Eric S White

List of Publications by Year
in descending order

161 papers	15,284 citations	13865 67 h-index	18647 119 g-index
166 all docs	166 docs citations	166 times ranked	17784 citing authors

#	Article	IF	Citations
1	Detection and Early Referral of Patients With Interstitial Lung Abnormalities. Chest, 2022, 161, 470-482.	0.8	26
2	Human antigen R promotes lung fibroblast differentiation to myofibroblasts and increases extracellular matrix production. Journal of Cellular Physiology, 2021, 236, 6836-6851.	4.1	17
3	Adipose Stromal Cell-Secretome Counteracts Profibrotic Signals From IPF Lung Matrices. Frontiers in Pharmacology, 2021, 12, 669037.	3.5	8
4	Adoption of Antifibrotic Medications: A Closer Look at the Data. Annals of the American Thoracic Society, 2021, 18, 1756-1757.	3.2	0
5	Interpretation of PFTs and Decline in PFTs. In Clinical Practice, 2021, , 139-149.	0.0	0
6	Type I Collagen Signaling Regulates Opposing Fibrotic Pathways through Î±₂Î²₁ Integrin. American Journal of Respiratory Cell and Molecular Biology, 2020, 63, 613-622.	2.9	12
7	Identification of a unique temporal signature in blood and BAL associated with IPF progression. Scientific Reports, 2020, 10, 12049.	3.3	10
8	Development and Initial Validation Analyses of the Living with Idiopathic Pulmonary Fibrosis Questionnaire. American Journal of Respiratory and Critical Care Medicine, 2020, 202, 1689-1697.	5.6	27
9	Changes in biomarkers with nintedanib and sildenafil in subjects with IPF in the INSTAGE trial: subgroup analysis by right heart dysfunction (RHD). , 2020, , .		0
10	Effects of nintedanib on markers of epithelial damage in subjects with IPF: data from the INMARK trial. , 2020, , .		1
11	Changes in biomarkers with nintedanib plus sildenafil in subjects with IPF by presence of emphysema in the INSTAGE trial. , 2020, , .		0
12	Radiographic Honeycombing and Altered Lung Microbiota in Patients with Idiopathic Pulmonary Fibrosis. American Journal of Respiratory and Critical Care Medicine, 2019, 200, 1544-1547.	5.6	20
13	Biomarkers of extracellular matrix turnover in patients with idiopathic pulmonary fibrosis given nintedanib (INMARK study): a randomised, placebo-controlled study. Lancet Respiratory Medicine,the, 2019, 7, 771-779.	10.7	65
14	Frailty and geriatric conditions in older patients with idiopathic pulmonary fibrosis. Respiratory Medicine, 2019, 148, 6-12.	2.9	31
15	The Fibrosis Across Organs Symposium: A Roadmap for Future Research Priorities. American Journal of the Medical Sciences, 2019, 357, 405-410.	1.1	1
16	Routine Chest Radiography for the Evaluation of Pneumothorax Following Bronchoscopy. Academic Radiology, 2019, 26, 585-590.	2.5	5
17	Lung Microbiota Contribute to Pulmonary Inflammation and Disease Progression in Pulmonary Fibrosis. American Journal of Respiratory and Critical Care Medicine, 2019, 199, 1127-1138.	5.6	205
18	Prevalence, Treatment, and Outcomes of Coexistent Pulmonary Hypertension and Interstitial Lung Disease in Systemic Sclerosis. Arthritis and Rheumatology, 2019, 71, 1339-1349.	5.6	54

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19	Hypersensitivity Pneumonitis. Chest, 2019, 155, 699-711.	0.8	148
20	Blood biomarkers predicting disease progression in patients with IPF: data from the INMARK trial. , 2019, , .		3
