

# Jaeok Park

## List of Publications by Year in descending order

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Version: 2024-02-01

23  
papers

489  
citations

623734

14  
h-index

713466

21  
g-index

23  
all docs

23  
docs citations

23  
times ranked

750  
citing authors

#	ARTICLE	IF	CITATIONS
1	Inhibition of farnesyl pyrophosphate (FPP) and/or geranylgeranyl pyrophosphate (GGPP) biosynthesis and its implication in the treatment of cancers. <i>Critical Reviews in Biochemistry and Molecular Biology</i> , 2019, 54, 41-60.	5.2	52
2	Design and Synthesis of Active Site Inhibitors of the Human Farnesyl Pyrophosphate Synthase: Apoptosis and Inhibition of ERK Phosphorylation in Multiple Myeloma Cells. <i>Journal of Medicinal Chemistry</i> , 2012, 55, 3201-3215.	6.4	46
3	Thienopyrimidine Bisphosphonate (ThPBP) Inhibitors of the Human Farnesyl Pyrophosphate Synthase: Optimization and Characterization of the Mode of Inhibition. <i>Journal of Medicinal Chemistry</i> , 2013, 56, 7939-7950.	6.4	43
4	Human isoprenoid synthase enzymes as therapeutic targets. <i>Frontiers in Chemistry</i> , 2014, 2, 50.	3.6	37
5	Unraveling the Prenylationâ€“Cancer Paradox in Multiple Myeloma with Novel Geranylgeranyl Pyrophosphate Synthase (GGPPS) Inhibitors. <i>Journal of Medicinal Chemistry</i> , 2018, 61, 6904-6917.	6.4	33
6	Human farnesyl pyrophosphate synthase is allosterically inhibited by its own product. <i>Nature Communications</i> , 2017, 8, 14132.	12.8	32
7	Maintenance of Native-like Protein Dynamics May Not Be Required for Engineering Functional Proteins. <i>Chemistry and Biology</i> , 2014, 21, 1330-1340.	6.0	29
8	Multistage Screening Reveals Chameleon Ligands of the Human Farnesyl Pyrophosphate Synthase: Implications to Drug Discovery for Neurodegenerative Diseases. <i>Journal of Medicinal Chemistry</i> , 2014, 57, 5764-5776.	6.4	29
9	Ternary complex structures of human farnesyl pyrophosphate synthase bound with a novel inhibitor and secondary ligands provide insights into the molecular details of the enzymeâ€™s active site closure. <i>BMC Structural Biology</i> , 2012, 12, 32.	2.3	21
10	Pharmacophore Mapping of Thienopyrimidine-Based Monophosphonate (ThP-MP) Inhibitors of the Human Farnesyl Pyrophosphate Synthase. <i>Journal of Medicinal Chemistry</i> , 2017, 60, 2119-2134.	6.4	21
11	A potential gain-of-function variant of SLC9A6 leads to endosomal alkalization and neuronal atrophy associated with Christianson Syndrome. <i>Neurobiology of Disease</i> , 2019, 121, 187-204.	4.4	21
12	Structural and functional insights into esterase-mediated macrolide resistance. <i>Nature Communications</i> , 2021, 12, 1732.	12.8	21
13	The Structural Dynamics of Engineered Î²-Lactamases Vary Broadly on Three Timescales yet Sustain Native Function. <i>Scientific Reports</i> , 2019, 9, 6656.	3.3	19
14	Phosphonate and Bisphosphonate Inhibitors of Farnesyl Pyrophosphate Synthases: A Structure-Guided Perspective. <i>Frontiers in Chemistry</i> , 2020, 8, 612728.	3.6	19
15	Probing the molecular and structural elements of ligands binding to the active site versus an allosteric pocket of the human farnesyl pyrophosphate synthase. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2015, 25, 1117-1123.	2.2	15
16	Assorted dysfunctions of endosomal alkali cation/proton exchanger SLC9A6 variants linked to Christianson syndrome. <i>Journal of Biological Chemistry</i> , 2020, 295, 7075-7095.	3.4	13
17	Chirality-Driven Mode of Binding of Î±-Aminophosphonic Acid-Based Allosteric Inhibitors of the Human Farnesyl Pyrophosphate Synthase (hFPPS). <i>Journal of Medicinal Chemistry</i> , 2019, 62, 9691-9702.	6.4	10
18	Structure of human farnesyl pyrophosphate synthase in complex with an aminopyridine bisphosphonate and two molecules of inorganic phosphate. <i>Acta Crystallographica Section F, Structural Biology Communications</i> , 2014, 70, 299-304.	0.8	9

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19	Structural and phylogenetic analyses of resistance to next-generation aminoglycosides conferred by AAC(2) enzymes. <i>Scientific Reports</i> , 2021, 11, 11614.	3.3	9
20	Crystallographic and thermodynamic characterization of phenylaminopyridine bisphosphonates binding to human farnesyl pyrophosphate synthase. <i>PLoS ONE</i> , 2017, 12, e0186447.	2.5	5
21	Synthesis and Evaluation of Structurally Diverse C-2-Substituted Thienopyrimidine-Based Inhibitors of the Human Geranylgeranyl Pyrophosphate Synthase. <i>Journal of Medicinal Chemistry</i> , 2022, 65, 2471-2496.	6.4	5
22	Crystal structures of hFPPS in complex with novel anticancer drug leads. <i>Acta Crystallographica Section A: Foundations and Advances</i> , 2014, 70, C712-C712.	0.1	0
23	Revisiting the Catalytic Cycle and Kinetic Mechanism of Aminoglycoside <i>O</i> -Nucleotidyltransferase(2): A Structural and Kinetic Study. <i>ACS Chemical Biology</i> , 2020, 15, 686-694.	3.4	0