

Marco Mura

List of Publications by Year in descending order

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Version: 2024-02-01

54
papers

2,101
citations

361045

20
h-index

233125

45
g-index

57
all docs

57
docs citations

57
times ranked

2897
citing authors

#	ARTICLE	IF	CITATIONS
1	Bile acid aspiration and the development of bronchiolitis obliterans after lung transplantation. <i>Journal of Thoracic and Cardiovascular Surgery</i> , 2005, 129, 1144-1152.	0.4	290
2	Acute lung injury and cell death: how many ways can cells die?. <i>American Journal of Physiology - Lung Cellular and Molecular Physiology</i> , 2008, 294, L632-L641.	1.3	197
3	Predicting survival in newly diagnosed idiopathic pulmonary fibrosis: a 3-year prospective study. <i>European Respiratory Journal</i> , 2012, 40, 101-109.	3.1	179
4	TNF α -Induced Long Pentraxin PTX3 Expression in Human Lung Epithelial Cells via JNK. <i>Journal of Immunology</i> , 2005, 175, 8303-8311.	0.4	166
5	Vascular endothelial growth factor and related molecules in acute lung injury. <i>Journal of Applied Physiology</i> , 2004, 97, 1605-1617.	1.2	165
6	Metabolomic Heterogeneity of Pulmonary Arterial Hypertension. <i>PLoS ONE</i> , 2014, 9, e88727.	1.1	111
7	Osteopontin lung gene expression is a marker of disease severity in pulmonary arterial hypertension. <i>Respirology</i> , 2019, 24, 1104-1110.	1.3	76
8	The presence of emphysema further impairs physiologic function in patients with idiopathic pulmonary fibrosis. <i>Respiratory Care</i> , 2006, 51, 257-65.	0.8	70
9	ANGIOTENSIN-CONVERTING ENZYME INHIBITOR CAPTOPRIL PREVENTS OLEIC ACID-INDUCED SEVERE ACUTE LUNG INJURY IN RATS. <i>Shock</i> , 2007, 28, 106-111.	1.0	66
10	Comprehensive gene expression profiling identifies distinct and overlapping transcriptional profiles in non-specific interstitial pneumonia and idiopathic pulmonary fibrosis. <i>Respiratory Research</i> , 2018, 19, 153.	1.4	66
11	The early responses of VEGF and its receptors during acute lung injury: implication of VEGF in alveolar epithelial cell survival. <i>Critical Care</i> , 2006, 10, R130.	2.5	65
12	HRCT and histopathological evaluation of fibrosis and tissue destruction in IPF associated with pulmonary emphysema. <i>Respiratory Medicine</i> , 2008, 102, 1753-1761.	1.3	54
13	Gene Expression Profiling in the Lungs of Patients With Pulmonary Hypertension Associated With Pulmonary Fibrosis. <i>Chest</i> , 2012, 141, 661-673.	0.4	49
14	Src tyrosine kinase inhibition prevents pulmonary ischemia-reperfusion-induced acute lung injury. <i>Intensive Care Medicine</i> , 2012, 38, 894-905.	3.9	49
15	Functional Predictors of Exertional Dyspnea, 6-min Walking Distance and HRCT Fibrosis Score in Idiopathic Pulmonary Fibrosis. <i>Respiration</i> , 2006, 73, 495-502.	1.2	44
16	High-volume ventilation induces pentraxin 3 expression in multiple acute lung injury models in rats. <i>American Journal of Physiology - Lung Cellular and Molecular Physiology</i> , 2007, 292, L144-L153.	1.3	42
17	Functions of Type II Pneumocyte-Derived Vascular Endothelial Growth Factor in Alveolar Structure, Acute Inflammation, and Vascular Permeability. <i>American Journal of Pathology</i> , 2010, 176, 1725-1734.	1.9	42
18	PTX3 as a potential biomarker of acute lung injury: supporting evidence from animal experimentation. <i>Intensive Care Medicine</i> , 2010, 36, 356-364.	3.9	40

