

# Pei-Pei Kung

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/6799564/publications.pdf>

Version: 2024-02-01

12  
papers

568  
citations

840776

11  
h-index

1125743

13  
g-index

15  
all docs

15  
docs citations

15  
times ranked

909  
citing authors

#	ARTICLE	IF	CITATIONS
1	Polycomb repressive complex 2 structure with inhibitor reveals a mechanism of activation and drug resistance. <i>Nature Communications</i> , 2016, 7, 11384.	12.8	137
2	Optimization of Orally Bioavailable Enhancer of Zeste Homolog 2 (EZH2) Inhibitors Using Ligand and Property-Based Design Strategies: Identification of Development Candidate ( <i>R</i> )-5,8-Dichloro-7-(methoxy(oxetan-3-yl)methyl)-2-((4-methoxy-6-methyl-2-oxo-1,2-dihydropyridin-3-yl)methyl)-3,4-dihydroisoquinoline-1-carboxamide (PF-06821497). <i>Journal of Medicinal Chemistry</i> , 2018, 61, 650-665.	6.4	91
3	Dihydroxyphenylisoindoline Amides as Orally Bioavailable Inhibitors of the Heat Shock Protein 90 (Hsp90) Molecular Chaperone. <i>Journal of Medicinal Chemistry</i> , 2010, 53, 499-503.	6.4	64
4	Design and Synthesis of Pyridone-Containing 3,4-Dihydroisoquinoline-1(2 <i>H</i> )-ones as a Novel Class of Enhancer of Zeste Homolog 2 (EZH2) Inhibitors. <i>Journal of Medicinal Chemistry</i> , 2016, 59, 8306-8325.	6.4	53
5	Dihydroxyphenyl amides as inhibitors of the Hsp90 molecular chaperone. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2008, 18, 6273-6278.	2.2	46
6	Design strategies to target crystallographic waters applied to the Hsp90 molecular chaperone. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2011, 21, 3557-3562.	2.2	43
7	Optimization of Potent, Selective, and Orally Bioavailable Pyrrolidinopyrimidine-Containing Inhibitors of Heat Shock Protein 90. Identification of Development Candidate 2-Amino-4-[4-chloro-2-[2-(4-fluoro-1 <i>H</i> -pyrazol-1-yl)ethoxy]-6-methylphenyl]- <i>N</i> -(2,2-difluoropropyl)-5,7-dihydro-6 <i>H</i> -pyrido[4,3- <i>b</i> ]pyridine-3-carboxamide (PF-06821497). <i>Journal of Medicinal Chemistry</i> , 2011, 54, 3368-3385.	6.4	40
8	Characterization of Specific <i>N</i> -Acetyltransferase 50 (Naa50) Inhibitors Identified Using a DNA Encoded Library. <i>ACS Medicinal Chemistry Letters</i> , 2020, 11, 1175-1184.	2.8	27
9	Rapid, microwave-assisted synthesis of <i>N</i> 1-substituted 3-amino-1,2,4-triazoles. <i>Tetrahedron Letters</i> , 2009, 50, 1667-1670.	1.4	17
10	Design, synthesis, and biological evaluation of novel human 5- <i>deoxy</i> -5-methylthioadenosine phosphorylase (MTAP) substrates. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2005, 15, 2829-2833.	2.2	14
11	Design and Characterization of a Pyridone-Containing EZH2 Inhibitor Phosphate Prodrug. <i>Journal of Medicinal Chemistry</i> , 2021, 64, 1725-1732.	6.4	11
12	Harnessing Ionic Selectivity in Acetyltransferase Chemoproteomic Probes. <i>ACS Chemical Biology</i> , 2021, 16, 27-34.	3.4	5