genes on breast cancer risk in BRCA1-positive women. <i>BMC Cancer</i> , 2019, 19, 787.	1.1	10
21	Assessment of Novel Genome-Wide Significant Gene Loci and Lesion Growth in Geographic Atrophy Secondary to Age-Related Macular Degeneration. <i>JAMA Ophthalmology</i> , 2019, 137, 867.	1.4	28
22	The Y227N mutation affects bestrophin-1 protein stability and impairs sperm function in a mouse model of Best vitelliform macular dystrophy. <i>Biology Open</i> , 2019, 8, .	0.6	5
23	Cost-effective molecular inversion probe-based <i>ABCA4</i> sequencing reveals deep intronic variants in Stargardt disease. <i>Human Mutation</i> , 2019, 40, 1749-1759.	1.1	39
24	Retinal Layer Thicknesses in Early Age-Related Macular Degeneration: Results From the German AugUR Study. , 2019, 60, 1581.		34
25	Identification of the retinoschisin-binding site on the retinal Na/K-ATPase. <i>PLoS ONE</i> , 2019, 14, e0216320.	1.1	13
26	On the differences between mega- and meta-imputation and analysis exemplified on the genetics of age-related macular degeneration. <i>Genetic Epidemiology</i> , 2019, 43, 559-576.	0.6	5
27	The agonistic TSPO ligand XBD173 attenuates the glial response thereby protecting inner retinal neurons in a murine model of retinal ischemia. <i>Journal of Neuroinflammation</i> , 2019, 16, 43.	3.1	35
28	Cell-Type-Specific Complement Expression in the Healthy and Diseased Retina. <i>Cell Reports</i> , 2019, 29, 2835-2848.e4.	2.9	81
29	Y chromosome mosaicism is associated with age-related macular degeneration. <i>European Journal of Human Genetics</i> , 2019, 27, 36-41.	1.4	49
30	Gene panel testing of 5589 <i>BRCA1/2</i> -negative index patients with breast cancer in a routine diagnostic setting: results of the German Consortium for Hereditary Breast and Ovarian Cancer. <i>Cancer Medicine</i> , 2018, 7, 1349-1358.	1.3	126
31	BEST1 protein stability and degradation pathways differ between autosomal dominant Best disease and autosomal recessive bestrophinopathy accounting for the distinct retinal phenotypes. <i>Human Molecular Genetics</i> , 2018, 27, 1630-1641.	1.4	32
32	Recurrent structural variation, clustered sites of selection, and disease risk for the complement factor H ( <i>CFH</i> ) gene family. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2018, 115, E4433-E4442.	3.3	43
33	A mega-analysis of expression quantitative trait loci (eQTL) provides insight into the regulatory architecture of gene expression variation in liver. <i>Scientific Reports</i> , 2018, 8, 5865.	1.6	52
34	A Deep Learning Algorithm for Prediction of Age-Related Eye Disease Study Severity Scale for Age-Related Macular Degeneration from Color Fundus Photography. <i>Ophthalmology</i> , 2018, 125, 1410-1420.	2.5	365
35	Mutational spectrum in a worldwide study of 29,700 families with <i>BRCA1</i> or <i>BRCA2</i> mutations. <i>Human Mutation</i> , 2018, 39, 593-620.	1.1	224
36	Pleiotropic Effects of Risk Factors in Age-Related Macular Degeneration and Seemingly Unrelated Complex Diseases. <i>Advances in Experimental Medicine and Biology</i> , 2018, 1074, 247-255.	0.8	1

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37	Views of ophthalmologists on the genetics of age-related macular degeneration: Results of a qualitative study. <i>PLoS ONE</i> , 2018, 13, e0209328.	1.1	9
38	Pathomechanism of mutated and secreted retinoschisin in X-linked juvenile retinoschisis. <i>Experimental Eye Research</i> , 2018, 177, 23-34.	1.2	14
39	Evaluation of serum sphingolipids and the influence of genetic risk factors in age-related macular degeneration. <i>PLoS ONE</i> , 2018, 13, e0200739.	1.1	19
40	Effects of genetic variants in the TSPO gene on protein structure and stability. <i>PLoS ONE</i> , 2018, 13, e0195627.	1.1	19
41	Prevalence of pathogenic BRCA1/2 germline mutations among 802 women with unilateral triple-negative breast cancer without family cancer history. <i>BMC Cancer</i> , 2018, 18, 265.	1.1	84
42	Choroidal Flow Signal in Late-Onset Stargardt Disease and Age-Related Macular Degeneration: An OCT-Angiography Study. , 2018, 59, AMD122.		38
43	Investigating the modulation of genetic effects on late AMD by age and sex: Lessons learned and two additional loci. <i>PLoS ONE</i> , 2018, 13, e0194321.	1.1	19
44	On the impact of different approaches to classify age-related macular degeneration: Results from the German AugUR study. <i>Scientific Reports</i> , 2018, 8, 8675.	1.6	31
45	Twenty-year follow-up of a familial case of PTH1R-associated primary failure of tooth eruption. <i>American Journal of Orthodontics and Dentofacial Orthopedics</i> , 2017, 151, 598-606.	0.8	6
46	Recombinant Haplotypes Narrow the ARMS2/HTRA1 Association Signal for Age-Related Macular Degeneration. <i>Genetics</i> , 2017, 205, 919-924.	1.2	65
47	Retinoschisin is linked to retinal Na/K-ATPase signaling and localization. <i>Molecular Biology of the Cell</i> , 2017, 28, 2178-2189.	0.9	26
48	The X-linked juvenile retinoschisis protein retinoschisin is a novel regulator of mitogen-activated protein kinase signalling and apoptosis in the retina. <i>Journal of Cellular and Molecular Medicine</i> , 2017, 21, 768-780.	1.6	18
49	Genetic pleiotropy between age-related macular degeneration and 16 complex diseases and traits. <i>Genome Medicine</i> , 2017, 9, 29.	3.6	52
50	An Eye on Age-Related Macular Degeneration: The Role of MicroRNAs in Disease Pathology. <i>Molecular Diagnosis and Therapy</i> , 2017, 21, 31-43.	1.6	78
51	Mutations in the Genes for Interphotoreceptor Matrix Proteoglycans, IMPG1 and IMPG2, in Patients with Vitelliform Macular Lesions. <i>Genes</i> , 2017, 8, 170.	1.0	24
52	Mutation Spectrum of the <i>ABCA4</i> Gene in 335 Stargardt Disease Patients From a Multicenter German Cohort—Impact of Selected Deep Intronic Variants and Common SNPs. , 2017, 58, 394.		104
53	Progression of Late-Onset Stargardt Disease. , 2016, 57, 5186.		57
54	Distinct Genetic Risk Profile of the Rapidly Progressing Diffuse-Trickling Subtype of Geographic Atrophy in Age-Related Macular Degeneration (AMD). , 2016, 57, 2463.		22

