

# James D Clark

## List of Publications by Year in Descending Order

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

47  
papers

4,597  
citations

27  
h-index

48  
g-index

48  
ext. papers

5,073  
ext. citations

9.2  
avg, IF

4.72  
L-index

#	Paper	IF	Citations
47	Demonstration of In Vitro to In Vivo Translation of a TYK2 Inhibitor That Shows Cross Species Potency Differences. <i>Scientific Reports</i> , <b>2020</b> , 10, 8974	4.9	4
46	Molecular and Cellular Responses to the TYK2/JAK1 Inhibitor PF-06700841 Reveal Reduction of Skin Inflammation in Plaque Psoriasis. <i>Journal of Investigative Dermatology</i> , <b>2020</b> , 140, 1546-1555.e4	4.3	21
45	Design and optimization of a series of 4-(3-azabicyclo[3.1.0]hexan-3-yl)pyrimidin-2-amines: Dual inhibitors of TYK2 and JAK1. <i>Bioorganic and Medicinal Chemistry</i> , <b>2020</b> , 28, 115481	3.4	3
44	Discovery of Tyrosine Kinase 2 (TYK2) Inhibitor (PF-06826647) for the Treatment of Autoimmune Diseases. <i>Journal of Medicinal Chemistry</i> , <b>2020</b> , 63, 13561-13577	8.3	19
43	Evaluation of the Short-, Mid-, and Long-Term Effects of Tofacitinib on Lymphocytes in Patients With Rheumatoid Arthritis. <i>Arthritis and Rheumatology</i> , <b>2019</b> , 71, 685-695	9.5	21
42	Reversibility of peripheral blood leukocyte phenotypic and functional changes after exposure to and withdrawal from tofacitinib, a Janus kinase inhibitor, in healthy volunteers. <i>Clinical Immunology</i> , <b>2018</b> , 191, 10-20	9	15
41	Identification of N-{cis-3-[Methyl(7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino]cyclobutyl}propane-1-sulfonamide (PF-04965842): A Selective JAK1 Clinical Candidate for the Treatment of Autoimmune Diseases. <i>Journal of Medicinal Chemistry</i> , <b>2018</b> , 61, 1438-1452	8.3	57
40	Dual Inhibition of TYK2 and JAK1 for the Treatment of Autoimmune Diseases: Discovery of ((S)-2,2-Difluorocyclopropyl)((1R,5S)-3-(2-((1-methyl-1H-pyrazol-4-yl)amino)pyrimidin-4-yl)-3,8-diazabicyclo[3.2.1]octan-8-yl)methanone (PF-06700841). <i>Journal of Medicinal Chemistry</i> , <b>2018</b> , 61, 8597-8612	8.3	54
39	Discovery of a JAK3-Selective Inhibitor: Functional Differentiation of JAK3-Selective Inhibition over pan-JAK or JAK1-Selective Inhibition. <i>ACS Chemical Biology</i> , <b>2016</b> , 11, 3442-3451	4.9	85
38	Tofacitinib attenuates pathologic immune pathways in patients with psoriasis: A randomized phase 2 study. <i>Journal of Allergy and Clinical Immunology</i> , <b>2016</b> , 137, 1079-1090	11.5	89
37	Parsing the Interferon Transcriptional Network and Its Disease Associations. <i>Cell</i> , <b>2016</b> , 164, 564-78	56.2	151
36	Network pharmacology of JAK inhibitors. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , <b>2016</b> , 113, 9852-7	11.5	44
35	The mechanism of action of tofacitinib - an oral Janus kinase inhibitor for the treatment of rheumatoid arthritis. <i>Clinical and Experimental Rheumatology</i> , <b>2016</b> , 34, 318-28	2.2	155
34	Discovery and development of Janus kinase (JAK) inhibitors for inflammatory diseases. <i>Journal of Medicinal Chemistry</i> , <b>2014</b> , 57, 5023-38	8.3	313
33	The balance of expression of PTPN22 splice forms is significantly different in rheumatoid arthritis patients compared with controls. <i>Genome Medicine</i> , <b>2012</b> , 4, 2	14.4	20
32	Variants within STAT genes reveal association with anticitrullinated protein antibody-negative rheumatoid arthritis in 2 European populations. <i>Journal of Rheumatology</i> , <b>2012</b> , 39, 1509-16	4.1	21
31	Preclinical evaluation of an inhibitor of cytosolic phospholipase A2 for the treatment of asthma. <i>Journal of Pharmacology and Experimental Therapeutics</i> , <b>2012</b> , 340, 656-65	4.7	26

