Anders Albrechtsen

List of Publications by Year in descending order

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26567 15218 18,916 129 56 126 citations h-index papers

g-index 149 149 149 23868 docs citations times ranked citing authors all docs

#	Article	IF	CITATIONS
1	ANGSD: Analysis of Next Generation Sequencing Data. BMC Bioinformatics, 2014, 15, 356.	1.2	1,935
2	Genotype and SNP calling from next-generation sequencing data. Nature Reviews Genetics, 2011, 12, 443-451.	7.7	1,238
3	Upper Palaeolithic Siberian genome reveals dual ancestry of Native Americans. Nature, 2014, 505, 87-91.	13.7	821
4	Ancient human genome sequence of an extinct Palaeo-Eskimo. Nature, 2010, 463, 757-762.	13.7	750
5	Recalibrating Equus evolution using the genome sequence of an early Middle Pleistocene horse. Nature, 2013, 499, 74-78.	13.7	717
6	An Aboriginal Australian Genome Reveals Separate Human Dispersals into Asia. Science, 2011, 334, 94-98.	6.0	675
7	Estimating Individual Admixture Proportions from Next Generation Sequencing Data. Genetics, 2013, 195, 693-702.	1.2	515
8	The genome of a Late Pleistocene human from a Clovis burial site in western Montana. Nature, 2014, 506, 225-229.	13.7	500
9	Genomic evidence for the Pleistocene and recent population history of Native Americans. Science, 2015, 349, aab3884.	6.0	449
10	A genomic history of Aboriginal Australia. Nature, 2016, 538, 207-214.	13.7	439
11	Low Physical Activity Accentuates the Effect of the <i>FTO</i> rs9939609 Polymorphism on Body Fat Accumulation. Diabetes, 2008, 57, 95-101.	0.3	431
12	Inferring Population Structure and Admixture Proportions in Low-Depth NGS Data. Genetics, 2018, 210, 719-731.	1.2	426
13	Genetic variant near IRS1 is associated with type 2 diabetes, insulin resistance and hyperinsulinemia. Nature Genetics, 2009, 41, 1110-1115.	9.4	418
14	Greenlandic Inuit show genetic signatures of diet and climate adaptation. Science, 2015, 349, 1343-1347.	6.0	397
15	A common Greenlandic TBC1D4 variant confers muscle insulin resistance and type 2 diabetes. Nature, 2014, 512, 190-193.	13.7	338
16	SNP Calling, Genotype Calling, and Sample Allele Frequency Estimation from New-Generation Sequencing Data. PLoS ONE, 2012, 7, e37558.	1,1	336
17	Ascertainment Biases in SNP Chips Affect Measures of Population Divergence. Molecular Biology and Evolution, 2010, 27, 2534-2547.	3.5	317
18	Terminal Pleistocene Alaskan genome reveals first founding population of Native Americans. Nature, 2018, 553, 203-207.	13.7	304

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19	Resequencing of 200 human exomes identifies an excess of low-frequency non-synonymous coding variants. Nature Genetics, 2010, 42, 969-972.	9.4	297
20	Genomic structure in Europeans dating back at least 36,200 years. Science, 2014, 346, 1113-1118.	6.0	287
21	The genetic prehistory of the New World Arctic. Science, 2014, 345, 1255832.	6.0	264
22	Ancient genomes show social and reproductive behavior of early Upper Paleolithic foragers. Science, 2017, 358, 659-662.	6.0	263
23	Prehistoric genomes reveal the genetic foundation and cost of horse domestication. Proceedings of the National Academy of Sciences of the United States of America, 2014, 111, E5661-9.	3.3	260
24	The ancestry and affiliations of Kennewick Man. Nature, 2015, 523, 455-458.	13.7	241
25	Ancient genomes revisit the ancestry of domestic and Przewalski's horses. Science, 2018, 360, 111-114.	6.0	241
26	Studies of Association of Variants Near the <i>HHEX</i> , <i>CDKN2A/B</i> , and <i>IGF2BP2</i> Genes With Type 2 Diabetes and Impaired Insulin Release in 10,705 Danish Subjects. Diabetes, 2007, 56, 3105-3111.	0.3	230
27	Genomic Analyses from Non-invasive Prenatal Testing Reveal Genetic Associations, Patterns of Viral Infections, and Chinese Population History. Cell, 2018, 175, 347-359.e14.	13.5	213
28	Calculation of Tajima's D and other neutrality test statistics from low depth next-generation sequencing data. BMC Bioinformatics, 2013, 14, 289.	1.2	211
29	Quantifying Population Genetic Differentiation from Next-Generation Sequencing Data. Genetics, 2013, 195, 979-992.	1.2	187
30	Ancient genomic changes associated with domestication of the horse. Science, 2017, 356, 442-445.	6.0	185
31	Speciation with gene flow in equids despite extensive chromosomal plasticity. Proceedings of the National Academy of Sciences of the United States of America, 2014, 111, 18655-18660.	3.3	183
32	Weighting sequence variants based on their annotation increases power of whole-genome association studies. Nature Genetics, 2016, 48, 314-317.	9.4	178
33	Estimation of allele frequency and association mapping using next-generation sequencing data. BMC Bioinformatics, 2011, 12, 231.	1.2	170
34	Next-generation biology: Sequencing and data analysis approaches for non-model organisms. Marine Genomics, 2016, 30, 3-13.	0.4	164
35	Evolutionary Genomics and Conservation of the Endangered Przewalski's Horse. Current Biology, 2015, 25, 2577-2583.	1.8	161
36	Loss-of-function variants in ADCY3 increase risk of obesity and type 2 diabetes. Nature Genetics, 2018, 50, 172-174.	9.4	156