21	Changes in biomarkers in patients with idiopathic pulmonary fibrosis (IPF) treated with nintedanib and sildenafil. , 2019, , .		3
22	Effect of nintedanib on blood biomarkers in patients with IPF in the INMARK trial. , 2019, , .		3
23	Interstitial pneumonia with autoimmune features (IPAF): a clinical entity?. , 2019, , .		0
24	Impact of comorbidities in interstitial pneumonia with autoimmune features (IPAF). , 2019, , .		0
25	Reliability, construct validity and responsiveness to change of the PROMIS-29 in systemic sclerosis-associated interstitial lung disease. Clinical and Experimental Rheumatology, 2019, 37 Suppl 119, 49-56.	0.8	1
26	Discoidin Domain Receptor 2 Signaling Regulates Fibroblast Apoptosis through PDK1/Akt. American Journal of Respiratory Cell and Molecular Biology, 2018, 59, 295-305.	2.9	35
27	Loss of CDKN2B Promotes Fibrosis via Increased Fibroblast Differentiation Rather Than Proliferation. American Journal of Respiratory Cell and Molecular Biology, 2018, 59, 200-214.	2.9	15
28	The ED-A domain enhances the capacity of fibronectin to store latent TGF-Î² binding protein-1 in the fibroblast matrix. Journal of Cell Science, 2018, 131, .	2.0	107
29	Increased circulating desmosine and age-dependent elastinolysis in idiopathic pulmonary fibrosis. Respiratory Research, 2018, 19, 45.	3.6	12
30	Extracellular matrix in lung development, homeostasis and disease. Matrix Biology, 2018, 73, 77-104.	3.6	200
31	Bioengineered lungs generated from human i <scp>PSC</scp> sâ€derived epithelial cells on native extracellular matrix. Journal of Tissue Engineering and Regenerative Medicine, 2018, 12, e1623-e1635.	2.7	35
32	InÂVitro Induction and InÂVivo Engraftment of Lung Bud Tip Progenitor Cells Derived from Human Pluripotent Stem Cells. Stem Cell Reports, 2018, 10, 101-119.	4.8	192
33	Investigating the effects of nintedanib on biomarkers of extracellular matrix turnover in patients with IPF: design of the randomised placebo-controlled INMARKÂ®trial. BMJ Open Respiratory Research, 2018, 5, e000325.	3.0	14
34	Optimising experimental research in respiratory diseases: an ERS statement. European Respiratory Journal, 2018, 51, 1702133.	6.7	98
35	Development and validation of a radiological diagnosis model for hypersensitivity pneumonitis. European Respiratory Journal, 2018, 52, 1800443.	6.7	55
36	Wilmsâ€™ tumor 1 drives fibroproliferation and myofibroblast transformation in severe fibrotic lung disease. JCI Insight, 2018, 3, .	5.0	32

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37	Î±vÎ²3 Integrin drives fibroblast contraction and strain stiffening of soft provisional matrix during progressive fibrosis. JCI Insight, 2018, 3, .	5.0	78
38	FOXM1 is a critical driver of lung fibroblast activation and fibrogenesis. Journal of Clinical Investigation, 2018, 128, 2389-2405.	8.2	88
39	Investigating effects of nintedanib on biomarkers of ECM turnover in patients with IPF: the INMARK study. , 2018, , .		0
40	Microbes Are Associated with Host Innate Immune Response in Idiopathic Pulmonary Fibrosis. American Journal of Respiratory and Critical Care Medicine, 2017, 196, 208-219.	5.6	130
41	An Official American Thoracic Society Workshop Report: Use of Animal Models for the Preclinical Assessment of Potential Therapies for Pulmonary Fibrosis. American Journal of Respiratory Cell and Molecular Biology, 2017, 56, 667-679.	2.9	267
