Paul J Wolters

List of Publications by Year
in descending order

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149 papers	16,548 citations	20817 60 h-index	17592 121 g-index

152 all docs	152 docs citations	152 times ranked	19430 citing authors

#	Article	IF	Citations
1	Interaction Between Epithelial and Mesenchymal Cells in Interstitial Lung Disease. , 2022, , 114-125.		0
2	Lung transplant recipients with idiopathic pulmonary fibrosis have impaired alloreactive immune responses. Journal of Heart and Lung Transplantation, 2022, 41, 641-653.	0.6	11
3	A simple method to generate human airway epithelial organoids with externally orientated apical membranes. American Journal of Physiology - Lung Cellular and Molecular Physiology, 2022, 322, L420-L437.	2.9	13
4	Common idiopathic pulmonary fibrosis risk variants are associated with hypersensitivity pneumonitis. Thorax, 2022, 77, 508-510.	5.6	14
5	Proteomic biomarkers of progressive fibrosing interstitial lung disease: a multicentre cohort analysis. Lancet Respiratory Medicine,the, 2022, 10, 593-602.	10.7	31
6	Oncostatin M expression induced by bacterial triggers drives airway inflammatory and mucus secretion in severe asthma. Science Translational Medicine, 2022, 14, eabf8188.	12.4	17
7	Human alveolar type 2 epithelium transdifferentiates into metaplastic KRT5+ basal cells. Nature Cell Biology, 2022, 24, 10-23.	10.3	108
8	Acquisition of cellular properties during alveolar formation requires differential activity and distribution of mitochondria. ELife, 2022, 11, .	6.0	9
9	Sine oculis homeobox homolog 1 plays a critical role in pulmonary fibrosis. JCI Insight, 2022, 7, .	5.0	4
10	Colocalization of Gene Expression and DNA Methylation with Genetic Risk Variants Supports Functional Roles of MUC5B and DSP in Idiopathic Pulmonary Fibrosis. American Journal of Respiratory and Critical Care Medicine, 2022, 206, 1259-1270.	5.6	12
11	Essential Components of an Interstitial Lung Disease Clinic. Chest, 2021, 159, 1517-1530.	0.8	18
12	Extracellular BMP1 is the major proteinase for COOH-terminal proteolysis of type I procollagen in lung fibroblasts. American Journal of Physiology - Cell Physiology, 2021, 320, C162-C174.	4.6	7
13	Leukocyte telomere length and mycophenolate therapy in chronic hypersensitivity pneumonitis. European Respiratory Journal, 2021, 57, 2002872.	6.7	32
14	Autoantibodies targeting telomere-associated proteins in systemic sclerosis. Annals of the Rheumatic Diseases, 2021, 80, 912-919.	0.9	19
15	Molecular mapping of interstitial lung disease reveals a phenotypically distinct senescent basal epithelial cell population. JCI Insight, 2021, 6, .	5.0	42
16	Molecular markers of telomere dysfunction and senescence are common findings in the usual interstitial pneumonia pattern of lung fibrosis. Histopathology, 2021, 79, 67-76.	2.9	25
17	MAV(S)erick mitochondria: an unconventional role for mitochondrial antiviral signalling protein in pulmonary fibrosis. European Respiratory Journal, 2021, 57, 2004500.	6.7	0
18	Genetically increased circulating FUT3 level leads to reduced risk of Idiopathic Pulmonary Fibrosis: a Mendelian Randomisation Study. European Respiratory Journal, 2021, , 2003979.	6.7	9

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19	Peripheral blood leucocyte telomere length is associated with progression of interstitial lung disease in systemic sclerosis. Thorax, 2021, 76, 1186-1192.	5.6	18
20	Invariant natural killer TÂcells coordinate removal of senescent cells. Med, 2021, 2, 938-950.e8.	4.4	28
