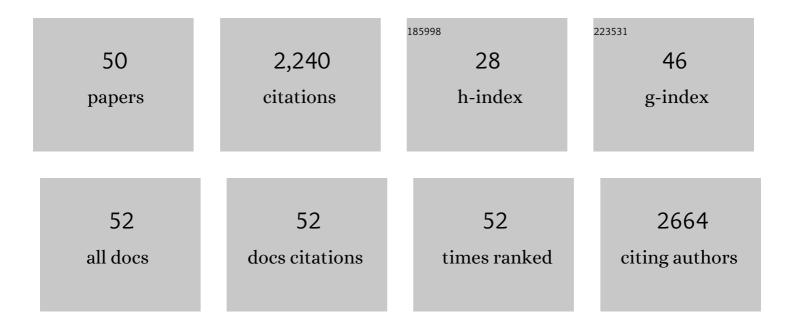
## olivier Tabary

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Alveolar epithelial cells: Master regulators of lung homeostasis. International Journal of Biochemistry and Cell Biology, 2013, 45, 2568-2573.	1.2	187
2	Selective Up-Regulation of Chemokine IL-8 Expression in Cystic Fibrosis Bronchial Gland Cells in Vivo and in Vitro. American Journal of Pathology, 1998, 153, 921-930.	1.9	158
3	Distinct cytokine production by lung and blood neutrophils from children with cystic fibrosis. American Journal of Physiology - Lung Cellular and Molecular Physiology, 2003, 284, L997-L1003.	1.3	108
4	Involvement of toll-like receptor 4 in the inflammatory reaction induced by hydroxyapatite particles. Biomaterials, 2007, 28, 400-404.	5.7	105
5	Airway epithelial cell inflammatory signalling in cystic fibrosis. International Journal of Biochemistry and Cell Biology, 2008, 40, 1703-1715.	1.2	102
6	Neutrophil Elastase Degrades Cystic Fibrosis Transmembrane Conductance Regulator via Calpains and Disables Channel Function <i>In Vitro</i> and <i>In Vivo</i> . American Journal of Respiratory and Critical Care Medicine, 2013, 187, 170-179.	2.5	97
7	High Susceptibility for Cystic Fibrosis Human Airway Gland Cells to Produce IL-8 Through the lκB Kinase α Pathway in Response to Extracellular NaCl Content. Journal of Immunology, 2000, 164, 3377-3384.	0.4	95
8	Genistein Inhibits Constitutive and Inducible NFκB Activation and Decreases IL-8 Production by Human Cystic Fibrosis Bronchial Gland Cells. American Journal of Pathology, 1999, 155, 473-481.	1.9	91
9	Calcium-dependent regulation of NF-κB activation in cystic fibrosis airway epithelial cells. Cellular Signalling, 2006, 18, 652-660.	1.7	77
10	Neutrophils in Cystic Fibrosis Display a Distinct Gene Expression Pattern. Molecular Medicine, 2008, 14, 36-44.	1.9	69
11	Lung disease modifier genes in cystic fibrosis. International Journal of Biochemistry and Cell Biology, 2014, 52, 83-93.	1.2	66
12	Transcription factors and miRNAs that regulate fetal to adult <i>CFTR</i> expression change are new targets for cystic fibrosis. European Respiratory Journal, 2015, 45, 116-128.	3.1	65
13	Enhanced IL-1β-induced IL-8 production in cystic fibrosis lung epithelial cells is dependent of both mitogen-activated protein kinases and NF-κB signaling. Biochemical and Biophysical Research Communications, 2007, 357, 402-407.	1.0	61
14	MicroRNA-9 downregulates the ANO1 chloride channel and contributes to cystic fibrosis lung pathology. Nature Communications, 2017, 8, 710.	5.8	56
15	Translating the genetics of cystic fibrosis to personalized medicine. Translational Research, 2016, 168, 40-49.	2.2	54
16	Relationship between I?B? deficiency, NF?B activity and interleukin-8 production in CF human airway epithelial cells. Pflugers Archiv European Journal of Physiology, 2001, 443, S40-S44.	1.3	52
17	Adherence of airway neutrophils and inflammatory response are increased in CF airway epithelial cell-neutrophil interactions. American Journal of Physiology - Lung Cellular and Molecular Physiology, 2006, 290, L588-L596.	1.3	50
18	Circulating and airway neutrophils in cystic fibrosis display different TLR expression and responsiveness to interleukin-10. Cytokine, 2008, 41, 54-60.	1.4	47

