Andrew J Cassidy

List of Publications by Year
in descending order

14	1,342	840585 11	1058333 14
papers	citations	h-index	g-index

14	14	14	2114
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	The sputum microbiome and clinical outcomes in patients with bronchiectasis: a prospective observational study. Lancet Respiratory Medicine,the, 2021, 9, 885-896.	5.2	63
2	Exome Sequencing Reveals Common and Rare Variants in F5 Associated With ACE Inhibitor and Angiotensin Receptor Blockerâ€“Induced Angioedema. Clinical Pharmacology and Therapeutics, 2020, 108, 1195-1202.	2.3	18
3	Neutrophil extracellular traps are associated with disease severity and microbiota diversity in patients with chronic obstructive pulmonary disease. Journal of Allergy and Clinical Immunology, 2018, 141, 117-127.	1.5	207
4	Genetic mannose binding lectin deficiency is associated with airway microbiota diversity and reduced exacerbation frequency in COPD. Thorax, 2018, 73, 510-518.	2.7	28
5	Novel <scp>TGM</scp>5 mutations in acral peeling skin syndrome. Experimental Dermatology, 2015, 24, 285-289.	1.4	11
6	Gauging NOTCH1 Activation in Cancer Using Immunohistochemistry. PLoS ONE, 2013, 8, e67306.	1.1	98
7	The Human Intermediate Filament Database: comprehensive information on a gene family involved in many human diseases. Human Mutation, 2008, 29, 351-360.	1.1	309
8	Prevalent and Rare Mutations in the Gene Encoding Filaggrin Cause Ichthyosis Vulgaris and Predispose Individuals to Atopic Dermatitis. Journal of Investigative Dermatology, 2006, 126, 1770-1775.	0.3	210
9	The Genetic Basis of Pachyonychia Congenita. Journal of Investigative Dermatology Symposium Proceedings, 2005, 10, 21-30.	0.8	98
10	Insights into Genotypeâ€“Phenotype Correlation in Pachyonychia Congenita from the Human Intermediate Filament Mutation Database. Journal of Investigative Dermatology Symposium Proceedings, 2005, 10, 31-36.	0.8	39
11	A Homozygous Missense Mutation in TGM5 Abolishes Epidermal Transglutaminase 5 Activity and Causes Acral Peeling Skin Syndrome. American Journal of Human Genetics, 2005, 77, 909-917.	2.6	122
12	Cloning and characterization of a novel human olfactory UDP-glucuronosyltransferase. Biochemical Journal, 1999, 340, 837-843.	1.7	95
13	Cloning and characterization of a novel human olfactory UDP-glucuronosyltransferase. Biochemical Journal, 1999, 340, 837.	1.7	37
14	6 Cloning of the human UGT1 gene complex in yeast artificial chromosomes: Novel aspects of gene structure and subchromosomal mapping to 2q37. Biochemical Society Transactions, 1997, 25, S562-S562.	1.6	7