21	TGFÎ²2 and TGFÎ²3 isoforms drive fibrotic disease pathogenesis. Science Translational Medicine, 2021, 13, .	12.4	56
22	The prognostic role of matrix metalloproteinase-7 in scleroderma-associated interstitial lung disease. European Respiratory Journal, 2021, 58, 2101560.	6.7	6
23	Treatment of fibrotic interstitial lung disease: current approaches and future directions. Lancet, The, 2021, 398, 1450-1460.	13.7	47
24	MUC5B promoter variant rs35705950 and rheumatoid arthritis associated interstitial lung disease survival and progression. Seminars in Arthritis and Rheumatism, 2021, 51, 996-1004.	3.4	17
25	Blocking LOXL2 and TGFÎ²1 signalling induces collagen I turnover in precision-cut lung slices derived from patients with idiopathic pulmonary fibrosis. Thorax, 2021, 76, 729-732.	5.6	28
26	Dual inhibition of Î±vÎ²6 and Î±vÎ²1 reduces fibrogenesis in lung tissue explants from patients with IPF. Respiratory Research, 2021, 22, 265.	3.6	28
27	Molecular programs of fibrotic change in aging human lung. Nature Communications, 2021, 12, 6309.	12.8	33
28	Cut From the Same Cloth: Similarities Between Hypersensitivity Pneumonitis and IPF. American Journal of Respiratory and Critical Care Medicine, 2021, , .	5.6	1
29	Peripheral blood leukocyte telomere length is associated with survival of sepsis patients. European Respiratory Journal, 2020, 55, 1901044.	6.7	27
30	Diaphragmatic Atrophy May Limit Progression of Idiopathic Pulmonary Fibrosis. American Journal of Respiratory and Critical Care Medicine, 2020, 201, e72-e73.	5.6	1
31	Lymphatic Proliferation Ameliorates Pulmonary Fibrosis after Lung Injury. American Journal of Pathology, 2020, 190, 2355-2375.	3.8	21
32	LUNG FUNCTION TRAJECTORY IN THE PIRFENIDONE FOR RESTRICTIVE CHRONIC LUNG ALLOGRAFT DYSFUNCTION (PIRCLAD) TRIAL: INTERIM ANALYSIS. Chest, 2020, 158, A2391-A2392.	0.8	0
33	Gli1+ mesenchymal stromal cells form a pathological niche to promote airway progenitor metaplasia in the fibrotic lung. Nature Cell Biology, 2020, 22, 1295-1306.	10.3	62
34	The Pulmonary Fibrosis Foundation Patient Registry. Rationale, Design, and Methods. Annals of the American Thoracic Society, 2020, 17, 1620-1628.	3.2	27
35	SAFETY AND TOLERABILITY OF PIRFENIDONE FOR RESTRICTIVE CHRONIC LUNG ALLOGRAFT DYSFUNCTION (PIRCLAD): INTERIM RESULTS. Chest, 2020, 158, A2389-A2390.	0.8	2
36	Osteopontin Links Myeloid Activation and Disease Progression in Systemic Sclerosis. Cell Reports Medicine, 2020, 1, 100140.	6.5	42

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37	Frailty after lung transplantation is associated with impaired health-related quality of life and mortality. Thorax, 2020, 75, 669-678.	5.6	24
38	Telomere length in patients with unclassifiable interstitial lung disease: a cohort study. European Respiratory Journal, 2020, 56, 2000268.	6.7	12
39	Long-term ozone exposure is positively associated with telomere length in critically ill patients. Environment International, 2020, 141, 105780.	10.0	18
40	Airway Epithelial Telomere Dysfunction Drives Remodeling Similar to Chronic Lung Allograft Dysfunction. American Journal of Respiratory Cell and Molecular Biology, 2020, 63, 490-501.	2.9	17
41	Reversal of TGFÎ²1-Driven Profibrotic State in Patients with Pulmonary Fibrosis. New England Journal of Medicine, 2020, 382, 1068-1070.	27.0	42
42	Chronic Hypersensitivity Pneumonitis, an Interstitial Lung Disease with Distinct Molecular Signatures. American Journal of Respiratory and Critical Care Medicine, 2020, 202, 1430-1444.	5.6	66