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19	A simple dyspnoea scale as part of the assessment to predict outcome across chronic interstitial lung disease. <i>Respirology</i> , 2017, 22, 501-507.	1.3	35
20	Review: New perspectives in the treatment of idiopathic pulmonary fibrosis. <i>Therapeutic Advances in Respiratory Disease</i> , 2008, 2, 75-93.	1.0	25
21	Anti-Human Tissue Factor Antibody Ameliorated Intestinal Ischemia Reperfusion-Induced Acute Lung Injury in Human Tissue Factor Knock-In Mice. <i>PLoS ONE</i> , 2008, 3, e1527.	1.1	21
22	Inflammatory activity is still present in the advanced stages of idiopathic pulmonary fibrosis. <i>Respirology</i> , 2005, 10, 609-614.	1.3	19
23	Metabolic syndrome and risk of pulmonary involvement. <i>Respiratory Medicine</i> , 2010, 104, 47-51.	1.3	18
24	Bone Marrow Cells Expressing Clara Cell Secretory Protein Increase Epithelial Repair After Ablation of Pulmonary Clara Cells. <i>Molecular Therapy</i> , 2013, 21, 1251-1258.	3.7	15
25	Depletion of Bone Marrow CCSP-Expressing Cells Delays Airway Regeneration. <i>Molecular Therapy</i> , 2015, 23, 561-569.	3.7	15
26	Longitudinal functional changes with clinically significant radiographic progression in idiopathic pulmonary fibrosis: are we following the right parameters?. <i>Respiratory Research</i> , 2020, 21, 119.	1.4	15
27	Multi-dimensional scores to predict mortality in patients with idiopathic pulmonary fibrosis undergoing lung transplantation assessment. <i>Respiratory Medicine</i> , 2017, 125, 65-71.	1.3	13
28	Bullous emphysema versus diffuse emphysema: a functional and radiologic comparison. <i>Respiratory Medicine</i> , 2005, 99, 171-178.	1.3	11
29	INTESTINAL ISCHEMIA-REPERFUSION-INDUCED ACUTE LUNG INJURY AND ONCOTIC CELL DEATH IN MULTIPLE ORGANS. <i>Shock</i> , 2007, 28, 227-238.	1.0	9
30	Early Decline in Six-Minute Walk Distance from the Time of Diagnosis Predicts Clinical Worsening in Pulmonary Arterial Hypertension. <i>Respiration</i> , 2015, 89, 365-373.	1.2	9
31	Multi-dimensional Assessment of IPF Across a Wide Range of Disease Severity. <i>Lung</i> , 2018, 196, 707-713.	1.4	9
32	Multi-dimensional indices to stage idiopathic pulmonary fibrosis: a systematic review. <i>Sarcoidosis Vasculitis and Diffuse Lung Diseases</i> , 2014, 31, 8-18.	0.2	9
33	Assessing the Therapeutic Response to Pirfenidone in Idiopathic Pulmonary Fibrosis: Can We Do Better than with Forced Vital Capacity Alone?. <i>Lung</i> , 2017, 195, 101-105.	1.4	8
34	Cluster analysis of transcriptomic datasets to identify endotypes of idiopathic pulmonary fibrosis. <i>Thorax</i> , 2023, 78, 551-558.	2.7	8
35	Does Technetium-99m Diethylenetriaminepentaacetate Clearance Predict the Clinical Course of Idiopathic Pulmonary Fibrosis?. <i>Canadian Respiratory Journal</i> , 2004, 11, 477-479.	0.8	7
36	Successful Treatment of Fibrosing Organising Pneumonia Causing Respiratory Failure with Mycophenolic Acid. <i>Respiration</i> , 2016, 92, 279-282.	1.2	7

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37	Pulmonary Artery Abnormalities in Ex-smokers with and without Airflow Obstruction. COPD: Journal of Chronic Obstructive Pulmonary Disease, 2016, 13, 224-234.	0.7	7
38	Associated Pulmonary Hypertension Is an Independent Contributor to Exercise Intolerance in Chronic Fibrosing Interstitial Pneumonias. Respiration, 2018, 96, 543-551.	1.2	7
39	Implementing an interstitial lung disease clinic improves survival without increasing health care resource utilization. Pulmonary Pharmacology and Therapeutics, 2019, 56, 94-99.	1.1	7
40	The CALIPER-Revised Version of the Composite Physiologic Index is a Better Predictor of Survival in IPF than the Original Version. Lung, 2020, 198, 169-172.	1.4	7
41	Fat-Free Mass Index Controlled for Age and Sex and Malnutrition Are Predictors of Survival in Interstitial Lung Disease. Respiration, 2021, 100, 379-386.	1.2	7
42	Use of nintedanib in interstitial lung disease other than idiopathic pulmonary fibrosis: much caution is warranted. Pulmonary Pharmacology and Therapeutics, 2021, 66, 101987.	1.1	6
43	A 53-year-old man with dyspnoea, respiratory failure, consistent with infliximab-induced acute interstitial pneumonitis after an accelerated induction dosing schedule. BMJ Case Reports, 2017, 2017, bcr-2017-219956.	0.2	6
44	Interstitial Lung Disease, Body Mass Index, Energy Expenditure and Malnutrition—a Review. Current Pulmonology Reports, 2017, 6, 70-74.	0.5	5
45	Tobacco Worker’s Lung: A Neglected Subtype of Hypersensitivity Pneumonitis. Lung, 2021, 199, 13-19.	1.4	5
46	Nutrition implications of intrinsic restrictive lung disease. Nutrition in Clinical Practice, 2022, 37, 239-255.	1.1	5
47	Exercise capacity and its relationship with body composition and nutrition status in patients with interstitial lung disease. Nutrition in Clinical Practice, 2021, 36, 891-898.	1.1	4
48	Digital quantification of p16-positive foci in fibrotic interstitial lung disease is associated with a phenotype of idiopathic pulmonary fibrosis with reduced survival. Respiratory Research, 2022, 23, .	1.4	3
49	Bronchiolitis obliterans syndrome as manifestation of lung GVHD: Not the only one. Respirology, 2019, 24, 702-702.	1.3	2
50	Invasive pulmonary aspergillosis in an immunocompetent, heavy smoker of marijuana with emphysema and chronic obstructive pulmonary disease. Canadian Journal of Respiratory, Critical Care, and Sleep Medicine, 2020, , 1-4.	0.2	2
51	TPMT and HLA-DQA1-HLA-DRB genetic profiling to guide the use of azathioprine in the treatment of interstitial lung disease: First experience. Pulmonary Pharmacology and Therapeutics, 2021, 66, 101988.	1.1	2
52	6-minute walk distance as a predictor of outcome in idiopathic pulmonary fibrosis. European Respiratory Journal, 2014, 43, 1822-1823.	3.1	1
53	Exercise Testing in Idiopathic Pulmonary Fibrosis: Expanding Our Options. Respiration, 2021, 100, 568-570.	1.2	1
54	Validation of the risk stratification score in idiopathic pulmonary fibrosis: study protocol of a prospective, multi-centre, observational, 3-year clinical trial. BMC Pulmonary Medicine, 2021, 21, 396.	0.8	0