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37	The GCKR rs780094 polymorphism is associated with elevated fasting serum triacylglycerol, reduced fasting and OGTT-related insulinaemia, and reduced risk of type 2 diabetes. Diabetologia, 2007, 51, 70-75.	2.9	153
38	Association Testing of Novel Type 2 Diabetes Risk Alleles in the <i>JAZF1</i> , <i>CDC123</i> / <i>CAMK1D</i> , <i>TSPAN8</i> , <i>THADA</i> , <i>ADAMTS9</i> , and <i>NOTCH2</i> Loci With Insulin Release, Insulin Sensitivity, and Obesity in a Population-Based Sample of 4,516 Glucose-Tolerant Middle-Aged Danes. Diabetes, 2008, 57, 2534-2540.	0.3	151
39	Natural Selection Affects Multiple Aspects of Genetic Variation at Putatively Neutral Sites across the Human Genome. PLoS Genetics, 2011, 7, e1002326.	1.5	146
40	Population genomics of the Viking world. Nature, 2020, 585, 390-396.	13.7	143
41	Darwinian and demographic forces affecting human protein coding genes. Genome Research, 2009, 19, 838-849.	2.4	139
42	Tracking the origins of Yakutian horses and the genetic basis for their fast adaptation to subarctic environments. Proceedings of the National Academy of Sciences of the United States of America, 2015, 112, E6889-97.	3.3	139
43	The origin and evolution of maize in the Southwestern United States. Nature Plants, 2015, 1, 14003.	4.7	138
44	Whole-Exome Sequencing of 2,000 Danish Individuals and the Role of Rare Coding Variants in Type 2 Diabetes. American Journal of Human Genetics, 2013, 93, 1072-1086.	2.6	124
45	Detection of internal N7-methylguanosine (m7G) RNA modifications by mutational profiling sequencing. Nucleic Acids Research, 2019, 47, e126-e126.	6.5	124
46	Natural Selection and the Distribution of Identity-by-Descent in the Human Genome. Genetics, 2010, 186, 295-308.	1.2	119
47	Exome sequencing-driven discovery of coding polymorphisms associated with common metabolic phenotypes. Diabetologia, 2013, 56, 298-310.	2.9	119
48	Genetic Architecture of Vitamin B12 and Folate Levels Uncovered Applying Deeply Sequenced Large Datasets. PLoS Genetics, 2013, 9, e1003530.	1.5	112
49	Studies of the relationship between the ENPP1 K121Q polymorphism and type 2 diabetes, insulin resistance and obesity in 7,333 Danish white subjects. Diabetologia, 2006, 49, 2097-2104.	2.9	111
50	Archaic adaptive introgression in <i>TBX15/WARS2</i> . Molecular Biology and Evolution, 2017, 34, msw283.	3.5	101
51	Relatedness mapping and tracts of relatedness for genomeâ€wide data in the presence of linkage disequilibrium. Genetic Epidemiology, 2009, 33, 266-274.	0.6	99
52	Estimating inbreeding coefficients from NGS data: Impact on genotype calling and allele frequency estimation. Genome Research, 2013, 23, 1852-1861.	2.4	89
53	Uncovering the Genetic History of the Present-Day Greenlandic Population. American Journal of Human Genetics, 2015, 96, 54-69.	2.6	85
54	Two ancient human genomes reveal Polynesian ancestry among the indigenous Botocudos of Brazil. Current Biology, 2014, 24, R1035-R1037.	1.8	73