42	The peripheral blood proteome signature of idiopathic pulmonary fibrosis is distinct from normal and is associated with novel immunological processes. Scientific Reports, 2017, 7, 46560.	3.3	51
43	MMP-7 is a predictive biomarker of disease progression in patients with idiopathic pulmonary fibrosis. ERJ Open Research, 2017, 3, 00074-2016.	2.6	86
44	The diagnosis of idiopathic pulmonary fibrosis: current and future approaches. Lancet Respiratory Medicine,the, 2017, 5, 61-71.	10.7	79
45	Joint SNMMIâ€“ASNC expert consensus document on the role of 18F-FDG PET/CT in cardiac sarcoid detection and therapy monitoring. Journal of Nuclear Cardiology, 2017, 24, 1741-1758.	2.1	132
46	Joint SNMMIâ€“ASNC Expert Consensus Document on the Role of ¹⁸F-FDG PET/CT in Cardiac Sarcoid Detection and Therapy Monitoring. Journal of Nuclear Medicine, 2017, 58, 1341-1353.	5.0	187
47	Lysyl oxidases regulate fibrillar collagen remodelling in idiopathic pulmonary fibrosis. DMM Disease Models and Mechanisms, 2017, 10, 1301-1312.	2.4	110
48	The instructive extracellular matrix of the lung: basic composition and alterations in chronic lung disease. European Respiratory Journal, 2017, 50, 1601805.	6.7	341
49	Utility of Transbronchial vsÂSurgical Lung Biopsy in the Diagnosis of Suspected Fibrotic Interstitial Lung Disease. Chest, 2017, 151, 389-399.	0.8	87
50	Hsp90 regulation of fibroblast activation in pulmonary fibrosis. JCI Insight, 2017, 2, e91454.	5.0	73
51	A bioengineered niche promotes in vivo engraftment and maturation of pluripotent stem cell derived human lung organoids. ELife, 2016, 5, .	6.0	162
52	Commentary: A Breath of Fresh Air on the Mesenchyme: Impact of Impaired Mesenchymal Development on the Pathogenesis of Bronchopulmonary Dysplasia. Frontiers in Medicine, 2016, 3, 13.	2.6	0
53	Netrinâ€1 Regulates Fibrocyte Accumulation in the Decellularized Fibrotic Sclerodermatous Lung Microenvironment and in Bleomycinâ€Induced Pulmonary Fibrosis. Arthritis and Rheumatology, 2016, 68, 1251-1261.	5.6	51
54	An American Thoracic Society Official Research Statement: Future Directions in Lung Fibrosis Research. American Journal of Respiratory and Critical Care Medicine, 2016, 193, 792-800.	5.6	22

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55	Plasma Surfactant Protein-D, Matrix Metalloproteinase-7, and Osteopontin Index Distinguishes Idiopathic Pulmonary Fibrosis from Other Idiopathic Interstitial Pneumonias. American Journal of Respiratory and Critical Care Medicine, 2016, 194, 1242-1251.	5.6	131
56	Comparative biology of decellularized lung matrix: Implications of species mismatch in regenerative medicine. Biomaterials, 2016, 102, 220-230.	11.4	68
57	Animal and cellular models of human disease. Journal of Pathology, 2016, 238, 137-140.	4.5	9
58	TGF-Î² Induced Collagen Remodelling by IPF Fibroblasts is Alleviated by Inhibition of Lysyl Oxidase Enzyme Activity. QJM - Monthly Journal of the Association of Physicians, 2016, , .	0.5	0
59	Recommendations for minimum information for publication ofÂexperimental pathology data: <scp>MINPEPA</scp> guidelines. Journal of Pathology, 2016, 238, 359-367.	4.5	31
60	IL-36Î³ is secreted in microparticles and exosomes by lung macrophages in response to bacteria and bacterial components. Journal of Leukocyte Biology, 2016, 100, 413-421.	3.3	47
