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List of Publications by Year
in descending order

34 papers	2,461 citations	471509 17 h-index	395702 33 g-index
36 all docs	36 docs citations	36 times ranked	5713 citing authors

#	Article	IF	Citations
1	A genome-wide association study identifies new psoriasis susceptibility loci and an interaction between HLA-C and ERAP1. Nature Genetics, 2010, 42, 985-990.	21.4	918
2	Genome-wide association study identifies five loci associated with lung function. Nature Genetics, 2010, 42, 36-44.	21.4	518
3	Genetic association study of QT interval highlights role for calcium signaling pathways in myocardial repolarization. Nature Genetics, 2014, 46, 826-836.	21.4	281
4	Effect of Five Genetic Variants Associated with Lung Function on the Risk of Chronic Obstructive Lung Disease, and Their Joint Effects on Lung Function. American Journal of Respiratory and Critical Care Medicine, 2011, 184, 786-795.	5.6	128
5	Genome-wide linkage analysis of Scandinavian affected sib-pairs supports presence of susceptibility loci for celiac disease on chromosomes 5 and 11. European Journal of Human Genetics, 2001, 9, 938-944.	2.8	80
6	Meta and pooled analysis of European coeliac disease data. European Journal of Human Genetics, 2003, 11, 828-834.	2.8	79
7	A Possible Mechanism behind Autoimmune Disorders Discovered By Genome-Wide Linkage and Association Analysis in Celiac Disease. PLoS ONE, 2013, 8, e70174.	2.5	51
8	Association study of IL2/IL21 and FcgRIIa: significant association with the IL2/IL21 region in Scandinavian coeliac disease families. Genes and Immunity, 2008, 9, 364-367.	4.1	35
9	Association analysis of GWAS and candidate gene loci in a Pakistani population with psoriasis. Molecular Immunology, 2015, 64, 190-194.	2.2	30
10	Characterisation of a Swedish cohort with orofacial granulomatosis with or without Crohn's disease. Oral Diseases, 2015, 21, e98-104.	3.0	28
11	A collaborative European search for non-DQA1*05-DQB1*02 celiac disease loci on HLA-DR3 haplotypes: analysis of transmission from homozygous parents. Human Immunology, 2003, 64, 350-358.	2.4	27
12	Single Nucleotide Polymorphisms in the FADS Gene Cluster but not the ELOVL2 Gene are Associated with Serum Polyunsaturated Fatty Acid Composition and Development of Allergy (in a Swedish Birth) Tj ETQq0 0 0 rgBT /Overlock 10 Tf	4.2	27
13	Validity of histology for the diagnosis of paediatric coeliac disease: a Swedish multicentre study. Scandinavian Journal of Gastroenterology, 2016, 51, 427-433.	1.5	25
14	Single nucleotide polymorphisms in theNOS2andNOS3genes are associated with exhaled nitric oxide. Journal of Medical Genetics, 2012, 49, 200-205.	3.2	24
15	A comprehensive screen for SNP associations on chromosome region 5q31â€“33 in Swedish/Norwegian celiac disease families. European Journal of Human Genetics, 2007, 15, 980-987.	2.8	20
16	Different <scp>DRB1</scp>*03:01â€<scp>DQB1</scp>*02:01 haplotypes confer different risk for celiac disease. Hla, 2017, 90, 95-101.	0.6	19
17	A family-based genome-wide association study of chronic rhinosinusitis with nasal polyps implicates several genes in the disease pathogenesis. PLoS ONE, 2017, 12, e0185244.	2.5	19
18	The IL12B gene does not confer susceptibility to coeliac disease. Tissue Antigens, 2002, 59, 70-72.	1.0	17

#	Article	IF	Citations
19	Altered peripheral amino acid profile indicate a systemic impact of active celiac disease and a possible role of amino acids in disease pathogenesis. PLoS ONE, 2018, 13, e0193764.	2.5	17
20	Association Between Genotypes and Phenotypes in Coeliac Disease. Journal of Pediatric Gastroenterology and Nutrition, 2009, 49, 165-169.	1.8	14
21	Shared Genetic Factors Involved in Celiac Disease, Type 2 Diabetes and Anorexia Nervosa Suggest Common Molecular Pathways for Chronic Diseases. PLoS ONE, 2016, 11, e0159593.	2.5	14
22	Genes involved in muscle contractility and nutrient signaling pathways within celiac disease risk loci show differential mRNA expression. BMC Medical Genetics, 2015, 16, 44.	2.1	13
23	Variation in the CTLA4/CD28 gene region confers an increased risk of coeliac disease. Annals of Human Genetics, 2002, 66, 125-37.	0.8	11
24	Meta-Analysis of Genome-Wide Linkage Studies in Celiac Disease. Human Heredity, 2009, 68, 223-230.	0.8	10
25	Heredity of nasal polyps. Rhinology, 2015, 53, 25-28.	1.3	10
26	Haplotypes of the inducible nitric oxide synthase gene are strongly associated with exhaled nitric oxide levels in adults: a population-based study. Journal of Medical Genetics, 2014, 51, 449-454.	3.2	9
27	Prediction and evaluation of the effect of pre-centrifugation sample management on the measurable untargeted LC-MS plasma metabolome. Analytica Chimica Acta, 2021, 1182, 338968.	5.4	9
28	Searching for genes influencing a complex disease: the case of coeliac disease. European Journal of Human Genetics, 2008, 16, 542-553.	2.8	7
29	The angiotensin-converting enzyme gene insertion polymorphism: a higher risk for psoriasis in male patients. British Journal of Dermatology, 2016, 175, 824-826.	1.5	6
30	A familyâ€based genomeâ€wide association study of recurrent aphthous stomatitis. Oral Diseases, 2020, 26, 1696-1705.	3.0	6
31	Role of proneurotensin as marker of paediatric coeliac disease. Clinical and Experimental Immunology, 2016, 186, 387-392.	2.6	5
32	Radiation of the urinary bladder attenuates the development of lipopolysaccharide-induced cystitis. International Immunopharmacology, 2020, 83, 106334.	3.8	3
33	The influence of heredity versus environment on coeliac disease. Gut, 2016, 65, 1779-1780.	12.1	1
34	Study Designs for Exploring the Non-HLA Genetics in Celiac Disease. Methods in Molecular Biology, 2015, 1326, 35-44.	0.9	0