43	A Systematically Derived Exposure Assessment Instrument for Chronic Hypersensitivity Pneumonitis. Chest, 2020, 157, 1506-1512.	0.8	33
44	Collagen-producing lung cell atlas identifies multiple subsets with distinct localization and relevance to fibrosis. Nature Communications, 2020, 11, 1920.	12.8	346
45	Age-dependent regulation of cell-mediated collagen turnover. JCI Insight, 2020, 5, .	5.0	26
46	Histopathological and molecular analysis of idiopathic pulmonary fibrosis lungs from patients treated with pirfenidone or nintedanib. Histopathology, 2019, 74, 341-349.	2.9	20
47	A nonlinear relationship between visceral adipose tissue and frailty in adult lung transplant candidates. American Journal of Transplantation, 2019, 19, 3155-3161.	4.7	25
48	Rare Protein-Altering Telomere-related Gene Variants in Patients with Chronic Hypersensitivity Pneumonitis. American Journal of Respiratory and Critical Care Medicine, 2019, 200, 1154-1163.	5.6	81
49	Prevalence and Clinical Significance ofÂAntineutrophil Cytoplasmic Antibodies inÂNorth American Patients With Idiopathic Pulmonary Fibrosis. Chest, 2019, 156, 715-723.	0.8	45
50	Regulatory T cells in skin are uniquely poised to suppress profibrotic immune responses. Science Immunology, 2019, 4, .	11.9	78
51	Resequencing Study Confirms That Host Defense and Cell Senescence Gene Variants Contribute to the Risk of Idiopathic Pulmonary Fibrosis. American Journal of Respiratory and Critical Care Medicine, 2019, 200, 199-208.	5.6	90
52	Molecular endpoints for establishing target engagement by novel idiopathic pulmonary fibrosis therapies. European Respiratory Journal, 2019, 53, 1900283.	6.7	0
53	Association between greenhouse working exposure and bronchial asthma: A pilot, cross-sectional survey of 5,420 greenhouse farmers from northeast China. Journal of Occupational and Environmental Hygiene, 2019, 16, 286-293.	1.0	3
54	Pulmonary physiology is poorly associated with radiological extent of disease in systemic sclerosis-associated interstitial lung disease. European Respiratory Journal, 2019, 53, 1802182.	6.7	11

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55	Significance of bronchiolocentric fibrosis in patients with histopathological usual interstitial pneumonia. Histopathology, 2019, 74, 1088-1097.	2.9	16
56	Reference-based analysis of lung single-cell sequencing reveals a transitional profibrotic macrophage. Nature Immunology, 2019, 20, 163-172.	14.5	2,330
57	Telomere length and genetic variant associations with interstitial lung disease progression and survival. European Respiratory Journal, 2019, 53, 1801641.	6.7	119
58	Air Pollution Exposure Is Associated With Lower Lung Function, but Not Changes in Lung Function, in Patients With Idiopathic Pulmonary Fibrosis. Chest, 2018, 154, 119-125.	0.8	76
59	Time for a change: is idiopathic pulmonary fibrosis still idiopathic and only fibrotic?. Lancet Respiratory Medicine,the, 2018, 6, 154-160.	10.7	137
60	Identification of Diagnostic Criteria for Chronic Hypersensitivity Pneumonitis. An International Modified Delphi Survey. American Journal of Respiratory and Critical Care Medicine, 2018, 197, 1036-1044.	5.6	174
61	Pirfenidone in the kaleidoscope: reflecting mechanisms through differentÂangles. European Respiratory Journal, 2018, 52, 1802046.	6.7	8
62	MUC5B Promoter Variant and Rheumatoid Arthritis with Interstitial Lung Disease. New England Journal of Medicine, 2018, 379, 2209-2219.	27.0	326
63	Response. Chest, 2018, 154, 727-728.	0.8	0