61	Mucking around in the Genome: MUC5B in Idiopathic Pulmonary Fibrosis. American Journal of Respiratory and Critical Care Medicine, 2016, 193, 355-357.	5.6	4
62	Senescent Cells Contribute to the Physiological Remodeling of Aged Lungs. Journals of Gerontology - Series A Biological Sciences and Medical Sciences, 2016, 71, 153-160.	3.6	65
63	Microbes mediated host innate immune response in idiopathic pulmonary fibrosis. , 2016, , .		1
64	Six-SOMAmer Index Relating to Immune, Protease and Angiogenic Functions Predicts Progression in IPF. PLoS ONE, 2016, 11, e0159878.	2.5	43
65	Plakoglobin expression in fibroblasts and its role in idiopathic pulmonary fibrosis. BMC Pulmonary Medicine, 2015, 15, 140.	2.0	3
66	In vitro generation of human pluripotent stem cell derived lung organoids. ELife, 2015, 4, .	6.0	605
67	Comparative Biology of the Normal LungÂExtracellular Matrix. , 2015, , 387-402.		2
68	Blue Journal Conference. Aging and Susceptibility to Lung Disease. American Journal of Respiratory and Critical Care Medicine, 2015, 191, 261-269.	5.6	149
69	Extensive Phenotyping of Individuals at Risk for Familial Interstitial Pneumonia Reveals Clues to the Pathogenesis of Interstitial Lung Disease. American Journal of Respiratory and Critical Care Medicine, 2015, 191, 417-426.	5.6	141
70	Lung Extracellular Matrix and Fibroblast Function. Annals of the American Thoracic Society, 2015, 12, S30-S33.	3.2	145
71	Inhibition of Myocardin-Related Transcription Factor/Serum Response Factor Signaling Decreases Lung Fibrosis and Promotes Mesenchymal Cell Apoptosis. American Journal of Pathology, 2015, 185, 969-986.	3.8	138
72	Generation of tissue-engineered small intestine using embryonic stem cell-derived human intestinal organoids. Biology Open, 2015, 4, 1462-1472.	1.2	151

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73	Conformational coupling of integrin and Thy-1 regulates Fyn priming and fibroblast mechanotransduction. Journal of Cell Biology, 2015, 211, 173-190.	5.2	78
74	Fibrocytes Regulate Wilms Tumor 1â€“Positive Cell Accumulation in Severe Fibrotic Lung Disease. Journal of Immunology, 2015, 195, 3978-3991.	0.8	29
75	TNF-Receptor Inhibitor Therapy for the Treatment of Children with Idiopathic Pneumonia Syndrome. A Joint Pediatric Blood and Marrow Transplant Consortium and Children's Oncology Group Study (ASCT0521). Biology of Blood and Marrow Transplantation, 2015, 21, 67-73.	2.0	62
76	Conformational coupling of integrin and Thy-1 regulates Fyn priming and fibroblast mechanotransduction. Journal of General Physiology, 2015, 146, 1465OIA57.	1.9	0
77	Up-Regulation of Heparan Sulfate 6-O-Sulfation in Idiopathic Pulmonary Fibrosis. American Journal of Respiratory Cell and Molecular Biology, 2014, 50, 106-114.	2.9	30
78	Lung Fibroblasts from Patients with Idiopathic Pulmonary Fibrosis Exhibit Genome-Wide Differences in DNA Methylation Compared to Fibroblasts from Nonfibrotic Lung. PLoS ONE, 2014, 9, e107055.	2.5	70
79	Chitinase 3â€“Like 1 Suppresses Injury and Promotes Fibroproliferative Responses in Mammalian Lung Fibrosis. Science Translational Medicine, 2014, 6, 240ra76.	12.4	162
80	Open-Access Biorepository for Idiopathic Pulmonary Fibrosis. The Way Forward. Annals of the American Thoracic Society, 2014, 11, 1171-1175.	3.2	15