30	The expression of splice forms for the rheumatoid arthritis risk associated gene PTPN22 is significantly different for patients compared to controls. <i>Annals of the Rheumatic Diseases</i> , <b>2012</b> , 71, A56.1-A56	2.4	
29	The cPLA2 inhibitor efipladib decreases nociceptive responses without affecting PGE2 levels in the cerebral spinal fluid. <i>Neuropharmacology</i> , <b>2011</b> , 60, 633-41	5.5	4
28	Cytosolic phospholipase A2 blockade abrogates disease during the tissue-damage effector phase of experimental autoimmune encephalomyelitis by its action on APCs. <i>Journal of Immunology</i> , <b>2011</b> , 187, 1986-97	5.3	10
27	Effect of perzinfotel and a proprietary phospholipase A(2) inhibitor on kinetic gait and subjective lameness scores in dogs with sodium urate-induced synovitis. <i>American Journal of Veterinary Research</i> , <b>2011</b> , 72, 757-63	1.1	7
26	A human CXCL13-induced actin polymerization assay measured by fluorescence plate reader. <i>Assay and Drug Development Technologies</i> , <b>2010</b> , 8, 73-84	2.1	5
25	Utility of cytosolic phospholipase A2 (cPLA2) inhibitors in the treatment of asthma. <i>Progress in Respiratory Research</i> , <b>2010</b> , 207-212		
24	Selective functional inhibition of JAK-3 is sufficient for efficacy in collagen-induced arthritis in mice. <i>Arthritis and Rheumatism</i> , <b>2010</b> , 62, 2283-93		68
23	Benzhydrylquinazolinediones: novel cytosolic phospholipase A2alpha inhibitors with improved physicochemical properties. <i>Bioorganic and Medicinal Chemistry</i> , <b>2009</b> , 17, 4383-405	3.4	26
22	Reactions of functionalized sulfonamides: application to lowering the lipophilicity of cytosolic phospholipase A2alpha inhibitors. <i>Journal of Medicinal Chemistry</i> , <b>2009</b> , 52, 1156-71	8.3	12
21	Blockade of cytosolic phospholipase A2 alpha prevents experimental autoimmune encephalomyelitis and diminishes development of Th1 and Th17 responses. <i>Journal of Neuroimmunology</i> , <b>2008</b> , 204, 29-37	3.5	53
20	Indole cytosolic phospholipase A2 alpha inhibitors: discovery and in vitro and in vivo characterization of 4-{3-[5-chloro-2-(2-[(3,4-dichlorobenzyl)sulfonyl]amino)ethyl]-1-(diphenylmethyl)-1H-indol-3-yl]propyl}benzoic acid, efipladib. <i>Journal of Medicinal Chemistry</i> , <b>2008</b> , 51, 3388-413	8.3	70
19	Benzenesulfonamide indole inhibitors of cytosolic phospholipase A2alpha: optimization of in vitro potency and rat pharmacokinetics for oral efficacy. <i>Bioorganic and Medicinal Chemistry</i> , <b>2008</b> , 16, 1345-58	3.4	29
18	Thermodynamic characterization of cytosolic phospholipase A2 alpha inhibitors. <i>Analytical Biochemistry</i> , <b>2008</b> , 383, 217-25	3.1	7
17	Pharmacologic inhibition of tpl2 blocks inflammatory responses in primary human monocytes, synoviocytes, and blood. <i>Journal of Biological Chemistry</i> , <b>2007</b> , 282, 33295-33304	5.4	66
16	Discovery of Ecopladib, an indole inhibitor of cytosolic phospholipase A2alpha. <i>Journal of Medicinal Chemistry</i> , <b>2007</b> , 50, 1380-400	8.3	70
15	Inhibition of cytosolic phospholipase A2alpha: hit to lead optimization. <i>Journal of Medicinal Chemistry</i> , <b>2006</b> , 49, 135-58	8.3	61
14	1,2,4-Oxadiazolidin-3,5-diones and 1,3,5-triazin-2,4,6-triones as cytosolic phospholipase A2alpha inhibitors. <i>Bioorganic and Medicinal Chemistry Letters</i> , <b>2006</b> , 16, 2978-81	2.9	17
13	A fluorescence-based assay for fatty acid amide hydrolase compatible with high-throughput screening. <i>Analytical Biochemistry</i> , <b>2005</b> , 343, 143-51	3.1	48

12	Cytosolic phospholipase A2 alpha-deficient mice are resistant to experimental autoimmune encephalomyelitis. <i>Journal of Experimental Medicine</i> , <b>2005</b> , 202, 841-51	16.6	120
11	Potential therapeutic uses of phospholipase A2 inhibitors. <i>Expert Opinion on Therapeutic Patents</i> , <b>2004</b> , 14, 937-950	6.8	27
10	Structure-activity relationships of indole cytosolic phospholipase A(2)alpha inhibitors: substrate mimetics. <i>Bioorganic and Medicinal Chemistry Letters</i> , <b>2003</b> , 13, 4501-4	2.9	20
9	Cytosolic phospholipase A2alpha-deficient mice are resistant to collagen-induced arthritis. <i>Journal of Experimental Medicine</i> , <b>2003</b> , 197, 1297-302	16.6	129
8	Crystal structure of human cytosolic phospholipase A2 reveals a novel topology and catalytic mechanism. <i>Cell</i> , <b>1999</b> , 97, 349-60	56.2	304
7	Trifluoromethyl ketones and methyl fluorophosphonates as inhibitors of group IV and VI phospholipases A(2): structure-function studies with vesicle, micelle, and membrane assays. <i>Biochimica Et Biophysica Acta - Biomembranes</i> , <b>1999</b> , 1420, 45-56	3.8	63
6	Characterization of Ca <sup>2+</sup> -dependent phospholipase A2 activity during zebrafish embryogenesis. <i>Journal of Biological Chemistry</i> , <b>1999</b> , 274, 19338-46	5.4	49
5	Solution structure and membrane interactions of the C2 domain of cytosolic phospholipase A2. <i>Journal of Molecular Biology</i> , <b>1998</b> , 280, 485-500	6.5	109
4	Independent folding and ligand specificity of the C2 calcium-dependent lipid binding domain of cytosolic phospholipase A2. <i>Journal of Biological Chemistry</i> , <b>1998</b> , 273, 1365-72	5.4	101
3	Cytosolic phospholipase A2. <i>Journal of Lipid Mediators and Cell Signalling</i> , <b>1995</b> , 12, 83-117		432
2	A novel arachidonic acid-selective cytosolic PLA2 contains a Ca(2+)-dependent translocation domain with homology to PKC and GAP. <i>Cell</i> , <b>1991</b> , 65, 1043-51	56.2	1552
1	Malate synthase: proof of a stepwise Claisen condensation using the double-isotope fractionation test. <i>Biochemistry</i> , <b>1988</b> , 27, 5961-71	3.2	40