Merry-Lynn Mcdonald

List of Publications by Year in descending order

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933447 1199594 13 836 10 12 citations g-index h-index papers 13 13 13 1875 docs citations times ranked citing authors all docs

#	Article	IF	CITATIONS
1	Longitudinal Association Between Muscle Loss and Mortality in Ever Smokers. Chest, 2022, 161, 960-970.	0.8	18
2	The Role of SNP Interactions when Determining Independence of Novel Signals in Genetic Association Studies $\hat{a} \in \text{``An Application to ARG1}$ and Bronchodilator Response. Journal of Personalized Medicine, 2021, 11, 145.	2.5	0
3	Diffuse Idiopathic Skeletal Hyperostosis in Smokers and Restrictive Spirometry Pattern: An Analysis of the COPDGene Cohort. Journal of Rheumatology, 2020, 47, 531-538.	2.0	6
4	Lungâ€Specific Risk Factors Associated With Incident Hip Fracture in Current and Former Smokers. Journal of Bone and Mineral Research, 2020, 35, 1952-1961.	2.8	6
5	Asthma Is a Risk Factor for Respiratory Exacerbations Without Increased Rate of Lung Function Decline. Chest, 2018, 153, 368-377.	0.8	14
6	Pectoralis muscle area and mortality in smokers without airflow obstruction. Respiratory Research, 2018, 19, 62.	3.6	41
7	Blood eosinophil count thresholds and exacerbations in patients with chronic obstructive pulmonary disease. Journal of Allergy and Clinical Immunology, 2018, 141, 2037-2047.e10.	2.9	138
8	A genome-wide association study identifies risk loci for spirometric measures among smokers of European and African ancestry. BMC Genetics, 2015 , 16 , 138 .	2.7	119
9	The clinical and genetic features of COPD-asthma overlap syndrome. European Respiratory Journal, 2014, 44, 341-350.	6.7	249
10	Chest CT Measures of Muscle and Adipose Tissue in COPD. Academic Radiology, 2014, 21, 1255-1261.	2.5	50
11	Splice-site mutations in the TRIC gene underlie autosomal recessive nonsyndromic hearing impairment in Pakistani families. Journal of Human Genetics, 2008, 53, 101-105.	2.3	45
12	Reduced endothelial secretion and plasma levels of transforming growth factor- \hat{l}^21 in patients with hereditary hemorrhagic telangiectasia type 1. Cardiovascular Research, 2005, 68, 155-164.	3.8	36
13	Potential Role of Modifier Genes Influencing Transforming Growth Factor- \hat{l}^21 Levels in the Development of Vascular Defects in Endoglin Heterozygous Mice with Hereditary Hemorrhagic Telangiectasia. American Journal of Pathology, 2001, 158, 2011-2020.	3.8	114