64	Prognostic and predictive biomarkers for patients with idiopathic pulmonary fibrosis treated with pirfenidone: post-hoc assessment of the CAPACITY and ASCEND trials. Lancet Respiratory Medicine,the, 2018, 6, 615-626.	10.7	87
65	Increased Extracellular Vesicles Mediate WNT5A Signaling in Idiopathic Pulmonary Fibrosis. American Journal of Respiratory and Critical Care Medicine, 2018, 198, 1527-1538.	5.6	127
66	Laparoscopic anti-reflux surgery for the treatment of idiopathic pulmonary fibrosis (WRAP-IPF): a multicentre, randomised, controlled phase 2 trial. Lancet Respiratory Medicine,the, 2018, 6, 707-714.	10.7	109
67	Analysis of protein-altering variants in telomerase genes and their association with MUC5B common variant status in patients with idiopathic pulmonary fibrosis: a candidate gene sequencing study. Lancet Respiratory Medicine,the, 2018, 6, 603-614.	10.7	133
68	Pathologic Findings and Prognosis in a LargeÂProspective Cohort of Chronic Hypersensitivity Pneumonitis. Chest, 2017, 152, 502-509.	0.8	131
69	The use of pretest probability increases the value of high-resolution CT in diagnosing usual interstitial pneumonia. Thorax, 2017, 72, 424-429.	5.6	103
70	CXCL14 is a candidate biomarker for Hedgehog signalling in idiopathic pulmonary fibrosis. Thorax, 2017, 72, 780-787.	5.6	47
71	Spontaneous Chitin Accumulation in Airways and Age-Related Fibrotic Lung Disease. Cell, 2017, 169, 497-509.e13.	28.9	87
72	The performance of the GAP model in patients with rheumatoid arthritis associated interstitial lung disease. Respiratory Medicine, 2017, 127, 51-56.	2.9	49

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73	Short lung transplant donor telomere length is associated with decreased CLAD-free survival. Thorax, 2017, 72, 1052-1054.	5.6	57
74	Mortality Risk Prediction in Scleroderma-Related Interstitial LungÂDisease. Chest, 2017, 152, 999-1007.	0.8	61
75	A recurring theme in pulmonary fibrosis genetics. European Respiratory Journal, 2017, 49, 1700545.	6.7	6
76	The MUC5B promoter polymorphism and telomere length in patients with chronic hypersensitivity pneumonitis: an observational cohort-control study. Lancet Respiratory Medicine,the, 2017, 5, 639-647.	10.7	206
77	Use of Mycophenolate Mofetil or Azathioprine for the Management of Chronic Hypersensitivity Pneumonitis. Chest, 2017, 151, 619-625.	0.8	177
78	Genome-wide imputation study identifies novel HLA locus for pulmonary fibrosis and potential role for auto-immunity in fibrotic idiopathic interstitial pneumonia. BMC Genetics, 2016, 17, 74.	2.7	84
79	miR-34 miRNAs Regulate Cellular Senescence in Type II Alveolar Epithelial Cells of Patients with Idiopathic Pulmonary Fibrosis. PLoS ONE, 2016, 11, e0158367.	2.5	106
80	Amplification of TGFÎ² Induced ITGB6 Gene Transcription May Promote Pulmonary Fibrosis. PLoS ONE, 2016, 11, e0158047.	2.5	34
81	Donor-Reactive Regulatory T Cell Frequency Increases During Acute Cellular Rejection of Lung Allografts. Transplantation, 2016, 100, 2090-2098.	1.0	15
82	Non-invasive Imaging of Idiopathic Pulmonary Fibrosis Using Cathepsin Protease Probes. Scientific Reports, 2016, 6, 19755.	3.3	97
83	A diagnostic model for chronic hypersensitivity pneumonitis. Thorax, 2016, 71, 951-954.	5.6	70
84	Brief Report: Wholeâ€Exome Sequencing for Identification of Potential Causal Variants for Diffuse Cutaneous Systemic Sclerosis. Arthritis and Rheumatology, 2016, 68, 2257-2262.	5.6	35
85	Telomere dysfunction in alveolar epithelial cells causes lung remodeling and fibrosis. JCI Insight, 2016, 1, e86704.	5.0	192
86	Integrated, multicohort analysis of systemic sclerosis identifies robust transcriptional signature of disease severity. JCI Insight, 2016, 1, e89073.	5.0	57