OLIVIER TABARY

#	Article	IF	CITATIONS
19	Azithromycin analogue <scp>CSY</scp> 0073 attenuates lung inflammation induced by <scp>LPS</scp> challenge. British Journal of Pharmacology, 2014, 171, 1783-1794.	2.7	44
20	Oxidative stress induces extracellular signal-regulated kinase 1/2 mitogen-activated protein kinase in cystic fibrosis lung epithelial cells: Potential mechanism for excessive IL-8 expression. International Journal of Biochemistry and Cell Biology, 2008, 40, 432-446.	1.2	41
21	Anoctamin 1 dysregulation alters bronchial epithelial repair in cystic fibrosis. Biochimica Et Biophysica Acta - Molecular Basis of Disease, 2013, 1832, 2340-2351.	1.8	40
22	Fluticasone reduces IL-6 and IL-8 production of cystic fibrosis bronchial epithelial cells <i>via</i> IKK-β kinase pathway. European Respiratory Journal, 2003, 21, 574-581.	3.1	39
23	New Insights about miRNAs in Cystic Fibrosis. American Journal of Pathology, 2015, 185, 897-908.	1.9	37
24	Small RNA and transcriptome sequencing reveal the role of miRâ€199aâ€3p in inflammatory processes in cystic fibrosis airways. Journal of Pathology, 2018, 245, 410-420.	2.1	35
25	Novel Anti-Inflammatory Approaches for Cystic Fibrosis Lung Disease: Identification of Molecular Targets and Design of Innovative Therapies. Frontiers in Pharmacology, 2020, 11, 1096.	1.6	30
26	Interleukin-10 Inhibits Elevated Chemokine Interleukin-8 and Regulated on Activation Normal T Cell Expressed and Secreted Production in Cystic Fibrosis Bronchial Epithelial Cells by Targeting the IkB Kinase α/β Complex. American Journal of Pathology, 2003, 162, 293-302.	1.9	29
27	Emerging microRNA Therapeutic Approaches for Cystic Fibrosis. Frontiers in Pharmacology, 2018, 9, 1113.	1.6	29
28	Glucocorticoid receptor gene polymorphisms associated with progression of lung disease in young patients with cystic fibrosis. Respiratory Research, 2007, 8, 88.	1.4	28
29	Hyperinflammation in airways of cystic fibrosis patients: what's new?. Expert Review of Molecular Diagnostics, 2008, 8, 359-363.	1.5	27
30	Enhanced F508del-CFTR Channel Activity Ameliorates Bone Pathology in Murine Cystic Fibrosis. American Journal of Pathology, 2014, 184, 1132-1141.	1.9	27
31	Intracellular colocalization and interaction of IGF-binding protein-2 with the cyclin-dependent kinase inhibitor p21CIP1/WAF1 during growth inhibition. Biochemical Journal, 2005, 392, 457-465.	1.7	25
32	Proinflammatory Effect of Sodium 4-Phenylbutyrate in ΔF508-Cystic Fibrosis Transmembrane Conductance Regulator Lung Epithelial Cells: Involvement of Extracellular Signal-Regulated Protein Kinase 1/2 and c-Jun-NH <sub>2</sub> -Terminal Kinase Signaling. Journal of Pharmacology and Experimental Therapeutics, 2008, 326, 949-956.	1.3	25
33	Glucocorticoids reduce inflammation in cystic fibrosis bronchial epithelial cells. Cellular Signalling, 2012, 24, 1093-1099.	1.7	25
34	Cystic Fibrosis and Bone Disease: Defective Osteoblast Maturation with the F508del Mutation in Cystic Fibrosis Transmembrane Conductance Regulator. American Journal of Respiratory and Critical Care Medicine, 2014, 189, 746-748.	2.5	24
35	Normal and Cystic Fibrosis Human Bronchial Epithelial Cells Infected with Pseudomonas aeruginosa Exhibit Distinct Gene Activation Patterns. PLoS ONE, 2015, 10, e0140979.	1.1	22
36	Macrolides: New therapeutic perspectives in lung diseases. International Journal of Biochemistry and Cell Biology, 2011, 43, 1241-1246.	1.2	21

OLIVIER TABARY

#	Article	IF	CITATIONS
37	Cystic fibrosis bone disease: is the CFTR corrector C18 an option for therapy?. European Respiratory Journal, 2015, 45, 845-848.	3.1	19
38	Physiomer® reduces the chemokine interleukin-8 production by activated human respiratory epithelial cells. European Respiratory Journal, 2001, 18, 661-666.	3.1	16
39	Gap Junctional Communication Does not Contribute to the Interaction Between Neutrophils and Airway Epithelial Cells. Cell Communication and Adhesion, 2006, 13, 1-12.	1.0	15
40	Azithromycin fails to reduce inflammation in cystic fibrosis airway epithelial cells. European Journal of Pharmacology, 2012, 674, 1-6.	1.7	15
41	Reduction of chemokine IL-8 and RANTES expression in human bronchial epithelial cells by a sea-water derived saline through inhibited nuclear factor-κB activation. Biochemical and Biophysical Research Communications, 2003, 309, 310-316.	1.0	14
42	Oxidative stress response results in increased p21WAF1/CIP1 degradation in cystic fibrosis lung epithelial cells. Free Radical Biology and Medicine, 2006, 40, 75-86.	1.3	13
43	Cystic Fibrosis Transmembrane Conductance Regulator Controls Lung Proteasomal Degradation and Nuclear Factor-I <sup>®</sup> B Activity in Conditions of Oxidative Stress. American Journal of Pathology, 2008, 172, 1184-1194.	1.9	13
44	Restoration of Chloride Efflux by Azithromycin in Airway Epithelial Cells of Cystic Fibrosis Patients. Antimicrobial Agents and Chemotherapy, 2011, 55, 1792-1793.	1.4	11
45	miR-636: A Newly-Identified Actor for the Regulation of Pulmonary Inflammation in Cystic Fibrosis. Frontiers in Immunology, 2019, 10, 2643.	2.2	11
46	Reduced expression of Tis7/IFRD1 protein in murine and human cystic fibrosis airway epithelial cell models homozygous for the F508del-CFTR mutation. Biochemical and Biophysical Research Communications, 2011, 411, 471-476.	1.0	9
47	Morphological analysis of the trachea and pattern of breathing in βENaC-Tg mice. Respiratory Physiology and Neurobiology, 2011, 178, 346-348.	0.7	6
48	TMEM16A/ANO1: Current Strategies and Novel Drug Approaches for Cystic Fibrosis. Cells, 2021, 10, 2867.	1.8	6
49	Opposite Expression of Hepatic and Pulmonary Corticosteroid-Binding Globulin in Cystic Fibrosis Patients. Frontiers in Pharmacology, 2018, 9, 545.	1.6	2
50	Weak anti-inflammatory effects of glucocorticoids on CF bronchial epithelial cells. Journal of Cystic Fibrosis, 2009, 8, S54.	0.3	0