David A Van Heel

List of Publications by Year
in descending order

86 papers	14,826 citations	28274 55 h-index	49909 87 g-index
96 all docs	96 docs citations	96 times ranked	21254 citing authors

#	Article	IF	Citations
1	Integrating polygenic risk scores in the prediction of type 2 diabetes risk and subtypes in British Pakistanis and Bangladeshis: A population-based cohort study. PLoS Medicine, 2022, 19, e1003981.	8.4	24
2	MC3R links nutritional state to childhood growth and the timing of puberty. Nature, 2021, 599, 436-441.	27.8	59
3	The power of genetic diversity in genome-wide association studies of lipids. Nature, 2021, 600, 675-679.	27.8	353
4	Fine-scale population structure and demographic history of British Pakistanis. Nature Communications, 2021, 12, 7189.	12.8	21
5	Cohort Profile: East London Genes & Health (ELGH), a community-based population genomics and health study in British Bangladeshi and British Pakistani people. International Journal of Epidemiology, 2020, 49, 20-21i.	1.9	71
6	A single nucleotide polymorphism genetic risk score to aid diagnosis of coeliac disease: a pilot study in clinical care. Alimentary Pharmacology and Therapeutics, 2020, 52, 1165-1173.	3.7	17
7	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
8	Evaluating drug targets through human loss-of-function genetic variation. Nature, 2020, 581, 459-464.	27.8	115
9	Characterising a healthy adult with a rare HAO1 knockout to support a therapeutic strategy for primary hyperoxaluria. ELife, 2020, 9, .	6.0	45
10	Associations of autozygosity with a broad range of human phenotypes. Nature Communications, 2019, 10, 4957.	12.8	84
11	Identification of rare sequence variation underlying heritable pulmonary arterial hypertension. Nature Communications, 2018, 9, 1416.	12.8	279
12	Formalising recall by genotype as an efficient approach to detailed phenotyping and causal inference. Nature Communications, 2018, 9, 711.	12.8	54
13	Exome Sequencing and Rare Variant Analysis RevealsÂMultiple Filaggrin Mutations in BangladeshiÂFamilies with Atopic Eczema andÂAdditional Risk Genes. Journal of Investigative Dermatology, 2018, 138, 2674-2677.	0.7	37
14	Association Between Telomere Length and Risk of Cancer and Non-Neoplastic Diseases. JAMA Oncology, 2017, 3, 636.	7.1	376
15	Estimating the human mutation rate from autozygous segments reveals population differences in human mutational processes. Nature Communications, 2017, 8, 303.	12.8	81
16	DNAJC21 Mutations Link a Cancer-Prone Bone Marrow Failure Syndrome to Corruption in 60S Ribosome Subunit Maturation. American Journal of Human Genetics, 2016, 99, 115-124.	6.2	85
17	Health and population effects of rare gene knockouts in adult humans with related parents. Science, 2016, 352, 474-477.	12.6	272
18	Investigating the Causal Relationship of C-Reactive Protein with 32 Complex Somatic and Psychiatric Outcomes: A Large-Scale Cross-Consortium Mendelian Randomization Study. PLoS Medicine, 2016, 13, e1001976.	8.4	150

#	Article	IF	Citations
19	Exome Sequencing of 75 Individuals from Multiply Affected Coeliac Families and Large Scale Resequencing Follow Up. PLoS ONE, 2015, 10, e0116845.	2.5	8
20	Fine mapping in the MHC region accounts for 18% additional genetic risk for celiac disease. Nature Genetics, 2015, 47, 577-578.	21.4	123
21	Widespread non-additive and interaction effects within HLA loci modulate the risk of autoimmune diseases. Nature Genetics, 2015, 47, 1085-1090.	21.4	164
22	Diagnosis and management of adult coeliac disease: guidelines from the British Society of Gastroenterology. Gut, 2014, 63, 1210-1228.	12.1	870
23	Improving coeliac disease risk prediction by testing non-HLA variants additional to HLA variants. Gut, 2014, 63, 415-422.	12.1	113
24	HLA-DQA1â€“HLA-DRB1 variants confer susceptibility to pancreatitis induced by thiopurine immunosuppressants. Nature Genetics, 2014, 46, 1131-1134.	21.4	165
25	Genetic insights into common pathways and complex relationships among immune-mediated diseases. Nature Reviews Genetics, 2013, 14, 661-673.	16.3	459
26	Negligible impact of rare autoimmune-locus coding-region variants on missing heritability. Nature, 2013, 498, 232-235.	27.8	184
27	Inflammatory Skin and Bowel Disease Linked toADAM17Deletion. New England Journal of Medicine, 2011, 365, 1502-1508.	27.0	285
28	Dense genotyping identifies and localizes multiple common and rare variant association signals in celiac disease. Nature Genetics, 2011, 43, 1193-1201.	21.4	682
29	Mutations in CSTA, Encoding Cystatin A, Underlie Exfoliative Ichthyosis and Reveal a Role for This Protease Inhibitor in Cell-Cell Adhesion. American Journal of Human Genetics, 2011, 89, 564-571.	6.2	89
30	Comparative methylomics reveals gene-body H3K36me3 in Drosophila predicts DNA methylation and CpG landscapes in other invertebrates. Genome Research, 2011, 21, 1841-1850.	5.5	57
31	Pervasive Sharing of Genetic Effects in Autoimmune Disease. PLoS Genetics, 2011, 7, e1002254.	3.5	540
32	Trans-eQTLs Reveal That Independent Genetic Variants Associated with a Complex Phenotype Converge on Intermediate Genes, with a Major Role for the HLA. PLoS Genetics, 2011, 7, e1002197.	3.5	324
33	A Meta-Analysis of Genome-Wide Association Scans Identifies IL18RAP, PTPN2, TAGAP, and PUS10 As Shared Risk Loci for Crohn's Disease and Celiac Disease. PLoS Genetics, 2011, 7, e1001283.	3.5	187
34	Evolutionary and Functional Analysis of Celiac Risk Loci Reveals SH2B3 as a Protective Factor against Bacterial Infection. American Journal of Human Genetics, 2010, 86, 970-977.	6.2	168
35	Multiple common variants for celiac disease influencing immune gene expression. Nature Genetics, 2010, 42, 295-302.	21.4	871
36	Comprehensive, Quantitative Mapping of T Cell Epitopes in Gluten in Celiac Disease. Science Translational Medicine, 2010, 2, 41ra51.	12.4	393

