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List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Laboratory evaluation of the IFN- \hat{l}^3 circuit for the molecular diagnosis of Mendelian susceptibility to mycobacterial disease. Critical Reviews in Clinical Laboratory Sciences, 2018, 55, 184-204.	6.1	43
2	Lethal Influenza in Two Related Adults with Inherited GATA2 Deficiency. Journal of Clinical Immunology, 2018, 38, 513-526.	3.8	29
3	Inhibition of endotoxin-induced airway epithelial cell injury by a novel family of pyrrol derivates. Laboratory Investigation, 2016, 96, 632-640.	3.7	13
4	Clinical Features of Candidiasis in Patients With Inherited Interleukin 12 Receptor \hat{l}^21 Deficiency. Clinical Infectious Diseases, 2014, 58, 204-213.	5.8	98
5	Anti-Inflammatory Activity of a Novel Family of Aryl Ureas Compounds in an Endotoxin-Induced Airway Epithelial Cell Injury Model. PLoS ONE, 2012, 7, e48468.	2.5	21
6	Variants at the promoter of the interleukin-6 gene are associated with severity and outcome of pneumococcal community-acquired pneumonia. Intensive Care Medicine, 2012, 38, 256-262.	8.2	61
7	Partial recessive IFN-Î ³ R1 deficiency: genetic, immunological and clinical features of 14 patients from 11 kindreds. Human Molecular Genetics, 2011, 20, 1509-1523.	2.9	102
8	Influence of genetic variability at the surfactant proteins A and D in community-acquired pneumonia: a prospective, observational, genetic study. Critical Care, 2011, 15, R57.	5.8	51
9	The Fcî³ receptor IIA-H/H131 genotype is associated with bacteremia in pneumococcal community-acquired pneumonia*. Critical Care Medicine, 2011, 39, 1388-1393.	0.9	52
10	Oesophageal squamous cell carcinoma in a young adult with IL-12RÂ1 deficiency. Journal of Medical Genetics, 2010, 47, 635-637.	3.2	25