

List of Publications by Citations

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

54 papers	990 citations	17 h-index	30 g-index
62 ext. papers	1,201 ext. citations	6.2 avg, IF	4.06 L-index

#	Paper	IF	Citations
54	Structure of human urokinase plasminogen activator in complex with its receptor. <i>Science</i> , <b>2006</b> , 311, 656-9	33.3	242
53	Quercetin-3-rutinoside Inhibits Protein Disulfide Isomerase by Binding to Its bW Domain. <i>Journal of Biological Chemistry</i> , <b>2015</b> , 290, 23543-52	5.4	57
52	Be Active or Not: the Relative Contribution of Active and Passive Tumor Targeting of Nanomaterials. <i>Nanotheranostics</i> , <b>2017</b> , 1, 346-357	5.6	56
51	A novel tumor targeting drug carrier for optical imaging and therapy. <i>Theranostics</i> , <b>2014</b> , 4, 642-59	12.1	54
50	Structural basis of specificity of a peptidyl urokinase inhibitor, upain-1. <i>Journal of Structural Biology</i> , <b>2007</b> , 160, 1-10	3.4	45
49	Structure of catalytic domain of Matriptase in complex with Sunflower trypsin inhibitor-1. <i>BMC Structural Biology</i> , <b>2011</b> , 11, 30	2.7	44
48	Re-engineering the Immune Response to Metastatic Cancer: Antibody-Recruiting Small Molecules Targeting the Urokinase Receptor. <i>Angewandte Chemie - International Edition</i> , <b>2016</b> , 55, 3642-6	16.4	43
47	Crystal structure of the urokinase receptor in a ligand-free form. <i>Journal of Molecular Biology</i> , <b>2012</b> , 416, 629-41	6.5	41
46	Crystal Structure of the Michaelis Complex between Tissue-type Plasminogen Activator and Plasminogen Activators Inhibitor-1. <i>Journal of Biological Chemistry</i> , <b>2015</b> , 290, 25795-804	5.4	28
45	Crystal structures of matriptase in complex with its inhibitor hepatocyte growth factor activator inhibitor-1. <i>Journal of Biological Chemistry</i> , <b>2013</b> , 288, 11155-64	5.4	28
44	A structural mechanism of flavonoids in inhibiting serine proteases. <i>Food and Function</i> , <b>2017</b> , 8, 2437-2443	4.31	27
43	Stabilizing a flexible interdomain hinge region harboring the SMB binding site drives uPAR into its closed conformation. <i>Journal of Molecular Biology</i> , <b>2015</b> , 427, 1389-1403	6.5	22
42	A long-acting PAI-1 inhibitor reduces thrombus formation. <i>Thrombosis and Haemostasis</i> , <b>2017</b> , 117, 1338-1347	13.47	20
41	Rezymogenation of active urokinase induced by an inhibitory antibody. <i>Biochemical Journal</i> , <b>2013</b> , 449, 161-6	3.8	20
40	A specific plasminogen activator inhibitor-1 antagonist derived from inactivated urokinase. <i>Journal of Cellular and Molecular Medicine</i> , <b>2016</b> , 20, 1851-60	5.6	20
39	Nanoparticle Binding to Urokinase Receptor on Cancer Cell Surface Triggers Nanoparticle Disintegration and Cargo Release. <i>Theranostics</i> , <b>2019</b> , 9, 884-899	12.1	17
38	Dissociation of zinc phthalocyanine aggregation on bacterial surface is key for photodynamic antimicrobial effect. <i>Journal of Porphyrins and Phthalocyanines</i> , <b>2018</b> , 22, 925-934	1.8	17

37	Re-engineering the Immune Response to Metastatic Cancer: Antibody-Recruiting Small Molecules Targeting the Urokinase Receptor. <i>Angewandte Chemie</i> , <b>2016</b> , 128, 3706-3710	3.6	15
36	Serum Levels of Soluble Platelet Endothelial Cell Adhesion Molecule 1 in COVID-19 Patients Are Associated With Disease Severity. <i>Journal of Infectious Diseases</i> , <b>2021</b> , 223, 178-179	7	13
35	A drug carrier targeting murine uPAR for photodynamic therapy and tumor imaging. <i>Acta Biomaterialia</i> , <b>2015</b> , 23, 116-126	10.8	12
34	Development of inhibitors for uPAR: blocking the interaction of uPAR with its partners. <i>Drug Discovery Today</i> , <b>2021</b> , 26, 1076-1085	8.8	12
33	Specifically targeting cancer proliferation and metastasis processes: the development of matriptase inhibitors. <i>Cancer and Metastasis Reviews</i> , <b>2019</b> , 38, 507-524	9.6	11
32	Protein expression and preliminary crystallographic analysis of amino-terminal fragment of urokinase-type plasminogen activator. <i>Protein Expression and Purification</i> , <b>2006</b> , 49, 71-7	2	11
31	Structural basis of sequence-specific Holliday junction cleavage by MOC1. <i>Nature Chemical Biology</i> , <b>2019</b> , 15, 1241-1248	11.7	10
30	Crystal structures of the ligand-binding region of uPARAP: effect of calcium ion binding. <i>Biochemical Journal</i> , <b>2016</b> , 473, 2359-68	3.8	10
29	Mapping the topographic epitope landscape on the urokinase plasminogen activator receptor (uPAR) by surface plasmon resonance and X-ray crystallography. <i>Data in Brief</i> , <b>2015</b> , 5, 107-13	1.2	9
28	A novel purification procedure for recombinant human serum albumin expressed in <i>Pichia pastoris</i> . <i>Protein Expression and Purification</i> , <b>2018</b> , 149, 37-42	2	8
27	Dimer conformation of soluble PECAM-1, an endothelial marker. <i>International Journal of Biochemistry and Cell Biology</i> , <b>2016</b> , 77, 102-108	5.6	8
26	Plasma levels of the active form of suPAR are associated with COVID-19 severity. <i>Critical Care</i> , <b>2020</b> , 24, 704	10.8	8
25	The crystal structure of a multidomain protease inhibitor (HAI-1) reveals the mechanism of its auto-inhibition. <i>Journal of Biological Chemistry</i> , <b>2017</b> , 292, 8412-8423	5.4	7
24	Halogen bonding for the design of inhibitors by targeting the S1 pocket of serine proteases.. <i>RSC Advances</i> , <b>2018</b> , 8, 28189-28197	3.7	7
23	A novel ELISA for the detection of active form of plasminogen activator inhibitor-1 based on a highly specific trapping agent. <i>Analytica Chimica Acta</i> , <b>2019</b> , 1053, 98-104	6.6	6
22	Molecular basis of rutin inhibition of protein disulfide isomerase (PDI) by combined and experimental methods.. <i>RSC Advances</i> , <b>2018</b> , 8, 18480-18491	3.7	6
21	Enhanced Antitumor Efficacy and Imaging Application of Photosensitizer-Formulated Paclitaxel. <i>ACS Applied Materials &amp; Interfaces</i> , <b>2020</b> , 12, 4221-4230	9.5	6
20	An ELISA method detecting the active form of suPAR. <i>Talanta</i> , <b>2016</b> , 160, 205-210	6.2	6

