

# Igor Kurinov

## List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/225835/publications.pdf>

Version: 2024-02-01

24  
papers

1,016  
citations

516710

16  
h-index

642732

23  
g-index

26  
all docs

26  
docs citations

26  
times ranked

1748  
citing authors

#	ARTICLE	IF	CITATIONS
1	Structural mechanism of ligand activation in human calcium-sensing receptor. <i>ELife</i> , 2016, 5, .	6.0	189
2	Structures of CRISPR Cas3 offer mechanistic insights into Cascade-activated DNA unwinding and degradation. <i>Nature Structural and Molecular Biology</i> , 2014, 21, 771-777.	8.2	167
3	Mechanism of Polyubiquitination by Human Anaphase-Promoting Complex: RING Repurposing for Ubiquitin Chain Assembly. <i>Molecular Cell</i> , 2014, 56, 246-260.	9.7	98
4	Functional characterization of a PROTAC directed against BRAF mutant V600E. <i>Nature Chemical Biology</i> , 2020, 16, 1170-1178.	8.0	80
5	Inhibition of SCF ubiquitin ligases by engineered ubiquitin variants that target the Cul1 binding site on the Skp1â€F-box interface. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2016, 113, 3527-3532.	7.1	61
6	Crystal Structure of Clustered Regularly Interspaced Short Palindromic Repeats (CRISPR)-associated Csn2 Protein Revealed Ca <sup>2+</sup> -dependent Double-stranded DNA Binding Activity*. <i>Journal of Biological Chemistry</i> , 2011, 286, 30759-30768.	3.4	49
7	Higher-Order Assembly of BRCC36â€KIAA0157 Is Required for DUB Activity and Biological Function. <i>Molecular Cell</i> , 2015, 59, 970-983.	9.7	44
8	Structural basis for the recruitment of glycogen synthase by glycogenin. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2014, 111, E2831-40.	7.1	43
9	Structural basis for auxiliary subunit KCTD16 regulation of the GABA <sub>B</sub> receptor. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2019, 116, 8370-8379.	7.1	32
10	Crystal structure of the human Políu B-subunit in complex with the C-terminal domain of the catalytic subunit. <i>Journal of Biological Chemistry</i> , 2017, 292, 15717-15730.	3.4	30
11	Structural Insights into the Induced-fit Inhibition of Fascin by a Small-Molecule Inhibitor. <i>Journal of Molecular Biology</i> , 2018, 430, 1324-1335.	4.2	28
12	A Structure-Based Strategy for Engineering Selective Ubiquitin Variant Inhibitors of Skp1-Cul1-F-Box Ubiquitin Ligases. <i>Structure</i> , 2018, 26, 1226-1236.e3.	3.3	27
13	Structure of human ADP-ribosyl-acceptor hydrolase 3 bound to ADP-ribose reveals a conformational switch that enables specific substrate recognition. <i>Journal of Biological Chemistry</i> , 2018, 293, 12350-12359.	3.4	27
14	Structural and functional characterization of KEOPS dimerization by Pcc1 and its role in t <sup>sup</sup> 6 <sub>A</sub> biosynthesis. <i>Nucleic Acids Research</i> , 2016, 44, 6971-6980.	14.5	26
15	Structural and Functional Analysis of Ubiquitin-based Inhibitors That Target the Backsides of E2 Enzymes. <i>Journal of Molecular Biology</i> , 2020, 432, 952-966.	4.2	22
16	Conformation-specific inhibitors of activated Ras GTPases reveal limited Ras dependency of patient-derived cancer organoids. <i>Journal of Biological Chemistry</i> , 2020, 295, 4526-4540.	3.4	19
17	Crystal structure of human PACRG in complex with MEIG1 reveals roles in axoneme formation and tubulin binding. <i>Structure</i> , 2021, 29, 572-586.e6.	3.3	19
18	Structural Basis for Auto-Inhibition of the NDR1 Kinase Domain by an Atypically Long Activation Segment. <i>Structure</i> , 2018, 26, 1101-1115.e6.	3.3	17

#	ARTICLE	IF	CITATIONS
19	Rigidification Dramatically Improves Inhibitor Selectivity for RAF Kinases. ACS Medicinal Chemistry Letters, 2019, 10, 1074-1080.	2.8	10
20	FAM105A/OTULINL Is a Pseudodeubiquitinase of the OTU-Class that Localizes to the ER Membrane. Structure, 2019, 27, 1000-1012.e6.	3.3	10
21	Engineered SH2 Domains for Targeted Phosphoproteomics. ACS Chemical Biology, 0, , .	3.4	6
22	Panel of Engineered Ubiquitin Variants Targeting the Family of Human Ubiquitin Interacting Motifs. ACS Chemical Biology, 2022, 17, 941-956.	3.4	5
23	Bipartite binding of the N terminus of Skp2 to cyclin A. Structure, 2021, 29, 975-988.e5.	3.3	2
24	Structural and biochemical analysis of human ADP-ribosyl-acceptor hydrolase 3 reveals the basis of metal selectivity and different roles for the two magnesium ions. Journal of Biological Chemistry, 2021, 296, 100692.	3.4	1