Ernst K Schonbrunn

List of Publications by Year
in descending order

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37 papers	1,593 citations	393982 19 h-index	329751 37 g-index
38 all docs	38 docs citations	38 times ranked	2687 citing authors

#	Article	IF	Citations
1	Discovery of Dual TAF1-ATR Inhibitors and Ligand-Induced Structural Changes of the TAF1 Tandem Bromodomain. Journal of Medicinal Chemistry, 2022, 65, 4182-4200.	2.9	10
2	Dihydropyridine Lactam Analogs Targeting BET Bromodomains. ChemMedChem, 2022, 17, e202100407.	1.6	1
3	Tetrahydroindazole inhibitors of CDK2/cyclin complexes. European Journal of Medicinal Chemistry, 2021, 214, 113232.	2.6	5
4	Development of Dimethylisoxazole-Attached Imidazo[1,2-a]pyridines as Potent and Selective CBP/P300 Inhibitors. Journal of Medicinal Chemistry, 2021, 64, 5787-5801.	2.9	15
5	New Design Rules for Developing Potent Cell-Active Inhibitors of the Nucleosome Remodeling Factor (NURF) via BPTF Bromodomain Inhibition. Journal of Medicinal Chemistry, 2021, 64, 13902-13917.	2.9	14
6	Synthesis and structural characterization of a monocarboxylic inhibitor for GRB2 SH2 domain. Bioorganic and Medicinal Chemistry Letters, 2021, 51, 128354.	1.0	5
7	Structural Insights into JAK2 Inhibition by Ruxolitinib, Fedratinib, and Derivatives Thereof. Journal of Medicinal Chemistry, 2021, 64, 2228-2241.	2.9	49
8	Differential BET Bromodomain Inhibition by Dihydropteridinone and Pyrimidodiazepinone Kinase Inhibitors. Journal of Medicinal Chemistry, 2021, 64, 15772-15786.	2.9	10
9	NMR Analyses of Acetylated H2A.Z Isoforms Identify Differential Binding Interactions with the Bromodomain of the NURF Nucleosome Remodeling Complex. Biochemistry, 2020, 59, 1871-1880.	1.2	11
10	New inhibitors for the BPTF bromodomain enabled by structural biology and biophysical assay development. Organic and Biomolecular Chemistry, 2020, 18, 5174-5182.	1.5	14
11	Development of WEE2 kinase inhibitors as novel non-hormonal female contraceptives that target meiosisâ€. Biology of Reproduction, 2020, 103, 368-377.	1.2	7
12	Structural Basis of Inhibitor Selectivity in the BRD7/9 Subfamily of Bromodomains. Journal of Medicinal Chemistry, 2020, 63, 3227-3237.	2.9	19
13	Identification and Screening of Selective WEE2 Inhibitors to Develop Nonâ€Hormonal Contraceptives that Specifically Target Meiosis. ChemistrySelect, 2019, 4, 13363-13369.	0.7	7
14	Ligand-mediated protein degradation reveals functional conservation among sequence variants of the CUL4-type E3 ligase substrate receptor cereblon. Journal of Biological Chemistry, 2018, 293, 6187-6200.	1.6	32
15	Structural Basis of ALDH1A2 Inhibition by Irreversible and Reversible Small Molecule Inhibitors. ACS Chemical Biology, 2018, 13, 582-590.	1.6	48
16	Molecular Basis for the N-Terminal Bromodomain-and-Extra-Terminal-Family Selectivity of a Dual Kinaseâ€“Bromodomain Inhibitor. Journal of Medicinal Chemistry, 2018, 61, 9316-9334.	2.9	56
17	Design, Synthesis, and Characterization of a Fluorescence Polarization Pan-BET Bromodomain Probe. ACS Medicinal Chemistry Letters, 2018, 9, 1223-1229.	1.3	8
18	BET Bromodomain Inhibitors with One-Step Synthesis Discovered from Virtual Screen. Journal of Medicinal Chemistry, 2017, 60, 4805-4817.	2.9	39