#	Article	IF	Citations
37	Common and different genetic background for rheumatoid arthritis and coeliac disease. Human Molecular Genetics, 2009, 18, 4195-4203.	2.9	128
38	Recent advances in coeliac disease genetics. Gut, 2009, 58, 473-476.	12.1	31
39	Association study of the IL18RAP locus in three European populations with coeliac disease. Human Molecular Genetics, 2009, 18, 1148-1155.	2.9	29
40	The genetics of chronic inflammatory diseases. Human Molecular Genetics, 2009, 18, R101-R106.	2.9	51
41	Replication of celiac disease UK genome-wide association study results in a US population. Human Molecular Genetics, 2009, 18, 4219-4225.	2.9	70
42	A genome-wide meta-analysis identifies 22 loci associated with eight hematological parameters in the HaemGen consortium. Nature Genetics, 2009, 41, 1182-1190.	21.4	481
43	Genetics and pathogenesis of coeliac disease. Seminars in Immunology, 2009, 21, 346-354.	5.6	49
44	Analysis of HLA and Non-HLA Alleles Can Identify Individuals at High Risk for Celiac Disease. Gastroenterology, 2009, 137, 834-840.e3.	1.3	126
45	Coeliac disease-associated risk variants in TNFAIP3 and REL implicate altered NF-ÂB signalling. Gut, 2009, 58, 1078-1083.	12.1	170
46	Detection, Imputation, and Association Analysis of Small Deletions and Null Alleles on Oligonucleotide Arrays. American Journal of Human Genetics, 2008, 82, 1316-1333.	6.2	40
47	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
48	Newly identified genetic risk variants for celiac disease related to the immune response. Nature Genetics, 2008, 40, 395-402.	21.4	599
49	New susceptibility genes for ulcerative colitis. Nature Genetics, 2008, 40, 686-688.	21.4	17
50	Translational Mini-Review Series on the Immunogenetics of Gut Disease: Immunogenetics of coeliac disease. Clinical and Experimental Immunology, 2008, 153, 162-173.	2.6	45
51	Shared and Distinct Genetic Variants in Type 1 Diabetes and Celiac Disease. New England Journal of Medicine, 2008, 359, 2767-2777.	27.0	654
52	Genetic Analysis of Innate Immunity in Crohn's Disease and Ulcerative Colitis Identifies Two Susceptibility Loci Harboring CARD9 and IL18RAP. American Journal of Human Genetics, 2008, 82, 1202-1210.	6.2	229
53	Effective Detection of Human Leukocyte Antigen Risk Alleles in Celiac Disease Using Tag Single Nucleotide Polymorphisms. PLoS ONE, 2008, 3, e2270.	2.5	136
54	Associations with tight junction genes PARD3 and MAGI2 in Dutch patients point to a common barrier defect for coeliac disease and ulcerative colitisAn unusual case of ascites. Gut, 2007, 57, 463-467.	12.1	142