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55	Allele frequencyâ€free inference of close familial relationships from genotypes or lowâ€depth sequencing data. Molecular Ecology, 2019, 28, 35-48.	2.0	73
56	Powerful Inference with the D-Statistic on Low-Coverage Whole-Genome Data. G3: Genes, Genomes, Genetics, 2018, 8, 551-566.	0.8	71
57	Studies of the CommonDIO2Thr92Ala Polymorphism and Metabolic Phenotypes in 7342 Danish White Subjects. Journal of Clinical Endocrinology and Metabolism, 2007, 92, 363-366.	1.8	62
58	Large BRCA1 and BRCA2 genomic rearrangements in Danish high risk breast-ovarian cancer families. Breast Cancer Research and Treatment, 2009, 115, 315-323.	1.1	61
59	RelateAdmix: a software tool for estimating relatedness between admixed individuals. Bioinformatics, 2014, 30, 1027-1028.	1.8	61
60	Combined Analyses of 20 Common Obesity Susceptibility Variants. Diabetes, 2010, 59, 1667-1673.	0.3	55
61	The Effect of an Extreme and Prolonged Population Bottleneck on Patterns of Deleterious Variation: Insights from the Greenlandic Inuit. Genetics, 2017, 205, 787-801.	1.2	54
62	Combined analysis of 19 common validated type 2 diabetes susceptibility gene variants shows moderate discriminative value and no evidence of gene–gene interaction. Diabetologia, 2009, 52, 1308-1314.	2.9	53
63	Association Testing for Nextâ€Generation Sequencing Data Using Score Statistics. Genetic Epidemiology, 2012, 36, 430-437.	0.6	53
64	Genomic diversity and novel genome-wide association with fruit morphology in Capsicum, from 746k polymorphic sites. Scientific Reports, 2019, 9, 10067.	1.6	53
65	Impact of polymorphisms in WFS1 on prediabetic phenotypes in a population-based sample of middle-aged people with normal and abnormal glucose regulation. Diabetologia, 2008, 51, 1646-1652.	2.9	44
66	Studies of the association of the GNB3 825C>T polymorphism with components of the metabolic syndrome in white Danes. Diabetologia, 2006, 49, 75-82.	2.9	43
67	Evaluation of model fit of inferred admixture proportions. Molecular Ecology Resources, 2020, 20, 936-949.	2.2	43
68	Variation in the peroxisome proliferator-activated receptor \hat{l} gene in relation to common metabolic traits in 7,495 middle-aged white people. Diabetologia, 2007, 50, 1201-1208.	2.9	42
69	A method for detecting IBD regions simultaneously in multiple individuals—with applications to disease genetics. Genome Research, 2011, 21, 1168-1180.	2.4	42
70	High genetic diversity and low differentiation reflect the ecological versatility of the African leopard. Current Biology, 2021, 31, 1862-1871.e5.	1.8	41
71	Studies of associations between the Arg389Gly polymorphism of the \hat{l}^21 -adrenergic receptor gene (ADRB1) and hypertension and obesity in 7677 Danish white subjects. Diabetic Medicine, 2007, 24, 392-397.	1.2	40
72	Estimating IBD tracts from low coverage NGS data. Bioinformatics, 2016, 32, 2096-2102.	1.8	36