81	Prostaglandin E2 Inhibits Î±-Smooth Muscle Actin Transcription during Myofibroblast Differentiation via Distinct Mechanisms of Modulation of Serum Response Factor and Myocardin-related Transcription Factor-A. Journal of Biological Chemistry, 2014, 289, 17151-17162.	3.4	84
82	Fibronectin ^{EDA} Promotes Chronic Cutaneous Fibrosis Through Toll-Like Receptor Signaling. Science Translational Medicine, 2014, 6, 232ra50.	12.4	195
83	Randomized, Double-Blind, Placebo-Controlled Trial of Soluble Tumor Necrosis Factor Receptor: Enbrel (Etanercept) for the Treatment of Idiopathic Pneumonia Syndrome after Allogeneic Stem Cell Transplantation: Blood and Marrow Transplant Clinical Trials Network Protocol. Biology of Blood and Marrow Transplantation, 2014, 20, 858-864.	2.0	78
84	Maternal Infection with Schistosoma japonicum Induces a Profibrotic Response in Neonates. Infection and Immunity, 2014, 82, 350-355.	2.2	16
85	Repair and Regeneration of the Respiratory System: Complexity, Plasticity, and Mechanisms of Lung Stem Cell Function. Cell Stem Cell, 2014, 15, 123-138.	11.1	748
86	Arsenic trioxide inhibits transforming growth factor-Î²1-induced fibroblast to myofibroblast differentiation in vitro and bleomycin induced lung fibrosis in vivo. Respiratory Research, 2014, 15, 51.	3.6	46
87	Future Directions in Idiopathic Pulmonary Fibrosis Research. An NHLBI Workshop Report. American Journal of Respiratory and Critical Care Medicine, 2014, 189, 214-222.	5.6	199
88	Lung microbiome and disease progression in idiopathic pulmonary fibrosis: an analysis of the COMET study. Lancet Respiratory Medicine,the, 2014, 2, 548-556.	10.7	353
89	Matrix Biology of Idiopathic Pulmonary Fibrosis. American Journal of Pathology, 2014, 184, 1643-1651.	3.8	91
90	miR-92a regulates TGF-Î²1-induced WISP1 expression in pulmonary fibrosis. International Journal of Biochemistry and Cell Biology, 2014, 53, 432-441.	2.8	95

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91	Fibrotic extracellular matrix activates a profibrotic positive feedback loop. Journal of Clinical Investigation, 2014, 124, 1622-1635.	8.2	444
92	Inflammation, wound repair, and fibrosis: reassessing the spectrum of tissue injury and resolution. Journal of Pathology, 2013, 229, 141-144.	4.5	91
93	The myofibroblast matrix: implications for tissue repair andÂfibrosis. Journal of Pathology, 2013, 229, 298-309.	4.5	560
94	X-Linked Inhibitor of Apoptosis Regulates Lung Fibroblast Resistance to Fas-Mediated Apoptosis. American Journal of Respiratory Cell and Molecular Biology, 2013, 49, 86-95.	2.9	60
95	Schistosoma japonicum Soluble Egg Antigens Attenuate Invasion in a First Trimester Human Placental Trophoblast Model. PLoS Neglected Tropical Diseases, 2013, 7, e2253.	3.0	8
96	Histone modifications are responsible for decreased Fas expression and apoptosis resistance in fibrotic lung fibroblasts. Cell Death and Disease, 2013, 4, e621-e621.	6.3	122
97	Human iPS cellÃ¢â‚¬â€œderived alveolar epithelium repopulates lung extracellular matrix. Journal of Clinical Investigation, 2013, 123, 4950-4962.	8.2	214
98	Periostin promotes fibrosis and predicts progression in patients with idiopathic pulmonary fibrosis. American Journal of Physiology - Lung Cellular and Molecular Physiology, 2012, 303, L1046-L1056.	2.9	223
