## **Eunyoung Park**

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/295675/publications.pdf

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687363 940533 1,547 16 13 16 citations h-index g-index papers 17 17 17 2404 docs citations times ranked citing authors all docs

#	Article	IF	CITATIONS
1	Allosteric MEK inhibitors act on BRAF/MEK complexes to block MEK activation. Proceedings of the National Academy of Sciences of the United States of America, 2021, 118, .	7.1	23
2	The structural basis of PTEN regulation by multi-site phosphorylation. Nature Structural and Molecular Biology, 2021, 28, 858-868.	8.2	20
3	Mutantâ€Selective Allosteric EGFR Degraders are Effective Against a Broad Range of Drugâ€Resistant Mutations. Angewandte Chemie - International Edition, 2020, 59, 14481-14489.	13.8	75
4	Mutantâ€Selective Allosteric EGFR Degraders are Effective Against a Broad Range of Drugâ€Resistant Mutations. Angewandte Chemie, 2020, 132, 14589-14597.	2.0	13
5	The Eya1 Phosphatase Mediates Shh-Driven Symmetric Cell Division of Cerebellar Granule Cell Precursors. Developmental Neuroscience, 2020, 42, 170-186.	2.0	10
6	Discovery and Optimization of Dibenzodiazepinones as Allosteric Mutant-Selective EGFR Inhibitors. ACS Medicinal Chemistry Letters, 2019, 10, 1549-1553.	2.8	47
7	Single and Dual Targeting of Mutant EGFR with an Allosteric Inhibitor. Cancer Discovery, 2019, 9, 926-943.	9.4	220
8	Architecture of autoinhibited and active BRAF–MEK1–14-3-3 complexes. Nature, 2019, 575, 545-550.	27.8	197
9	Discovery of a Highly Potent and Broadly Effective Epidermal Growth Factor Receptor and HER2 Exon 20 Insertion Mutant Inhibitor. Angewandte Chemie - International Edition, 2018, 57, 11629-11633.	13.8	20
10	Discovery of a Highly Potent and Broadly Effective Epidermal Growth Factor Receptor and HER2 Exon 20 Insertion Mutant Inhibitor. Angewandte Chemie, 2018, 130, 11803-11807.	2.0	4
11	Leveraging Gas-Phase Fragmentation Pathways for Improved Identification and Selective Detection of Targets Modified by Covalent Probes. Analytical Chemistry, 2016, 88, 12248-12254.	6.5	31
12	Overcoming EGFR(T790M) and EGFR(C797S) resistance with mutant-selective allosteric inhibitors. Nature, 2016, 534, 129-132.	27.8	637
13	SPLINTS: small-molecule protein ligand interface stabilizers. Current Opinion in Structural Biology, 2016, 37, 115-122.	5.7	37
14	Structure of a Bud6/Actin Complex Reveals a Novel WH2-like Actin Monomer Recruitment Motif. Structure, 2015, 23, 1492-1499.	3.3	16
15	Structure and mechanism of activity-based inhibition of the EGF receptor by Mig6. Nature Structural and Molecular Biology, 2015, 22, 703-711.	8.2	72
16	PARP1-Driven Poly-ADP-Ribosylation Regulates BRCA1 Function in Homologous Recombination–Mediated DNA Repair. Cancer Discovery, 2014, 4, 1430-1447.	9.4	125