

# Pei-Pei Kung

## List of Publications by Year in descending order

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12  
papers

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times ranked

909  
citing authors

| #  | ARTICLE   | IF   | CITATIONS |
|----|---|------|-----------|
| 1  | Design and Characterization of a Pyridone-Containing EZH2 Inhibitor Phosphate Prodrug. <i>Journal of Medicinal Chemistry</i> , 2021, 64, 1725-1732.   | 6.4  | 11        |
| 2  | Harnessing Ionic Selectivity in Acetyltransferase Chemoproteomic Probes. <i>ACS Chemical Biology</i> , 2021, 16, 27-34.   | 3.4  | 5         |
| 3  | 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        |
| 4  | Optimization of Orally Bioavailable Enhancer of Zeste Homolog 2 (EZH2) Inhibitors Using Ligand and Property-Based Design Strategies: Identification of Development Candidate (R)-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 (PF-06821497). <i>Journal of Medicinal Chemistry</i> , 2018, 61, 650-665.   | 6.4  | 91        |
| 5  | 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        |
| 6  | 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       |
| 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>d</i> ]pyrimidin-2(1 <i>H</i> )-one (PF-04961859). <i>Journal of Medicinal Chemistry</i> , 2011, 54, 3368-3385. | 6.4  | 40        |
| 8  | 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        |
| 9  | 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        |
| 10 | 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        |
| 11 | Dihydroxyphenyl amides as inhibitors of the Hsp90 molecular chaperone. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2008, 18, 6273-6278.   | 2.2  | 46        |
| 12 | Design, synthesis, and biological evaluation of novel human 5'-deoxy-5'-methylthioadenosine phosphorylase (MTAP) substrates. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2005, 15, 2829-2833.   | 2.2  | 14        |