

Kathryn M Ferguson

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/6066826/publications.pdf>

Version: 2024-02-01

21
papers

4,597
citations

516710

16
h-index

839539

18
g-index

22
all docs

22
docs citations

22
times ranked

5791
citing authors

#	ARTICLE	IF	CITATIONS
1	Structural basis for inhibition of the epidermal growth factor receptor by cetuximab. <i>Cancer Cell</i> , 2005, 7, 301-311.	16.8	949
2	An Open-and-Shut Case? Recent Insights into the Activation of EGF/ErbB Receptors. <i>Molecular Cell</i> , 2003, 12, 541-552.	9.7	774
3	EGF Activates Its Receptor by Removing Interactions that Autoinhibit Ectodomain Dimerization. <i>Molecular Cell</i> , 2003, 11, 507-517.	9.7	675
4	Signal-dependent membrane targeting by pleckstrin homology (PH) domains. <i>Biochemical Journal</i> , 2000, 350, 1-18.	3.7	656
5	The EGFR Family: Not So Prototypical Receptor Tyrosine Kinases. <i>Cold Spring Harbor Perspectives in Biology</i> , 2014, 6, a020768-a020768.	5.5	345
6	Structure-Based View of Epidermal Growth Factor Receptor Regulation. <i>Annual Review of Biophysics</i> , 2008, 37, 353-373.	10.0	306
7	EGFR Ligands Differentially Stabilize Receptor Dimers to Specify Signaling Kinetics. <i>Cell</i> , 2017, 171, 683-695.e18.	28.9	276
8	Epidermal Growth Factor Receptor Dimerization and Activation Require Ligand-Induced Conformational Changes in the Dimer Interface. <i>Molecular and Cellular Biology</i> , 2005, 25, 7734-7742.	2.3	247
9	Complex Relationship between Ligand Binding and Dimerization in the Epidermal Growth Factor Receptor. <i>Cell Reports</i> , 2014, 9, 1306-1317.	6.4	78
10	Scratching the surface with the PH domain. <i>Nature Structural and Molecular Biology</i> , 1995, 2, 715-718.	8.2	59
11	Molecular Basis for Necitumumab Inhibition of EGFR Variants Associated with Acquired Cetuximab Resistance. <i>Molecular Cancer Therapeutics</i> , 2018, 17, 521-531.	4.1	45
12	Comparison of <i>Saccharomyces cerevisiae</i> F-BAR Domain Structures Reveals a Conserved Inositol Phosphate Binding Site. <i>Structure</i> , 2015, 23, 352-363.	3.3	40
13	Glioblastoma mutations alter EGFR dimer structure to prevent ligand bias. <i>Nature</i> , 2022, 602, 518-522.	27.8	36
14	Insulin and epidermal growth factor receptor family members share parallel activation mechanisms. <i>Protein Science</i> , 2020, 29, 1331-1344.	7.6	31
15	Dimerization of Tie2 mediated by its membrane-proximal FNIII domains. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2017, 114, 4382-4387.	7.1	29
16	Molecular determinants of KA1 domain-mediated autoinhibition and phospholipid activation of MARK1 kinase. <i>Biochemical Journal</i> , 2017, 474, 385-398.	3.7	21
17	Intramolecular autoinhibition of checkpoint kinase 1 is mediated by conserved basic motifs of the C-terminal kinase-associated 1 domain. <i>Journal of Biological Chemistry</i> , 2017, 292, 19024-19033.	3.4	15
18	Structural Basis for MARK1 Kinase Autoinhibition by Its KA1 Domain. <i>Structure</i> , 2018, 26, 1137-1143.e3.	3.3	15

#	ARTICLE	IF	CITATIONS
19	Discoidin Discoveries. Structure, 2012, 20, 568-570.	3.3	0
20	Structural aspects of extracellular EGFR signaling. FASEB Journal, 2009, 23, 198.3.	0.5	0
21	The Mechanism of Ligand-Induced Activation of the Tie Family of Receptor Tyrosine Kinases. FASEB Journal, 2019, 33, 809.10.	0.5	0