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73	Variants in the $5\hat{a} \in \mathbb{Z}^2$ region of the neuropeptide Y receptor Y2 gene (NPY2R) are associated with obesity in 5,971 white subjects. Diabetologia, 2006, 49, 2653-2658.	2.9	35
74	Novel de novo BRCA2mutation in a patient with a family history of breast cancer. BMC Medical Genetics, 2008, 9, 58.	2.1	33
75	The â^'250G>A Promoter Variant in Hepatic Lipase Associates with Elevated Fasting Serum High-Density Lipoprotein Cholesterol Modulated by Interaction with Physical Activity in a Study of 16,156 Danish Subjects. Journal of Clinical Endocrinology and Metabolism, 2008, 93, 2294-2299.	1.8	33
76	Segregation distortion in chicken and the evolutionary consequences of female meiotic drive in birds. Heredity, 2010, 105, 290-298.	1.2	33
77	Life-threatening viral disease in a novel form of autosomal recessive <i>IFNAR2</i> deficiency in the Arctic. Journal of Experimental Medicine, 2022, 219, .	4.2	33
78	A southern African origin and cryptic structure in the highly mobile plains zebra. Nature Ecology and Evolution, 2018, 2, 491-498.	3.4	32
79	fastNGSadmix: admixture proportions and principal component analysis of a single NGS sample. Bioinformatics, 2017, 33, 3148-3150.	1.8	31
80	A variant in the G6PC2/ABCB11 locus is associated with increased fasting plasma glucose, increased basal hepatic glucose production and increased insulin release after oral and intravenous glucose loads. Diabetologia, 2009, 52, 2122-2129.	2.9	29
81	Variation and association to diabetes in 2000 full mtDNA sequences mined from an exome study in a Danish population. European Journal of Human Genetics, 2014, 22, 1040-1045.	1.4	26
82	Ancestryâ€specific association mapping in admixed populations. Genetic Epidemiology, 2019, 43, 506-521.	0.6	26
83	Testing for Hardy–Weinberg equilibrium in structured populations using genotype or lowâ€depth next generation sequencing data. Molecular Ecology Resources, 2019, 19, 1144-1152.	2.2	26
84	Functional <i>SOCS1 </i> polymorphisms are associated with variation in obesity in whites. Diabetes, Obesity and Metabolism, 2009, 11, 196-203.	2.2	25
85	Interleukin-6 autoantibodies are involved in the pathogenesis of a subset of type 2 diabetes. Journal of Endocrinology, 2010, 204, 265-273.	1.2	25
86	Genetics of Type 2 Diabetes: the Power of Isolated Populations. Current Diabetes Reports, 2016, 16, 65.	1.7	25
87	The Validation and Assessment of Machine Learning: A Game of Prediction from High-Dimensional Data. PLoS ONE, 2009, 4, e6287.	1.1	22
88	Genetic analysis of the estrogen-related receptor \hat{l}_{\pm} and studies of association with obesity and type 2 diabetes. International Journal of Obesity, 2007, 31, 365-370.	1.6	21
89	DamMet: ancient methylome mapping accounting for errors, true variants, and post-mortem DNA damage. GigaScience, 2019, 8, .	3.3	20
90	Identification of Novel Genetic Determinants of Erythrocyte Membrane Fatty Acid Composition among Greenlanders. PLoS Genetics, 2016, 12, e1006119.	1.5	20

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91	A common Greenlandic Inuit BRCA1 RING domain founder mutation. Breast Cancer Research and Treatment, 2009, 115, 69-76.	1.1	19
92	Genetic screening of the FLCN gene identify six novel variants and a Danish founder mutation. Journal of Human Genetics, 2017, 62, 151-157.	1.1	19
93	Joint identification of sex and sexâ€linked scaffolds in nonâ€model organisms using low depth sequencing data. Molecular Ecology Resources, 2022, 22, 458-467.	2.2	19
94	Family and Population-Based Studies of Variation within the Ghrelin Receptor Locus in Relation to Measures of Obesity. PLoS ONE, 2010, 5, e10084.	1.1	18
95	Genetic and environmental determinants of 25-hydroxyvitamin D levels in multiple sclerosis. Multiple Sclerosis Journal, 2015, 21, 1414-1422.	1.4	18
96	Large-scale inference of population structure in presence of missingness using PCA. Bioinformatics, 2021, 37, 1868-1875.	1.8	17
97	Founder Effect of the <i>RET^{C611Y}</i> Mutation in Multiple Endocrine Neoplasia 2A in Denmark: A Nationwide Study. Thyroid, 2017, 27, 1505-1510.	2.4	16
98	Identification of novel high-impact recessively inherited type 2 diabetes risk variants in the Greenlandic population. Diabetologia, 2018, 61, 2005-2015.	2.9	14
99	A likelihood method for estimating present-day human contamination in ancient male samples using low-depth X-chromosome data. Bioinformatics, 2020, 36, 828-841.	1.8	14
100	A Novel -192c/g Mutation in the Proximal P2 Promoter of the Hepatocyte Nuclear Factor-4Â Gene (HNF4A) Associates With Late-Onset Diabetes. Diabetes, 2006, 55, 1869-1873.	0.3	12
101	A Bayesian Multilocus Association Method: Allowing for Higher-Order Interaction in Association Studies. Genetics, 2007, 176, 1197-1208.	1.2	12
102	A referenceâ€free approach to analyse RADseq data using standard next generation sequencing toolkits. Molecular Ecology Resources, 2021, 21, 1085-1097.	2.2	12
103	NGSremix: a software tool for estimating pairwise relatedness between admixed individuals from next-generation sequencing data. G3: Genes, Genomes, Genetics, 2021, 11, .	0.8	12
104	Vicariance followed by secondary gene flow in a young gazelle species complex. Molecular Ecology, 2021, 30, 528-544.	2.0	11
105	Diabetes in Population Isolates: Lessons from Greenland. Review of Diabetic Studies, 2015, 12, 320-329.	0.5	11
106	Warthog Genomes Resolve an Evolutionary Conundrum and Reveal Introgression of Disease Resistance Genes. Molecular Biology and Evolution, 2022, 39, .	3.5	11
107	A ?243A?G polymorphism upstream of the gene encoding GAD65 associates with lower levels of body mass index and glycaemia in a population-based sample of 5857 middle-aged White subjects. Diabetic Medicine, 2007, 24, 702-706.	1.2	10
108	Identification of a novel BRCA1 nucleotide 4803delCC/c.4684delCC mutation and a nucleotide 249T>A/c.130T>A (p.Cys44Ser) mutation in two Greenlandic Inuit families: implications for genetic screening of Greenlandic Inuit families with high risk for breast and/or ovarian cancer. Breast Cancer Research and Treatment, 2010, 124, 259-264.	1.1	10