87	Interstitial lung diseases in the hospitalized patient. BMC Medicine, 2015, 13, 245.	5.5	41
88	Classification of usual interstitial pneumonia in patients with interstitial lung disease: assessment of a machine learning approach using high-dimensional transcriptional data. Lancet Respiratory Medicine,the, 2015, 3, 473-482.	10.7	112
89	The effect of bronchodilators on forced vital capacity measurement in patients with idiopathic pulmonary fibrosis. Respiratory Medicine, 2015, 109, 1058-1062.	2.9	9
90	Lung Transplantation for Hypersensitivity Pneumonitis. Chest, 2015, 147, 1558-1565.	0.8	67

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91	Overexpression of Inhibitor of DNA-Binding 2 Attenuates Pulmonary Fibrosis through Regulation of c-Abl and Twist. American Journal of Pathology, 2015, 185, 1001-1011.	3.8	31
92	Transforming Growth Factor-Î² and Interleukin-1Î² Signaling Pathways Converge on the Chemokine CCL20 Promoter. Journal of Biological Chemistry, 2015, 290, 14717-14728.	3.4	22
93	Survival in interstitial pneumonia with features of autoimmune disease: A comparison of proposed criteria. Respiratory Medicine, 2015, 109, 1326-1331.	2.9	40
94	Frailty Phenotypes, Disability, and Outcomes in Adult Candidates for Lung Transplantation. American Journal of Respiratory and Critical Care Medicine, 2015, 192, 1325-1334.	5.6	181
95	Heterogeneous gene expression signatures correspond to distinct lung pathologies and biomarkers of disease severity in idiopathic pulmonary fibrosis. Thorax, 2015, 70, 48-56.	5.6	207
96	Accumulation of BDCA1+ Dendritic Cells in Interstitial Fibrotic Lung Diseases and Th2-High Asthma. PLoS ONE, 2014, 9, e99084.	2.5	34
97	Subacute Onset of Pulmonary Langerhans Cell Histiocytosis with Resolution after Smoking Cessation. American Journal of Respiratory and Critical Care Medicine, 2014, 190, e64-e64.	5.6	9
98	Selective Targeting of TGF-Î² Activation to Treat Fibroinflammatory Airway Disease. Science Translational Medicine, 2014, 6, 241ra79.	12.4	79
99	Endogenously Expressed IL-13RÎ±2 Attenuates IL-13â€“Mediated Responses but Does Not Activate Signaling in Human Lung Fibroblasts. Journal of Immunology, 2014, 193, 111-119.	0.8	69
100	Pathogenesis of Idiopathic Pulmonary Fibrosis. Annual Review of Pathology: Mechanisms of Disease, 2014, 9, 157-179.	22.4	621
101	Effect of telomere length on survival in patients with idiopathic pulmonary fibrosis: an observational cohort study with independent validation. Lancet Respiratory Medicine,the, 2014, 2, 557-565.	10.7	225
102	Dipeptidyl peptidase I controls survival from Klebsiella pneumoniae lung infection by processing surfactant protein D. Biochemical and Biophysical Research Communications, 2014, 450, 818-823.	2.1	8
103	Diagnosis of idiopathic pulmonary fibrosis with high-resolution CT. Lancet Respiratory Medicine,the, 2014, 2, e5.	10.7	8
104	Fibulin-1 Predicts Disease Progression in Patients With Idiopathic Pulmonary Fibrosis. Chest, 2014, 146, 1055-1063.	0.8	42
105	Predicting Survival Across Chronic Interstitial Lung Disease. Chest, 2014, 145, 723-728.	0.8	366
106	Serum IgE clearance is facilitated by human FcÎµRI internalization. Journal of Clinical Investigation, 2014, 124, 1187-1198.	8.2	74
107	Increased susceptibility to Klebsiella pneumonia and mortality in GSNOR-deficient mice. Biochemical and Biophysical Research Communications, 2013, 442, 122-126.	2.1	13