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19	Dual Targeting of WEE1 and PLK1 by AZD1775 Elicits Single Agent Cellular Anticancer Activity. ACS Chemical Biology, 2017, 12, 1883-1892.	1.6	57
20	Potent Dual BET Bromodomain-Kinase Inhibitors as Value-Added Multitargeted Chemical Probes and Cancer Therapeutics. Molecular Cancer Therapeutics, 2017, 16, 1054-1067.	1.9	40
21	Structureâ€“Activity Studies of Nâ€Butylâ€1â€deoxynojirimycin (NBâ€DNJ) Analogues: Discovery of Potent and Selective Aminocyclopentitol Inhibitors of GBA1 and GBA2. ChemMedChem, 2017, 12, 1977-1984.	1.6	13
22	Advances of small molecule targeting of kinases. Current Opinion in Chemical Biology, 2017, 39, 126-132.	2.8	44
23	Structural Basis of Wee Kinases Functionality and Inactivation by Diverse Small Molecule Inhibitors. Journal of Medicinal Chemistry, 2017, 60, 7863-7875.	2.9	68
24	Discovery of Diverse Smallâ€Molecule Inhibitors of Mammalian Sterile20â€like Kinaseâ€…3 (MST3). ChemMedChem, 2016, 11, 1137-1144.	1.6	22
25	An Advanced Tool To Interrogate BRD9. Journal of Medicinal Chemistry, 2016, 59, 4459-4461.	2.9	13
26	Stability of the Human Hsp90-p50Cdc37 Chaperone Complex against Nucleotides and Hsp90 Inhibitors, and the Influence of Phosphorylation by Casein Kinase 2. Molecules, 2015, 20, 1643-1660.	1.7	12
27	Fluorinated Aromatic Amino Acids Are Sensitive 19F NMR Probes for Bromodomain-Ligand Interactions. ACS Chemical Biology, 2014, 9, 2755-2760.	1.6	79
28	Differential antibacterial properties of the MurA inhibitors terreic acid and fosfomycin. Journal of Basic Microbiology, 2014, 54, 322-326.	1.8	22
29	Acetyl-lysine Binding Site of Bromodomain-Containing Protein 4 (BRD4) Interacts with Diverse Kinase Inhibitors. ACS Chemical Biology, 2014, 9, 1160-1171.	1.6	188
30	Development of Highly Potent and Selective Diaminothiazole Inhibitors of Cyclin-Dependent Kinases. Journal of Medicinal Chemistry, 2013, 56, 3768-3782.	2.9	73
31	Cyclin-Dependent Kinase Inhibitor Dinaciclib Interacts with the Acetyl-Lysine Recognition Site of Bromodomains. ACS Chemical Biology, 2013, 8, 2360-2365.	1.6	132
32	Synthesis and Evaluation of Eight- and Four-Membered Iminosugar Analogues as Inhibitors of Testicular Ceramide-Specific Glucosyltransferase, Testicular Î²-Glucosidase 2, and Other Glycosidases. Journal of Organic Chemistry, 2012, 77, 3082-3098.	1.7	38
33	A Novel Approach to the Discovery of Smallâ€Molecule Ligands of CDK2. ChemBioChem, 2012, 13, 2128-2136.	1.3	65
34	A Novel Mechanism by Which Small Molecule Inhibitors Induce the DFG Flip in Aurora A. ACS Chemical Biology, 2012, 7, 698-706.	1.6	58
35	Discovery of a Potential Allosteric Ligand Binding Site in CDK2. ACS Chemical Biology, 2011, 6, 492-501.	1.6	151
36	The Fungal Product Terreic Acid Is a Covalent Inhibitor of the Bacterial Cell Wall Biosynthetic Enzyme UDP-N-Acetylglucosamine 1-Carboxyvinyltransferase (MurA),. Biochemistry, 2010, 49, 4276-4282.	1.2	50

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37	Evidence That the Fosfomycin Target Cys115 in UDP-N-acetylglucosamine Enolpyruvyl Transferase (MurA) Is Essential for Product Release. Journal of Biological Chemistry, 2005, 280, 3757-3763.	1.6	117