99	Repeated Exposure to Aspergillus fumigatus Conidia Results in CD4⁺T Cell-Dependent and -Independent Pulmonary Arterial Remodeling in a Mixed Th1/Th2/Th17 Microenvironment That Requires Interleukin-4 (IL-4) and IL-10. Infection and Immunity, 2012, 80, 388-397.	2.2	32
100	Acellular Normal and Fibrotic Human Lung Matrices as a Culture System for In Vitro Investigation. American Journal of Respiratory and Critical Care Medicine, 2012, 186, 866-876.	5.6	552
101	Interleukin-17 Drives Pulmonary Eosinophilia following Repeated Exposure to Aspergillus fumigatus Conidia. Infection and Immunity, 2012, 80, 1424-1436.	2.2	76
102	PTEN Directly Activates the Actin Depolymerization Factor Cofilin-1 During PGE ₂ -Mediated Inhibition of Phagocytosis of Fungi. Science Signaling, 2012, 5, ra12.	3.6	61
103	Fibrotic and Sclerotic Manifestations of Chronic Graft-versus-Host Disease. Biology of Blood and Marrow Transplantation, 2012, 18, S46-S52.	2.0	20
104	Understanding Chronic GVHD from Different Angles. Biology of Blood and Marrow Transplantation, 2012, 18, S184-S188.	2.0	20
105	Soluble Tumor Necrosis Factor Receptor: Enbrel (Etanercept) for Subacute Pulmonary Dysfunction Following Allogeneic Stem Cell Transplantation. Biology of Blood and Marrow Transplantation, 2012, 18, 1044-1054.	2.0	48
106	Survivin expression induced by endothelin-1 promotes myofibroblast resistance to apoptosis. International Journal of Biochemistry and Cell Biology, 2012, 44, 158-169.	2.8	73
107	Recipientâ€derived EDA fibronectin promotes cardiac allograft fibrosis. Journal of Pathology, 2012, 226, 609-618.	4.5	50
108	Increased survivin expression contributes to apoptosis-resistance in IPF fibroblasts. Advances in Bioscience and Biotechnology (Print), 2012, 03, 657-664.	0.7	61

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109	Fibronectin splice variants: Understanding their multiple roles in health and disease using engineered mouse models. IUBMB Life, 2011, 63, 538-546.	3.4	141
110	PTEN Limits Alveolar Macrophage Function againstPseudomonas aeruginosaafter Bone Marrow Transplantation. American Journal of Respiratory Cell and Molecular Biology, 2011, 45, 1050-1058.	2.9	24
111	Accessory Cardiac Bronchus. American Journal of Respiratory and Critical Care Medicine, 2011, 183, 825-825.	5.6	6
112	Tissue Inhibitor of Matrix-Metalloproteaseâ€“1 Predicts Risk of Hepatic Fibrosis in Human Schistosoma japonicum Infection. Journal of Infectious Diseases, 2011, 203, 707-714.	4.0	31
113	Coevolution of T_H1, T_H2, and T_H17 Responses during Repeated Pulmonary Exposure toAspergillus fumigatusConidia. Infection and Immunity, 2011, 79, 125-135.	2.2	86
114	Innate Pathways Shape Sarcoidosis Signaling. American Journal of Respiratory and Critical Care Medicine, 2011, 183, 425-427.	5.6	4
115	Downregulation of FAK-related non-kinase mediates the migratory phenotype of human fibrotic lung fibroblasts. Experimental Cell Research, 2010, 316, 1600-1609.	2.6	41
116	Control of fibroblast fibronectin expression and alternative splicing via the PI3K/Akt/mTOR pathway. Experimental Cell Research, 2010, 316, 2644-2653.	2.6	59
117	EDAâ€containing cellular fibronectin induces fibroblast differentiation through binding to Î±₄Î²₇integrin receptor and MAPK/Erk 1/2â€dependent signaling. FASEB Journal, 2010, 24, 4503-4512.	0.5	130
118	Targeted Injury of Type II Alveolar Epithelial Cells Induces Pulmonary Fibrosis. American Journal of Respiratory and Critical Care Medicine, 2010, 181, 254-263.	5.6	399