108	Genome-wide association study identifies multiple susceptibility loci for pulmonary fibrosis. Nature Genetics, 2013, 45, 613-620.	21.4	667

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109	BPIFB1 Is a Lung-Specific Autoantigen Associated with Interstitial Lung Disease. Science Translational Medicine, 2013, 5, 206ra139.	12.4	87
110	Noradrenergic Neurons Regulate Monocyte Trafficking and Mortality during Gram-Negative Peritonitis in Mice. Journal of Immunology, 2013, 190, 4717-4724.	0.8	28
111	Prevalence and prognosis of unclassifiable interstitial lung disease. European Respiratory Journal, 2013, 42, 750-757.	6.7	238
112	Inflection points in sepsis biology: from local defense to systemic organ injury. American Journal of Physiology - Lung Cellular and Molecular Physiology, 2012, 303, L355-L363.	2.9	111
113	In search of the fibrotic epithelial cell: opportunities for a collaborative network. Thorax, 2012, 67, 179-182.	5.6	16
114	Relative versus absolute change in forced vital capacity in idiopathic pulmonary fibrosis. Thorax, 2012, 67, 407-411.	5.6	160
115	Cleaved cytokeratin-18 is a mechanistically informative biomarker in idiopathic pulmonary fibrosis. Respiratory Research, 2012, 13, 105.	3.6	32
116	Calcium-activated chloride channel TMEM16A modulates mucin secretion and airway smooth muscle contraction. Proceedings of the National Academy of Sciences of the United States of America, 2012, 109, 16354-16359.	7.1	290
117	Lung mast cell density defines a subpopulation of patients with idiopathic pulmonary fibrosis. Histopathology, 2012, 61, 98-106.	2.9	48
118	Integrin Î±9Î²1 in airway smooth muscle suppresses exaggerated airway narrowing. Journal of Clinical Investigation, 2012, 122, 2916-2927.	8.2	44
119	Gastroesophageal Reflux Therapy Is Associated with Longer Survival in Patients with Idiopathic Pulmonary Fibrosis. American Journal of Respiratory and Critical Care Medicine, 2011, 184, 1390-1394.	5.6	382
120	Viral Infection in Acute Exacerbation of Idiopathic Pulmonary Fibrosis. American Journal of Respiratory and Critical Care Medicine, 2011, 183, 1698-1702.	5.6	230
121	Parasitic Infection Improves Survival from Septic Peritonitis by Enhancing Mast Cell Responses to Bacteria in Mice. PLoS ONE, 2011, 6, e27564.	2.5	18
122	Interleukin-1Î² Induces Increased Transcriptional Activation of the Transforming Growth Factor-Î²-activating Integrin Subunit Î²8 through Altering Chromatin Architecture. Journal of Biological Chemistry, 2011, 286, 36864-36874.	3.4	35
123	Systemic mast cell degranulation increases mortality during polymicrobial septic peritonitis in mice. Journal of Leukocyte Biology, 2011, 90, 591-597.	3.3	22
124	Mouse and human lung fibroblasts regulate dendritic cell trafficking, airway inflammation, and fibrosis through integrin Î±vÎ²8â€“mediated activation of TGF-Î². Journal of Clinical Investigation, 2011, 121, 2863-2875.	8.2	157
125	Alveolar epithelial cells express mesenchymal proteins in patients with idiopathic pulmonary fibrosis. American Journal of Physiology - Lung Cellular and Molecular Physiology, 2011, 301, L71-L78.	2.9	135
126	Undifferentiated Connective Tissue Disease-Associated Interstitial Lung Disease: Changes in Lung Function. Lung, 2010, 188, 143-149.	3.3	50

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127	Transcription of the Transforming Growth Factor Î² Activating Integrin Î²8 Subunit Is Regulated by SP3, AP-1, and the p38 Pathway. Journal of Biological Chemistry, 2010, 285, 24695-24706.	3.4	27
128	Neutrophil-Derived IL-6 Limits Alveolar Barrier Disruption in Experimental Ventilator-Induced Lung Injury. Journal of Immunology, 2009, 182, 8056-8062.	0.8	49