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109	Genetic study of the Arctic CPT1A variant suggests that its effect on fatty acid levels is modulated by traditional Inuit diet. European Journal of Human Genetics, 2020, 28, 1592-1601.	1.4	10
110	Predictors and trajectories of treatment response to SSRIs in patients suffering from PTSD. Psychiatry Research, 2021, 301, 113964.	1.7	10
111	Genome-Wide Association Study of Genetic Variants in LPS-Stimulated IL-6, IL-8, IL-10, IL-1ra and TNF-α Cytokine Response in a Danish Cohort. PLoS ONE, 2013, 8, e66262.	1.1	10
112	Large-scale study of the ?232C�>�G polymorphism of PCK1 in Type�2 diabetes. Diabetic Medicine, 2006, 21140-1144.	23, 1:2	9
113	A large-scale genome-wide gene expression analysis in peripheral blood identifies very few differentially expressed genes related to antidepressant treatment and response in patients with major depressive disorder. Neuropsychopharmacology, 2021, 46, 1324-1332.	2.8	9
114	The genetic history of Greenlandic-European contact. Current Biology, 2021, 31, 2214-2219.e4.	1.8	9
115	Loss of Sucrase-Isomaltase Function Increases Acetate Levels and Improves Metabolic Health in Greenlandic Cohorts. Gastroenterology, 2022, 162, 1171-1182.e3.	0.6	9
116	Partial USH2A deletions contribute to Usher syndrome in Denmark. European Journal of Human Genetics, 2015, 23, 1646-1651.	1.4	8
117	Identification of a novel locus for a USH3 like syndrome combined with congenital cataract. Clinical Genetics, 2010, 78, 388-397.	1.0	6
118	Genetic determinants of glycated hemoglobin levels in the Greenlandic Inuit population. European Journal of Human Genetics, 2018, 26, 868-875.	1.4	6
119	Omega-3 fatty acids and risk of cardiovascular disease in Inuit: First prospective cohort study. Atherosclerosis, 2020, 312, 28-34.	0.4	6
120	Estimating narrow-sense heritability using family data from admixed populations. Heredity, 2020, 124, 751-762.	1.2	6
121	Physical activity attenuates postprandial hyperglycaemia in homozygous TBC1D4 loss-of-function mutation carriers. Diabetologia, 2021, 64, 1795-1804.	2.9	6
122	Efficient approaches for large-scale GWAS with genotype uncertainty. G3: Genes, Genomes, Genetics, 2022, 12, .	0.8	5
123	The derived allele of a novel intergenic variant at chromosome 11 associates with lower body mass index and a favorable metabolic phenotype in Greenlanders. PLoS Genetics, 2020, 16, e1008544.	1.5	4
124	Detecting selection in low-coverage high-throughput sequencing data using principal component analysis. BMC Bioinformatics, 2021, 22, 470.	1.2	4
125	An LDLR missense variant poses high risk of familial hypercholesterolemia in 30% of Greenlanders and offers potential of early cardiovascular disease intervention. Human Genetics and Genomics Advances, 2022, 3, 100118.	1.0	4
126	Genetic architecture of obesity and related metabolic traits $\hat{a}\in$ " recent insights from isolated populations. Current Opinion in Genetics and Development, 2018, 50, 74-78.	1.5	3

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127	How robust are cross-population signatures of polygenic adaptation in humans?. , 0, 1, .		3
128	Low Pass Genomes of 141,431 Chinese Reveal Patterns of Viral Infection, Novel Phenotypic Associations, and the Genetic History of China. SSRN Electronic Journal, $0, , .$	0.4	2
129	246-OR: A Loss-of-Function Mutation in the Sucrase-Isomaltase Gene Is Linked to a Markedly Healthier Metabolic Profile in Greenlanders. Diabetes, 2020, 69, .	0.3	O