## HélÃ"ne Choquet

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/2317655/publications.pdf

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32 papers 4,060 citations

394421 19 h-index 395702 33 g-index

39 all docs

39 docs citations

39 times ranked 7722 citing authors

#	Article	IF	Citations
1	Ancestry- and sex-specific effects underlying inguinal hernia susceptibility identified in a multiethnic genome-wide association study meta-analysis. Human Molecular Genetics, 2022, 31, 2279-2293.	2.9	6
2	Genome-Wide Association Study Identifies Two Common Loci Associated with Pigment Dispersion Syndrome/Pigmentary Glaucoma and Implicates Myopia in its Development. Ophthalmology, 2022, 129, 626-636.	5.2	10
3	Genome-wide association study of actinic keratosis identifies new susceptibility loci implicated in pigmentation and immune regulation pathways. Communications Biology, 2022, 5, 386.	4.4	9
4	Genome-wide meta-analysis identifies 127 open-angle glaucoma loci with consistent effect across ancestries. Nature Communications, 2021, 12, 1258.	12.8	196
5	Cigarette smoking behaviors and the importance of ethnicity and genetic ancestry. Translational Psychiatry, 2021, 11, 120.	4.8	9
6	A multi-ethnic genome-wide association study implicates collagen matrix integrity and cell differentiation pathways in keratoconus. Communications Biology, 2021, 4, 266.	4.4	36
7	Whole-genome sequencing association analysis of quantitative red blood cell phenotypes: The NHLBI TOPMed program. American Journal of Human Genetics, 2021, 108, 874-893.	6.2	28
8	Genetic variation affects morphological retinal phenotypes extracted from UK Biobank optical coherence tomography images. PLoS Genetics, 2021, 17, e1009497.	<b>3.</b> 5	50
9	A large multiethnic GWAS meta-analysis of cataract identifies new risk loci and sex-specific effects. Nature Communications, 2021, 12, 3595.	12.8	39
10	New and sex-specific migraine susceptibility loci identified from a multiethnic genome-wide meta-analysis. Communications Biology, 2021, 4, 864.	4.4	21
11	Transcriptome-Wide Association Study of Blood Cell Traits in African Ancestry and Hispanic/Latino Populations. Genes, 2021, 12, 1049.	2.4	11
12	GLIS1 regulates trabecular meshwork function and intraocular pressure and is associated with glaucoma in humans. Nature Communications, 2021, 12, 4877.	12.8	20
13	Clinical implications of recent advances in primary open-angle glaucoma genetics. Eye, 2020, 34, 29-39.	2.1	48
14	The Polygenic and Monogenic Basis of Blood Traits and Diseases. Cell, 2020, 182, 1214-1231.e11.	28.9	388
15	Trans-ethnic and Ancestry-Specific Blood-Cell Genetics in 746,667 Individuals from 5 Global Populations. Cell, 2020, 182, 1198-1213.e14.	28.9	353
16	A multiethnic genome-wide analysis of 44,039 individuals identifies 41 new loci associated with central corneal thickness. Communications Biology, 2020, 3, 301.	4.4	28
17	Meta-analysis of 542,934 subjects of European ancestry identifies new genes and mechanisms predisposing to refractive error and myopia. Nature Genetics, 2020, 52, 401-407.	21.4	180
18	Meta-Analysis of 26 638 Individuals Identifies Two Genetic Loci Associated With Left Ventricular Ejection Fraction. Circulation Genomic and Precision Medicine, 2020, 13, e002804.	3.6	10

#	Article	IF	Citations
19	Functional validity, role, and implications of heavy alcohol consumption genetic loci. Science Advances, 2020, 6, eaay5034.	10.3	47
20	Genetic and environmental factors underlying keratinocyte carcinoma risk. JCI Insight, 2020, 5, .	5.0	17
21	Genetic ancestry, skin pigmentation, and the risk of cutaneous squamous cell carcinoma in Hispanic/Latino and non-Hispanic white populations. Communications Biology, 2020, 3, 765.	4.4	6
22	Genome-wide Genotyping of Cerebral Cavernous Malformation Type 1 Individuals to Identify Genetic Modifiers of Disease Severity. Methods in Molecular Biology, 2020, 2152, 77-84.	0.9	4
23	Contribution of rare coding mutations in CD36 to type 2 diabetes and cardio-metabolic complications. Scientific Reports, 2019, 9, 17123.	3.3	8
24	Use of >100,000 NHLBI Trans-Omics for Precision Medicine (TOPMed) Consortium whole genome sequences improves imputation quality and detection of rare variant associations in admixed African and Hispanic/Latino populations. PLoS Genetics, 2019, 15, e1008500.	3.5	203
25	Association studies of up to 1.2 million individuals yield new insights into the genetic etiology of tobacco and alcohol use. Nature Genetics, 2019, 51, 237-244.	21.4	1,307
26	Common Mitochondrial Haplogroups and Cutaneous Squamous Cell Carcinoma Risk. Cancer Epidemiology Biomarkers and Prevention, 2018, 27, 838-841.	2.5	2
27	A Large Multiethnic Genome-Wide Association Study of Adult Body Mass Index Identifies Novel Loci. Genetics, 2018, 210, 499-515.	2.9	131
28	A multiethnic genome-wide association study of primary open-angle glaucoma identifies novel risk loci. Nature Communications, 2018, 9, 2278.	12.8	124
29	A large multi-ethnic genome-wide association study identifies novel genetic loci for intraocular pressure. Nature Communications, 2017, 8, 2108.	12.8	80
30	Cytochrome P450 and matrix metalloproteinase genetic modifiers of disease severity in Cerebral Cavernous Malformation type 1. Free Radical Biology and Medicine, 2016, 92, 100-109.	2.9	47
31	Polymorphisms in Inflammatory and Immune Response Genes Associated with Cerebral Cavernous Malformation Type 1 Severity. Cerebrovascular Diseases, 2014, 38, 433-440.	1.7	57
32	Dysfunction of lipid sensor GPR120 leads to obesity in both mouse and human. Nature, 2012, 483, 350-354.	27.8	572