#	Article	IF	Citations
55	Selective NOD1 Agonists Cause Shock and Organ Injury/DysfunctionIn Vivo. American Journal of Respiratory and Critical Care Medicine, 2007, 175, 595-603.	5.6	58
56	A Structural and Immunological Basis for the Role of Human Leukocyte Antigen DQ8 in Celiac Disease. Immunity, 2007, 27, 23-34.	14.3	157
57	Novel Association in Chromosome 4q27 Region with Rheumatoid Arthritis and Confirmation of Type 1 Diabetes Point to a General Risk Locus for Autoimmune Diseases. American Journal of Human Genetics, 2007, 81, 1284-1288.	6.2	189
58	Human Pancreatic Secretory Trypsin Inhibitor Stabilizes Intestinal Mucosa against Noxious Agents. American Journal of Pathology, 2007, 171, 1462-1473.	3.8	12
59	NOD2 activity modulates the phenotype of LPS-stimulated dendritic cells to promote the development of T-helper type 2-like lymphocytes â€" Possible implications for NOD2-associated Crohn's disease. Journal of Crohn's and Colitis, 2007, 1, 106-115.	1.3	17
60	A genome-wide association study for celiac disease identifies risk variants in the region harboring IL2 and IL21. Nature Genetics, 2007, 39, 827-829.	21.4	592
61	Recent advances in coeliac disease. Gut, 2006, 55, 1037-1046.	12.1	266
62	Detection of muramyl dipeptide-sensing pathway defects in patients with CrohnÊ¼s disease. Inflammatory Bowel Diseases, 2006, 12, 598-605.	1.9	21
63	Calcium Channel TRPV6 Expression in Human Duodenum: Different Relationships to the Vitamin D System and Aging in Men and Women. Journal of Bone and Mineral Research, 2006, 21, 1770-1777.	2.8	53
64	Modulation of dendritic cell phenotype and functionin an in vitro model of the intestinal epithelium. European Journal of Immunology, 2006, 36, 864-874.	2.9	71
65	Normal responses to specific NOD1-activating peptidoglycan agonists in the presence of the NOD2 frameshift and other mutations in Crohn's disease. European Journal of Immunology, 2006, 36, 1629-1635.	2.9	14
66	Interleukin 15: its role in intestinal inflammation. Gut, 2006, 55, 444-445.	12.1	37
67	Antagonists and non-toxic variants of the dominant wheat gliadin T cell epitope in coeliac disease. Gut, 2006, 55, 485-491.	12.1	56
68	Lack of association of MYO9B genetic variants with coeliac disease in a British cohort. Gut, 2006, 55, 969-972.	12.1	58
69	A common CTLA4 haplotype associated with coeliac disease. European Journal of Human Genetics, 2005, 13, 440-444.	2.8	76
70	Synergistic enhancement of Toll-like receptor responses by NOD1 activation. European Journal of Immunology, 2005, 35, 2471-2476.	2.9	135
71	Genetics in coeliac disease. Bailliere's Best Practice and Research in Clinical Gastroenterology, 2005, 19, 323-339.	2.4	103
72	Association between a complex insertion/deletion polymorphism in NOD1 (CARD4) and susceptibility to inflammatory bowel disease. Human Molecular Genetics, 2005, 14, 1245-1250.	2.9	299

#	Article	IF	Citations
73	T cells in peripheral blood after gluten challenge in coeliac disease. Gut, 2005, 54, 1217-1223.	12.1	110
74	Synergy between TLR9 and NOD2 innate immune responses is lost in genetic Crohn's disease. Gut, 2005, 54, 1553-1557.	12.1	111
75	Muramyl dipeptide and toll-like receptor sensitivity in NOD2-associated Crohn's disease. Lancet, The, 2005, 365, 1794-1796.	13.7	305
76	Inflammatory bowel disease susceptibility loci defined by genome scan meta-analysis of 1952 affected relative pairs. Human Molecular Genetics, 2004, 13, 763-770.	2.9	219
77	Further Evidence of IBD5/CARD15 (NOD2) Epistasis in the Susceptibility to Ulcerative Colitis. American Journal of Human Genetics, 2003, 73, 1465-1466.	6.2	59
78	Analysis of the IBD5 locus and potential gene-gene interactions in Crohn's disease. Gut, 2003, 52, 541-546.	12.1	96
79	Detecting the risks of osteoporotic fractures in coeliac disease. Gut, 2003, 52, 1229-a-1230.	12.1	5
80	The IBD6 Crohn's disease locus demonstrates complex interactions with CARD15 and IBD5 disease-associated variants. Human Molecular Genetics, 2003, 12, 2569-2575.	2.9	57
81	Association of TNF-alpha-857C with inflammatory bowel disease in the Australian population. Scandinavian Journal of Gastroenterology, 2003, 38, 533-4.	1.5	8
82	Inflammatory bowel disease is associated with a TNF polymorphism that affects an interaction between the OCT1 and NF-kappaB transcription factors. Human Molecular Genetics, 2002, 11, 1281-1289.	2.9	250
83	Crohn's disease: genetic susceptibility, bacteria, and innate immunity. Lancet, The, 2001, 357, 1902-1904.	13.7	66
84	Identification of novel polymorphisms in the Î²7 integrin gene: family-based association studies in inflammatory bowel disease. Genes and Immunity, 2001, 2, 455-460.	4.1	21
85	NOD2 (CARD15), the first susceptibility gene for Crohn's disease. Gut, 2001, 49, 752-754.	12.1	61
86	Colonoscopic appearances and diagnosis of intussusception due to large-bowel lipoma. Endoscopy, 1999, 31, 508.	1.8	3