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List of Publications by Year in descending order

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Version: 2024-02-01

44
papers

2,778
citations

279798

23
h-index

243625

44
g-index

47
all docs

47
docs citations

47
times ranked

4657
citing authors

#	ARTICLE	IF	CITATIONS
1	Mortality in pulmonary arterial hypertension: prediction by the 2015 European pulmonary hypertension guidelines risk stratification model. <i>European Respiratory Journal</i> , 2017, 50, 1700740.	6.7	489
2	Genomic analyses inform on migration events during the peopling of Eurasia. <i>Nature</i> , 2016, 538, 238-242.	27.8	360
3	A recent bottleneck of Y chromosome diversity coincides with a global change in culture. <i>Genome Research</i> , 2015, 25, 459-466.	5.5	348
4	BMPR2 mutations and survival in pulmonary arterial hypertension: an individual participant data meta-analysis. <i>Lancet Respiratory Medicine</i> , 2016, 4, 129-137.	10.7	307
5	Pulmonary Arterial Hypertension: A Current Perspective on Established and Emerging Molecular Genetic Defects. <i>Human Mutation</i> , 2015, 36, 1113-1127.	2.5	185
6	Genome-Wide Analysis of Cold Adaptation in Indigenous Siberian Populations. <i>PLoS ONE</i> , 2014, 9, e98076.	2.5	128
7	ERS statement on exercise training and rehabilitation in patients with severe chronic pulmonary hypertension. <i>European Respiratory Journal</i> , 2019, 53, 1800332.	6.7	110
8	Selective sweep on human amylase genes postdates the split with Neanderthals. <i>Scientific Reports</i> , 2016, 6, 37198.	3.3	67
9	The Andean Adaptive Toolkit to Counteract High Altitude Maladaptation: Genome-Wide and Phenotypic Analysis of the Collas. <i>PLoS ONE</i> , 2014, 9, e93314.	2.5	55
10	Therapeutic potential of KLF2-induced exosomal microRNAs in pulmonary hypertension. <i>Nature Communications</i> , 2020, 11, 1185.	12.8	52
11	Standardized exercise training is feasible, safe, and effective in pulmonary arterial and chronic thromboembolic pulmonary hypertension: results from a large European multicentre randomized controlled trial. <i>European Heart Journal</i> , 2021, 42, 2284-2295.	2.2	51
12	Change of right heart size and function by long-term therapy with riociguat in patients with pulmonary arterial hypertension and chronic thromboembolic pulmonary hypertension. <i>International Journal of Cardiology</i> , 2015, 195, 19-26.	1.7	46
13	Right ventricular size and function under riociguat in pulmonary arterial hypertension and chronic thromboembolic pulmonary hypertension (the RIVER study). <i>Respiratory Research</i> , 2018, 19, 258.	3.6	39
14	First identification of <i>Krüppel-like factor 2</i> mutation in heritable pulmonary arterial hypertension. <i>Clinical Science</i> , 2017, 131, 689-698.	4.3	38
15	Early treatment with ambrisentan of mildly elevated mean pulmonary arterial pressure associated with systemic sclerosis: a randomized, controlled, double-blind, parallel group study (EDITA study). <i>Arthritis Research and Therapy</i> , 2019, 21, 217.	3.5	34
16	EIF2AK4 mutation as a second hit in hereditary pulmonary arterial hypertension. <i>Respiratory Research</i> , 2016, 17, 141.	3.6	33
17	Gender-related differences in pulmonary arterial hypertension targeted drugs administration. <i>Pharmacological Research</i> , 2016, 114, 103-109.	7.1	33
18	Positive selection of AS3MT to arsenic water in Andean populations. <i>Mutation Research - Fundamental and Molecular Mechanisms of Mutagenesis</i> , 2015, 780, 97-102.	1.0	32