129	Genetic deficiency and pharmacological stabilization of mast cells reduce diet-induced obesity and diabetes in mice. Nature Medicine, 2009, 15, 940-945.	30.7	663
130	Epithelial cell Î±3Î²1 integrin links Î²-catenin and Smad signaling to promote myofibroblast formation and pulmonary fibrosis. Journal of Clinical Investigation, 2009, 119, 213-24.	8.2	342
131	Mfge8 diminishes the severity of tissue fibrosis in mice by binding and targeting collagen for uptake by macrophages. Journal of Clinical Investigation, 2009, 119, 3713-3722.	8.2	194
132	Mast Cell IL-6 Improves Survival from Klebsiella Pneumonia and Sepsis by Enhancing Neutrophil Killing. Journal of Immunology, 2008, 181, 5598-5605.	0.8	134
133	Relationships between Early Inflammatory Response to Bleomycin and Sensitivity to Lung Fibrosis. American Journal of Respiratory and Critical Care Medicine, 2007, 176, 1098-1107.	5.6	22
134	Idiopathic Nonspecific Interstitial Pneumonia. American Journal of Respiratory and Critical Care Medicine, 2007, 176, 691-697.	5.6	345
135	Anchors Away. New England Journal of Medicine, 2007, 356, 504-509.	27.0	9
136	Mast cells promote atherosclerosis by releasing proinflammatory cytokines. Nature Medicine, 2007, 13, 719-724.	30.7	379
137	Mast cells modulate the pathogenesis of elastase-induced abdominal aortic aneurysms in mice. Journal of Clinical Investigation, 2007, 117, 3359-3368.	8.2	209
138	Squamous metaplasia amplifies pathologic epithelial-mesenchymal interactions in COPD patients. Journal of Clinical Investigation, 2007, 117, 3551-3562.	8.2	222
139	Alveolar epithelial cell mesenchymal transition develops in vivo during pulmonary fibrosis and is regulated by the extracellular matrix. Proceedings of the National Academy of Sciences of the United States of America, 2006, 103, 13180-13185.	7.1	1,118
140	Mast Cells Protect Mice from Mycoplasma Pneumonia. American Journal of Respiratory and Critical Care Medicine, 2006, 173, 219-225.	5.6	78
141	Neutrophil histamine contributes to inflammation in mycoplasma pneumonia. Journal of Experimental Medicine, 2006, 203, 2907-2917.	8.5	89
142	Cutting Edge: Deficiency of Macrophage Migration Inhibitory Factor Impairs Murine Airway Allergic Responses. Journal of Immunology, 2006, 177, 5779-5784.	0.8	33
143	Mast cell dipeptidyl peptidase I mediates survival from sepsis. Journal of Clinical Investigation, 2004, 113, 628-634.	8.2	75
144	Mast cell dipeptidyl peptidase I mediates survival from sepsis. Journal of Clinical Investigation, 2004, 113, 628-634.	8.2	127

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145	Structure and Activity of Human Pancreasin, a Novel Tryptic Serine Peptidase Expressed Primarily by the Pancreas. Journal of Biological Chemistry, 2003, 278, 3363-3371.	3.4	25
146	Dipeptidyl Peptidase I Is Essential for Activation of Mast Cell Chymases, but Not Tryptases, in Mice. Journal of Biological Chemistry, 2001, 276, 18551-18556.	3.4	176
147	Characterization of Human Î³-Tryptases, Novel Members of the Chromosome 16p Mast Cell Tryptase and Prostasin Gene Families. Journal of Immunology, 2000, 164, 6566-6575.	0.8	111
148	Dipeptidyl Peptidase I Cleaves Matrix-Associated Proteins and Is Expressed Mainly by Mast Cells in Normal Dog Airways. American Journal of Respiratory Cell and Molecular Biology, 2000, 22, 183-190.	2.9	49
149	Regulated Expression, Processing, and Secretion of Dog Mast Cell Dipeptidyl Peptidase I. Journal of Biological Chemistry, 1998, 273, 15514-15520.	3.4	53