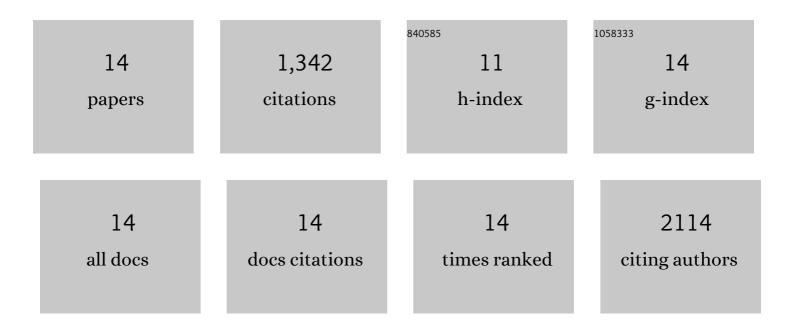
## Andrew J Cassidy

List of Publications by Year in descending order

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| #  | 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 <i>F5</i> Associated With ACE Inhibitor and Angiotensin Receptor Blocker–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<br>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<br>Proceedings, 2005, 10, 21-30.                                                                                         | 0.8 | 98        |
| 10 | Insights into Genotype–Phenotype Correlation in Pachyonychia Congenita from the Human<br>Intermediate Filament Mutation Database. Journal of Investigative Dermatology Symposium<br>Proceedings, 2005, 10, 31-36.    | 0.8 | 39        |
| 11 | A Homozygous Missense Mutation in TGM5 Abolishes Epidermal Transglutaminase 5 Activity and Causes<br>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<br>Journal, 1999, 340, 837-843.                                                                                     | 1.7 | 95        |
| 13 | Cloning and characterization of a novel human olfactory UDP-glucuronosyltransferase. Biochemical<br>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         |