

# James C Geoghegan

## List of Publications by Year in descending order

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Version: 2024-02-01

19  
papers

2,868  
citations

430874

18  
h-index

794594

19  
g-index

19  
all docs

19  
docs citations

19  
times ranked

4446  
citing authors

#	ARTICLE	IF	CITATIONS
1	Broad and potent activity against SARS-like viruses by an engineered human monoclonal antibody. <i>Science</i> , 2021, 371, 823-829.	12.6	285
2	Prolonged evolution of the human B cell response to SARS-CoV-2 infection. <i>Science Immunology</i> , 2021, 6, .	11.9	153
3	Mitigation of reversible self-association and viscosity in a human IgG1 monoclonal antibody by rational, structure-guided Fv engineering. <i>MAbs</i> , 2016, 8, 941-950.	5.2	51
4	Inhibition of CD73 AMP hydrolysis by a therapeutic antibody with a dual, non-competitive mechanism of action. <i>MAbs</i> , 2016, 8, 454-467.	5.2	91
5	Chondroitin Sulfate is the Primary Receptor for a Peptide-Modified AAV That Targets Brain Vascular Endothelium In Vivo. <i>Molecular Therapy - Nucleic Acids</i> , 2014, 3, e202.	5.1	12
6	Sialic Acid Deposition Impairs the Utility of AAV9, but Not Peptide-modified AAVs for Brain Gene Therapy in a Mouse Model of Lysosomal Storage Disease. <i>Molecular Therapy</i> , 2012, 20, 1393-1399.	8.2	40
7	Gene Silencing Mediated by siRNA-binding Fusion Proteins Is Attenuated by Double-stranded RNA-binding Domain Structure. <i>Molecular Therapy - Nucleic Acids</i> , 2012, 1, e53.	5.1	28
8	Isolation of phosphatidylethanolamine as a solitary cofactor for prion formation in the absence of nucleic acids. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2012, 109, 8546-8551.	7.1	211
9	Dissociation of Infectivity from Seeding Ability in Prions with Alternate Docking Mechanism. <i>PLoS Pathogens</i> , 2011, 7, e1002128.	4.7	43
10	Species-Dependent Differences in Cofactor Utilization for Formation of the Protease-Resistant Prion Protein in Vitro. <i>Biochemistry</i> , 2010, 49, 3928-3934.	2.5	85
11	Trans-Dominant Inhibition of Prion Propagation In Vitro Is Not Mediated by an Accessory Cofactor. <i>PLoS Pathogens</i> , 2009, 5, e1000535.	4.7	52
12	Selective Incorporation of Polyanionic Molecules into Hamster Prions. <i>Journal of Biological Chemistry</i> , 2007, 282, 36341-36353.	3.4	100
13	Copper (II) ions potently inhibit purified PrPres amplification. <i>Journal of Neurochemistry</i> , 2006, 96, 1409-1415.	3.9	53
14	Protease-resistant Prion Protein Amplification Reconstituted with Partially Purified Substrates and Synthetic Polyanions. <i>Journal of Biological Chemistry</i> , 2005, 280, 26873-26879.	3.4	177
15	Sclerostin Inhibition of Wnt-3a-induced C3H10T1/2 Cell Differentiation Is Indirect and Mediated by Bone Morphogenetic Proteins. <i>Journal of Biological Chemistry</i> , 2005, 280, 2498-2502.	3.4	160
16	Noggin and Sclerostin Bone Morphogenetic Protein Antagonists Form a Mutually Inhibitory Complex. <i>Journal of Biological Chemistry</i> , 2004, 279, 36293-36298.	3.4	88
17	Unique regulation of SOST, the sclerosteosis gene, by BMPs and steroid hormones in human osteoblasts. <i>Bone</i> , 2004, 35, 448-454.	2.9	78
18	Sclerostin promotes the apoptosis of human osteoblastic cells: a novel regulation of bone formation. <i>Bone</i> , 2004, 35, 828-835.	2.9	198

#	ARTICLE	IF	CITATIONS
19	Osteocyte control of bone formation via sclerostin, a novel BMP antagonist. EMBO Journal, 2003, 22, 6267-6276.	7.8	963