

Andreas L Marzinzik

List of Publications by Year in descending order

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9
papers

1,148
citations

1307594

7
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1474206

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docs citations

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times ranked

1507
citing authors

#	ARTICLE	IF	CITATIONS
1	Discovery, X-ray structure and CPP-conjugation enabled uptake of p53/MDM2 macrocyclic peptide inhibitors. <i>RSC Chemical Biology</i> , 2021, 2, 1661-1668.	4.1	7
2	Fragment-Based Discovery of Non-Bisphosphonate Binders of <i>Trypanosoma brucei</i> Farnesyl Pyrophosphate Synthase. <i>ChemBioChem</i> , 2020, 21, 3096-3111.	2.6	8
3	Discovery of Potent and Selective Antibody-Drug Conjugates with Eg5 Inhibitors through Linker and Payload Optimization. <i>ACS Medicinal Chemistry Letters</i> , 2019, 10, 1674-1679.	2.8	10
4	Discovery of Asciminib (ABL001), an Allosteric Inhibitor of the Tyrosine Kinase Activity of BCR-ABL1. <i>Journal of Medicinal Chemistry</i> , 2018, 61, 8120-8135.	6.4	275
5	The allosteric inhibitor ABL001 enables dual targeting of BCR-ABL1. <i>Nature</i> , 2017, 543, 733-737.	27.8	389
6	Inhibitors of the Abl kinase directed at either the ATP- or myristate-binding site. <i>Biochimica Et Biophysica Acta - Proteins and Proteomics</i> , 2010, 1804, 454-462.	2.3	59
7	Allosteric non-bisphosphonate FPPS inhibitors identified by fragment-based discovery. <i>Nature Chemical Biology</i> , 2010, 6, 660-666.	8.0	110
8	Binding or Bending: Distinction of Allosteric Abl Kinase Agonists from Antagonists by an NMR-Based Conformational Assay. <i>Journal of the American Chemical Society</i> , 2010, 132, 7043-7048.	13.7	95
9	Library Design for Fragment Based Screening. <i>Current Topics in Medicinal Chemistry</i> , 2005, 5, 751-762.	2.1	195