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55	Features of Age-Related Macular Degeneration in the General Adults and Their Dependency on Age, Sex, and Smoking: Results from the German KORA Study. PLoS ONE, 2016, 11, e0167181.	1.1	27
56	Prevalence of <i>BRCA1/2</i> germline mutations in 21401 families with breast and ovarian cancer. Journal of Medical Genetics, 2016, 53, 465-471.	1.5	179
57	Differentiation of murine models of $\alpha$ -negative ERG by single and repetitive light stimuli. Documenta Ophthalmologica, 2016, 132, 101-109.	1.0	17
58	Long-Term Macular Changes in the First Proband of Autosomal Dominant Vitreoretinopathy (ADVIRC) Due to a Newly Identified Mutation in <i>BEST1</i> . Ophthalmic Genetics, 2016, 37, 102-108.	0.5	16
59	Multiallelic copy number variation in the complement component 4A (C4A) gene is associated with late-stage age-related macular degeneration (AMD). Journal of Neuroinflammation, 2016, 13, 81.	3.1	31
60	Mutations in <i>EXOSC2</i> are associated with a novel syndrome characterised by retinitis pigmentosa, progressive hearing loss, premature ageing, short stature, mild intellectual disability and distinctive gestalt. Journal of Medical Genetics, 2016, 53, 419-425.	1.5	69
61	A large genome-wide association study of age-related macular degeneration highlights contributions of rare and common variants. Nature Genetics, 2016, 48, 134-143.	9.4	1,167
62	In-Depth Functional Diagnostics of Mouse Models by Single-Flash and Flicker Electroretinograms without Adapting Background Illumination. Advances in Experimental Medicine and Biology, 2016, 854, 619-625.	0.8	5
63	Monoallelic <i>ABCA4</i> Mutations Appear Insufficient to Cause Retinopathy: A Quantitative Autofluorescence Study. , 2015, 56, 8179.		38
64	Migraine and Vasospasm in Glaucoma: Age-Related Evaluation of 2027 Patients With Glaucoma or Ocular Hypertension. , 2015, 56, 7999.		51
65	The genetics of age-related macular degeneration (AMD) – Novel targets for designing treatment options?. European Journal of Pharmaceutics and Biopharmaceutics, 2015, 95, 194-202.	2.0	33
66	Variation of the group 5 grass pollen allergen content of airborne pollen in relation to geographic location and time in season. Journal of Allergy and Clinical Immunology, 2015, 136, 87-95.e6.	1.5	155
67	Reticular Pseudodrusen in Sorsby Fundus Dystrophy. Ophthalmology, 2015, 122, 1555-1562.	2.5	58
68	Common synonymous variants in <i>ABCA4</i> are protective for chloroquine induced maculopathy (toxic Tj ETQq0 0 0 rBT /Overlock 10 Tf	0.6	36
69	Sorsby Fundus Dystrophy: Novel Mutations, Novel Phenotypic Characteristics, and Treatment Outcomes. , 2015, 56, 2664.		55
70	Synonymous variants in <i>HTRA1</i> implicated in AMD susceptibility impair its capacity to regulate TGF- $\beta$ signaling. Human Molecular Genetics, 2015, 24, 6361-6373.	1.4	42
71	Clinical and Genetic Factors Associated with Progression of Geographic Atrophy Lesions in Age-Related Macular Degeneration. PLoS ONE, 2015, 10, e0126636.	1.1	61
72	The Role of the Complement System in Age-Related Macular Degeneration. Deutsches A&#x0308;rztblatt International, 2014, 111, 133-8.	0.6	47

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73	A Novel Antibody against Human Properdin Inhibits the Alternative Complement System and Specifically Detects Properdin from Blood Samples. PLoS ONE, 2014, 9, e96371.	1.1	44
74	A Circulating MicroRNA Profile Is Associated with Late-Stage Neovascular Age-Related Macular Degeneration. PLoS ONE, 2014, 9, e107461.	1.1	62
75	Modelling the Genetic Risk in Age-Related Macular Degeneration. PLoS ONE, 2012, 7, e37979.	1.1	79
76	Release of Bet v 1 from birch pollen from 5 European countries. Results from the HIALINE study. Atmospheric Environment, 2012, 55, 496-505.	1.9	141