119	Hypermethylation of PTGER2 Confers Prostaglandin E2 Resistance in Fibrotic Fibroblasts from Humans and Mice. American Journal of Pathology, 2010, 177, 2245-2255.	3.8	127
120	The antifibrotic effects of plasminogen activation occur via prostaglandin E2 synthesis in humans and mice. Journal of Clinical Investigation, 2010, 120, 1950-1960.	8.2	138
121	Prostaglandin E₂induces fibroblast apoptosis by modulating multiple survival pathways. FASEB Journal, 2009, 23, 4317-4326.	0.5	132
122	Phosphatase and Tensin Homologue on Chromosome 10 (PTEN) Directs Prostaglandin E2-mediated Fibroblast Responses via Regulation of E Prostanoid 2 Receptor Expression. Journal of Biological Chemistry, 2009, 284, 32264-32271.	3.4	20
123	The tumor suppressor protein PTEN inhibits rat hepatic stellate cell activation. Journal of Gastroenterology, 2009, 44, 847-855.	5.1	44
124	Infliximab therapy rescues cyclophosphamide failure in severe central nervous system sarcoidosis. Respiratory Medicine, 2009, 103, 268-273.	2.9	85
125	An Essential Role for Fibronectin Extra Type III Domain A in Pulmonary Fibrosis. American Journal of Respiratory and Critical Care Medicine, 2008, 177, 638-645.	5.6	257
126	The impact of soluble tumor necrosis factor receptor etanercept on the treatment of idiopathic pneumonia syndrome after allogeneic hematopoietic stem cell transplantation. Blood, 2008, 112, 3073-3081.	1.4	117

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127	Bronchoscopic Evaluation of Pulmonary Complications in Patients Undergoing Reduced-Intensity Versus Full-Intensity Transplants.. Blood, 2008, 112, 2163-2163.	1.4	3
128	PGE2 inhibition of TGF-Î²1-induced myofibroblast differentiation is Smad-independent but involves cell shape and adhesion-dependent signaling. American Journal of Physiology - Lung Cellular and Molecular Physiology, 2007, 293, L417-L428.	2.9	90
129	Pulmonary Vasculitis. , 2007, 36, 196-211.		0
130	Pulmonary Sarcoidosis. Seminars in Respiratory and Critical Care Medicine, 2007, 28, 053-074.	2.1	87
131	Diversity of the Angiogenic Phenotype in Nonâ€“Small Cell Lung Cancer. American Journal of Respiratory Cell and Molecular Biology, 2007, 36, 343-350.	2.9	32
132	Current and emerging strategies for the management of sarcoidosis. Expert Opinion on Pharmacotherapy, 2007, 8, 1293-1311.	1.8	21
133	Activation of Phosphatase and Tensin Homolog on Chromosome 10 Mediates the Inhibition of FcÎ³R Phagocytosis by Prostaglandin E2 in Alveolar Macrophages. Journal of Immunology, 2007, 179, 8350-8356.	0.8	44
134	Combinatorial activation of FAK and AKT by transforming growth factor-Î²1 confers an anoikis-resistant phenotype to myofibroblasts. Cellular Signalling, 2007, 19, 761-771.	3.6	220
135	Pharmacological Therapy for Wegener???s Granulomatosis. Drugs, 2006, 66, 1209-1228.	10.9	43
136	Infliximab in Sarcoidosis. American Journal of Respiratory and Critical Care Medicine, 2006, 174, 732-733.	5.6	11
137	Usual Interstitial Pneumonia. Seminars in Respiratory and Critical Care Medicine, 2006, 27, 634-651.	2.1	42
138	Negative Regulation of Myofibroblast Differentiation by PTEN (Phosphatase and Tensin Homolog) Tj ETQq0 0 0 rgBT /Overlock 10 Tf 50 112-121.	5.6	186