19	A general strategy to inhibit serine protease by targeting its autolysis loop. <i>FASEB Journal</i> , <b>2021</b> , 35, e21259	0.9	6
18	Improved therapeutic efficacy of quercetin-loaded polymeric nanoparticles on triple-negative breast cancer by inhibiting uPA.. <i>RSC Advances</i> , <b>2020</b> , 10, 34517-34526	3.7	5
17	Crystal structure of the unoccupied murine urokinase-type plasminogen activator receptor (uPAR) reveals a tightly packed DII-DIII unit. <i>FEBS Letters</i> , <b>2019</b> , 593, 1236-1247	3.8	4
16	Inhibition of the Citrus Canker Pathogen Using a Photosensitizer Assisted by Sunlight Irradiation. <i>Frontiers in Microbiology</i> , <b>2020</b> , 11, 571691	5.7	4
15	Crystal Structures of Human C4.4A Reveal the Unique Association of Ly6/uPAR/Eheurotoxin Domain. <i>International Journal of Biological Sciences</i> , <b>2020</b> , 16, 981-993	11.2	3
14	Structural basis of specific inhibition of tissue-type plasminogen activator by plasminogen activators inhibitor-1. <i>Data in Brief</i> , <b>2016</b> , 6, 550-5	1.2	2
13	Solution Structure of SpoIVB Reveals Mechanism of PDZ Domain-Regulated Protease Activity. <i>Frontiers in Microbiology</i> , <b>2019</b> , 10, 1232	5.7	2
12	Expression and crystallographic studies of the ligand-binding region of the human endocytic collagen receptor uPARAP. <i>Acta Crystallographica Section F, Structural Biology Communications</i> , <b>2015</b> , 71, 1442-7	1.1	2
11	tPA Point Mutation at Autolysis Loop Enhances Resistance to PAI-1 Inhibition and Catalytic Activity. <i>Thrombosis and Haemostasis</i> , <b>2019</b> , 119, 77-86	7	2
10	Potent inhibition of Severe Acute Respiratory Syndrome Coronavirus 2 by photosensitizers compounds. <i>Dyes and Pigments</i> , <b>2021</b> , 194, 109570	4.6	2
9	Crystal structure and cellular functions of uPAR dimer.. <i>Nature Communications</i> , <b>2022</b> , 13, 1665	17.4	2
8	Recombinant hepatocyte growth factor activator inhibitor 1: expression in Drosophila S2 cells, purification and crystallization. <i>Acta Crystallographica Section F, Structural Biology Communications</i> , <b>2017</b> , 73, 45-50	1.1	1
7	Expression and crystallographic studies of the D1D2 domains of C4.4A, a homologous protein to the urokinase receptor. <i>Acta Crystallographica Section F, Structural Biology Communications</i> , <b>2017</b> , 73, 486-490	1.1	1
6	Vascular thiol isomerases: Structures, regulatory mechanisms, and inhibitor development. <i>Drug Discovery Today</i> , <b>2021</b> , 27, 626-626	8.8	1
5	A supramolecular nanocarrier for efficient cancer imaging and therapy by targeting at matriptase. <i>Journal of Controlled Release</i> , <b>2021</b> , 334, 153-163	11.7	1
4	Development of a Potent Antimicrobial Peptide With Photodynamic Activity. <i>Frontiers in Microbiology</i> , <b>2021</b> , 12, 624465	5.7	1
3	A Clot-Homing Near-Infrared Probe for In Vivo Imaging of Murine Thromboembolic Models.. <i>Advanced Healthcare Materials</i> , <b>2022</b> , e2102213	10.1	0
2	A versatile insertion point on albumin to accommodate peptides and maintain their activities.. <i>International Journal of Biological Macromolecules</i> , <b>2022</b> , 205, 49-49	7.9	0

- 1      Structural determination of group A Streptococcal surface dehydrogenase and characterization of its interaction with urokinase-type plasminogen activator receptor. *Biochemical and Biophysical Research Communications*, **2019**, 510, 539-544 3.4