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19	Evidence of Early-Stage Selection on EPAS1 and GPR126 Genes in Andean High Altitude Populations. <i>Scientific Reports</i> , 2017, 7, 13042.	3.3	29
20	Bayesian Inference Associates Rare <i>KDR</i> Variants With Specific Phenotypes in Pulmonary Arterial Hypertension. <i>Circulation Genomic and Precision Medicine</i> , 2021, 14, .	3.6	29
21	Pathobiology, pathology and genetics of pulmonary hypertension: Update from the Cologne Consensus Conference 2018. <i>International Journal of Cardiology</i> , 2018, 272, 4-10.	1.7	26
22	Mutation in BMPR2 Promoter: A "Second Hit"™ for Manifestation of Pulmonary Arterial Hypertension?. <i>PLoS ONE</i> , 2015, 10, e0133042.	2.5	26
23	Identification of genetic defects in pulmonary arterial hypertension by a new gene panel diagnostic tool. <i>Clinical Science</i> , 2016, 130, 2043-2052.	4.3	25
24	NADPH oxidase subunit NOXO1 is a target for emphysema treatment in COPD. <i>Nature Metabolism</i> , 2020, 2, 532-546.	11.9	23
25	Exercise Training and Rehabilitation in Pulmonary Hypertension. <i>Heart Failure Clinics</i> , 2018, 14, 425-430.	2.1	21
26	Genetic Predisposition to High-Altitude Pulmonary Edema. <i>High Altitude Medicine and Biology</i> , 2020, 21, 28-36.	0.9	21
27	Genetic and phenotypic differentiation of an Andean intermediate altitude population. <i>Physiological Reports</i> , 2015, 3, e12376.	1.7	18
28	Supervised Exercise Training in Patients with Chronic Thromboembolic Pulmonary Hypertension as Early Follow-Up Treatment after Pulmonary Endarterectomy: A Prospective Cohort Study. <i>Respiration</i> , 2020, 99, 577-588.	2.6	18
29	Gene panel diagnostics reveals new pathogenic variants in pulmonary arterial hypertension. <i>Respiratory Research</i> , 2022, 23, 74.	3.6	18
30	Genetics of pulmonary hypertension and high-altitude pulmonary edema. <i>Journal of Applied Physiology</i> , 2020, 128, 1432-1438.	2.5	15
31	Epigenetic reactivation of transcriptional programs orchestrating fetal lung development in human pulmonary hypertension. <i>Science Translational Medicine</i> , 2022, 14, .	12.4	15
32	Circulating MicroRNA Markers for Pulmonary Hypertension in Supervised Exercise Intervention and Nightly Oxygen Intervention. <i>Frontiers in Physiology</i> , 2018, 9, 955.	2.8	14
33	Risk stratification and prognostic factors in patients with pulmonary arterial hypertension and comorbidities a cross-sectional cohort study with survival follow-up. <i>Respiratory Research</i> , 2020, 21, 127.	3.6	14
34	Myeloproliferative Diseases as Possible Risk Factor for Development of Chronic Thromboembolic Pulmonary Hypertension—A Genetic Study. <i>International Journal of Molecular Sciences</i> , 2020, 21, 3339.	4.1	13
35	Mutually reinforcing effects of genetic variants and interferon- β 1a therapy for pulmonary arterial hypertension development in multiple sclerosis patients. <i>Pulmonary Circulation</i> , 2019, 9, 1-6.	1.7	9
36	Analysis of full-length mitochondrial DNA D-loop sequences from <i>Macaca fascicularis</i> of different geographical origin reveals novel haplotypes. <i>Journal of Medical Primatology</i> , 2015, 44, 125-136.	0.6	6

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37	Right Heart Size and Right Ventricular Reserve in Pulmonary Hypertension: Impact on Management and Prognosis. <i>Diagnostics</i> , 2020, 10, 1110.	2.6	6
38	Prognostic impact of hypochromic erythrocytes in patients with pulmonary arterial hypertension. <i>Respiratory Research</i> , 2021, 22, 288.	3.6	6
39	The role of rehabilitation in patients with pulmonary arterial hypertension. <i>Current Opinion in Pulmonary Medicine</i> , 2019, 25, 398-404.	2.6	5
40	BMPR2 Promoter Variants Effect Gene Expression in Pulmonary Arterial Hypertension Patients. <i>Genes</i> , 2020, 11, 1168.	2.4	3
41	The effect of exercise training and physiotherapy on left and right heart function in heart failure with preserved ejection fraction: a systematic literature review. <i>Heart Failure Reviews</i> , 2023, 28, 193-206.	3.9	3
42	Multicentre trials on specialised exercise training and rehabilitation are useful in patients with pulmonary hypertension. <i>European Respiratory Journal</i> , 2019, 54, 1901631.	6.7	2
43	Reduction of BMPR2 mRNA Expression in Peripheral Blood of Pulmonary Arterial Hypertension Patients: A Marker for Disease Severity?. <i>Genes</i> , 2022, 13, 759.	2.4	2
44	The Experience, Prerequisites, and the Barriers in Organizing a Specialized Rehabilitation Program for Patients with Pulmonary Hypertension. <i>Respiration</i> , 2021, 100, 1-9.	2.6	0