## **Douglas S Jones**

List of Publications by Year in descending order

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DOLICIAS S IONES

#	Article	IF	CITATIONS
1	Cell surface–tethered IL-12 repolarizes the tumor immune microenvironment to enhance the efficacy of adoptive T cell therapy. Science Advances, 2022, 8, eabi8075.	10.3	21
2	A participant-derived xenograft model of HIV enables long-term evaluation of autologous immunotherapies. Journal of Experimental Medicine, 2021, 218, .	8.5	9
3	Inflammatory but not mitogenic contexts prime synovial fibroblasts for compensatory signaling responses to p38 inhibition. Science Signaling, 2018, 11, .	3.6	24
4	Structure-guided development of covalent TAK1 inhibitors. Bioorganic and Medicinal Chemistry, 2017, 25, 838-846.	3.0	28
5	Studies of TAK1-centered polypharmacology with novel covalent TAK1 inhibitors. Bioorganic and Medicinal Chemistry, 2017, 25, 1320-1328.	3.0	17
6	Profiling drugs for rheumatoid arthritis that inhibit synovial fibroblast activation. Nature Chemical Biology, 2017, 13, 38-45.	8.0	56
7	Inhibition of the CAS6/AXL pathway augments the efficacy of chemotherapies. Journal of Clinical Investigation, 2016, 127, 183-198.	8.2	86
8	An engineered dimeric fragment of hepatocyte growth factor is a potent câ€MET agonist. FEBS Letters, 2014, 588, 4831-4837.	2.8	23
9	An engineered Axl 'decoy receptor' effectively silences the Gas6-Axl signaling axis. Nature Chemical Biology, 2014, 10, 977-983.	8.0	117
10	Functional Mutation of Multiple Solvent-Exposed Loops in the Ecballium elaterium Trypsin Inhibitor-II Cystine Knot Miniprotein. PLoS ONE, 2011, 6, e16112.	2.5	37
11	Engineering hepatocyte growth factor fragments with high stability and activity as Met receptor agonists and antagonists. Proceedings of the National Academy of Sciences of the United States of America, 2011, 108, 13035-13040.	7.1	53
12	Developing therapeutic proteins by engineering ligand–receptor interactions. Trends in Biotechnology, 2008, 26, 498-505.	9.3	40