139	PTEN Regulates Fibroblast Elimination during Collagen Matrix Contraction. Journal of Biological Chemistry, 2006, 281, 33291-33301.	3.4	41
140	Management of Wegener Granulomatosis. Clinical Pulmonary Medicine, 2005, 12, 220-231.	0.3	1
141	Prostaglandin E2Inhibits Fibroblast Migration by E-Prostanoid 2 Receptorâ€“Mediated Increase in PTEN Activity. American Journal of Respiratory Cell and Molecular Biology, 2005, 32, 135-141.	2.9	124
142	Bleomycin-Induced E Prostanoid Receptor Changes Alter Fibroblast Responses to Prostaglandin E2. Journal of Immunology, 2005, 174, 5644-5649.	0.8	123
143	Corticosteroid Therapy for Sarcoidosis. Lung Biology in Health and Disease, 2005, , 689-716.	0.1	1
144	Wegener's Granulomatosis: Evolving Concepts in Treatment. Seminars in Respiratory and Critical Care Medicine, 2004, 25, 491-521.	2.1	30

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145	Plasminogen Activator Inhibitorâ€“1 Impairs Alveolar Epithelial Repair by Binding to Vitronectin. American Journal of Respiratory Cell and Molecular Biology, 2004, 31, 672-678.	2.9	62
146	Mechanisms of Pulmonary Fibrosis. Annual Review of Medicine, 2004, 55, 395-417.	12.2	640
147	Pathogenetic mechanisms in usual interstitial pneumonia/idiopathic pulmonary fibrosis. Journal of Pathology, 2003, 201, 343-354.	4.5	166
148	Pneumocystis Pneumonia Increases the Susceptibility of Mice to Sublethal Hyperoxia. Infection and Immunity, 2003, 71, 5970-5978.	2.2	10
149	Leptin Modulates Neutrophil Phagocytosis of Klebsiella pneumoniae. Infection and Immunity, 2003, 71, 4182-4185.	2.2	93
150	Myofibroblast Differentiation by Transforming Growth Factor-à¥†1 Is Dependent on Cell Adhesion and Integrin Signaling via Focal Adhesion Kinase. Journal of Biological Chemistry, 2003, 278, 12384-12389.	3.4	547
151	Integrin Î±₄Î²₁ Regulates Migration across Basement Membranes by Lung Fibroblasts. American Journal of Respiratory and Critical Care Medicine, 2003, 168, 436-442.	5.6	128
152	Bronchiolar Complications of Connective Tissue Diseases. Seminars in Respiratory and Critical Care Medicine, 2003, 24, 543-566.	2.1	51
153	Macrophage migration inhibitory factor and CXC chemokine expression in non-small cell lung cancer: role in angiogenesis and prognosis. Clinical Cancer Research, 2003, 9, 853-60.	7.0	117
154	Corticosteroids in idiopathic pulmonary fibrosis. Current Opinion in Pulmonary Medicine, 2001, 7, 298-308.	2.6	36
155	Sarcoidosis Involving Multiple Systems. Chest, 2001, 119, 1593-1597.	0.8	14
156	Improved survival in tumor-bearing SCID mice treated with interferon-Î³-inducible protein 10 (IP-10/CXCL10). Cancer Immunology, Immunotherapy, 2001, 50, 533-538.	4.2	63
157	Monocyte-Fibronectin Interactions, Via Î±5Î²1 Integrin, Induce Expression of CXC Chemokine-Dependent Angiogenic Activity. Journal of Immunology, 2001, 167, 5362-5366.	0.8	33
158	Non-Small Cell Lung Cancer Cells Induce Monocytes to Increase Expression of Angiogenic Activity. Journal of Immunology, 2001, 166, 7549-7555.	0.8	90
159	Timing of Lung Transplantation for Patients with Fibrotic Lung Diseases. Seminars in Respiratory and Critical Care Medicine, 2001, 22, 517-532.	2.1	8
160	Usual Interstitial Pneumonia. Seminars in Respiratory and Critical Care Medicine, 2001, 22, 357-386.	2.1	34
161	Dietary influences on morphine-induced analgesia in rats. Pharmacology Biochemistry and Behavior, 1991, 38, 681-684.	2.9	60