Graham A Heap

List of Publications by Year
in descending order

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22 papers	5,902 citations	331259 21 h-index	642321 23 g-index
26 all docs	26 docs citations	26 times ranked	11096 citing authors

#	Article	IF	Citations
1	Genome-wide association study implicates immune activation of multiple integrin genes in inflammatory bowel disease. Nature Genetics, 2017, 49, 256-261.	9.4	943
2	Multiple common variants for celiac disease influencing immune gene expression. Nature Genetics, 2010, 42, 295-302.	9.4	871
3	Dense genotyping identifies and localizes multiple common and rare variant association signals in celiac disease. Nature Genetics, 2011, 43, 1193-1201.	9.4	682
4	Shared and Distinct Genetic Variants in Type 1 Diabetes and Celiac Disease. New England Journal of Medicine, 2008, 359, 2767-2777.	13.9	654
5	Newly identified genetic risk variants for celiac disease related to the immune response. Nature Genetics, 2008, 40, 395-402.	9.4	599
6	Predictors of anti-TNF treatment failure in anti-TNF-naive patients with active luminal Crohn's disease: a prospective, multicentre, cohort study. The Lancet Gastroenterology and Hepatology, 2019, 4, 341-353.	3.7	431
7	The pattern and outcome of acute severe colitis. Journal of Crohn's and Colitis, 2010, 4, 431-437.	0.6	276
8	HLA-DQA1*05 Carriage Associated With Development of Anti-Drug Antibodies to Infliximab and Adalimumab in Patients With Crohnâ€™s Disease. Gastroenterology, 2020, 158, 189-199.	0.6	249
9	Genome-wide analysis of allelic expression imbalance in human primary cells by high-throughput transcriptome resequencing. Human Molecular Genetics, 2010, 19, 122-134.	1.4	197
10	Coeliac disease-associated risk variants in TNFAIP3 and REL implicate altered NF-ÂB signalling. Gut, 2009, 58, 1078-1083.	6.1	170
11	HLA-DQA1â€“HLA-DRB1 variants confer susceptibility to pancreatitis induced by thiopurine immunosuppressants. Nature Genetics, 2014, 46, 1131-1134.	9.4	165
12	Association of Genetic Variants in NUDT15 With Thiopurine-Induced Myelosuppression in Patients With Inflammatory Bowel Disease. JAMA - Journal of the American Medical Association, 2019, 321, 773.	3.8	129
13	Complex nature of SNP genotype effects on gene expression in primary human leucocytes. BMC Medical Genomics, 2009, 2, 1.	0.7	86
14	Clinical Features and HLA Association of 5-Aminosalicylate (5-ASA)-induced Nephrotoxicity in Inflammatory Bowel Disease. Journal of Crohn's and Colitis, 2016, 10, 149-158.	0.6	85
15	Insights into the genetic epidemiology of Crohn's and rare diseases in the Ashkenazi Jewish population. PLoS Genetics, 2018, 14, e1007329.	1.5	66
16	Comparative methylomics reveals gene-body H3K36me3 in Drosophila predicts DNA methylation and CpG landscapes in other invertebrates. Genome Research, 2011, 21, 1841-1850.	2.4	57
17	The genetics of chronic inflammatory diseases. Human Molecular Genetics, 2009, 18, R101-R106.	1.4	51
18	A protein-truncating R179X variant in RNF186 confers protection against ulcerative colitis. Nature Communications, 2016, 7, 12342.	5.8	50

#	Article	IF	Citations
19	Genetics and pathogenesis of coeliac disease. Seminars in Immunology, 2009, 21, 346-354.	2.7	49
20	The immunogenicity of biosimilar infliximab: can we extrapolate the data across indications?. Expert Review of Gastroenterology and Hepatology, 2015, 9, 27-34.	1.4	42
21	Association study of the IL18RAP locus in three European populations with coeliac disease. Human Molecular Genetics, 2009, 18, 1148-1155.	1.4	29
22	Clinical Features and Genetic Risk of Demyelination Following Anti-TNF Treatment. Journal of Crohn's and Colitis, 2020, 14, 1653-1661.	0.6	9