Mattia Mori

List of Publications by Year
in descending order

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129	2,607	212478	325983
papers	citations	28 h-index	40 g-index

132	132	132	4398
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Conformational insights into the C-terminal mutations of human rhodopsin in retinitis pigmentosa. Journal of Molecular Graphics and Modelling, 2022, 110, 108076.	1.3	1
2	The Triprenylated Anthranoid Ferruginin A, a Promising Scaffold for the Development of Novel Antibiotics against Gram-Positive Bacteria. Antibiotics, 2022, 11, 84.	1.5	0
3	Acipimox inhibits human carbonic anhydrases. Journal of Enzyme Inhibition and Medicinal Chemistry, 2022, 37, 672-679.	2.5	5
4	SARS-CoV-2 Nsp13 encodes for an HLA-E-stabilizing peptide that abrogates inhibition of NKG2A-expressing NK cells. Cell Reports, 2022, 38, 110503.	2.9	31
5	Rational design and synthesis of a novel BODIPY-based probe for selective imaging of tau tangles in human iPSC-derived cortical neurons. Scientific Reports, 2022, 12, 5257.	1.6	11
6	Hidden in Plantsâ€"A Review of the Anticancer Potential of the Solanaceae Family in In Vitro and In Vivo Studies. Cancers, 2022, 14, 1455.	1.7	13
7	Discovery of spirooxadiazoline oxindoles with dual-stage antimalarial activity. European Journal of Medicinal Chemistry, 2022, 236, 114324.	2.6	9
8	Esc peptides as novel potentiators of defective cystic fibrosis transmembrane conductance regulator: an unprecedented property of antimicrobial peptides. Cellular and Molecular Life Sciences, 2022, 79, 1.	2.4	4
9	Thienoguanosine, a unique non-perturbing reporter for investigating rotational dynamics of DNA duplexes and their complexes with proteins. International Journal of Biological Macromolecules, 2022, 213, 210-225.	3.6	5
10	Potency and Selectivity Optimization of Tryptophanolâ€Derived Oxazoloisoindolinones: Novel p53 Activators in Human Colorectal Cancer. ChemMedChem, 2021, 16, 250-258.	1.6	6
11	Glabrescione B delivery by self-assembling micelles efficiently inhibits tumor growth in preclinical models of Hedgehog-dependent medulloblastoma. Cancer Letters, 2021, 499, 220-231.	3.2	22
12	Repurposing drugs for the management of COVID-19. Expert Opinion on Therapeutic Patents, 2021, 31, 295-307.	2.4	49
13	A unique high-diversity natural product collection as a reservoir of new therapeutic leads. Organic Chemistry Frontiers, 2021, 8, 996-1025.	2.3	20
14	Sofosbuvir Selects for Drug-Resistant Amino Acid Variants in the Zika Virus RNA-Dependent RNA-Polymerase Complex In Vitro. International Journal of Molecular Sciences, 2021, 22, 2670.	1.8	4
15	Design and Synthesis of Piperazine-Based Compounds Conjugated to Humanized Ferritin as Delivery System of siRNA in Cancer Cells. Bioconjugate Chemistry, 2021, 32, 1105-1116.	1.8	14
16	Active Components from Cassia abbreviata Prevent HIV-1 Entry by Distinct Mechanisms of Action. International Journal of Molecular Sciences, 2021, 22, 5052.	1.8	6
17	Design and Synthesis of New Withaferin A Inspired Hedgehog Pathway Inhibitors. Chemistry - A European Journal, 2021, 27, 8350-8357.	1.7	5
18	Hepatic miR-144 Drives Fumarase Activity Preventing NRF2 Activation During Obesity. Gastroenterology, 2021, 161, 1982-1997.e11.	0.6	34

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19	A Selective Competitive Inhibitor of Aldehyde Dehydrogenase 1A3 Hinders Cancer Cell Growth, Invasiveness and Stemness In Vitro. Cancers, 2021, 13, 356.	1.7	21
20	Identification of Effective Anticancer G-Quadruplex-Targeting Chemotypes through the Exploration of a High Diversity Library of Natural Compounds. Pharmaceutics, 2021, 13, 1611.	2.0	12
21	Pharmacological Treatment of Malaria. Topics in Medicinal Chemistry, 2021, , 219-240.	0.4	1
22	Challenges and Promises for Obtaining New Antiprotozoal Drugs: Whatâ€™s Going Wrong?. Topics in Medicinal Chemistry, 2021, , 321-329.	0.4	3
23	Statins interfere with the attachment of S. cerevisiae mtDNA to the inner mitochondrial membrane. Journal of Enzyme Inhibition and Medicinal Chemistry, 2020, 35, 129-138.	2.5	9
24	Inhibition of Melanoma Cell Migration and Invasion Targeting the Hypoxic Tumor Associated CAXII. Cancers, 2020, 12, 3018.	1.7	13
25	A Class of Potent Inhibitors of the HIV-1 Nucleocapsid Protein Based on Aminopyrrolic Scaffolds. ACS Medicinal Chemistry Letters, 2020, 11, 698-705.	1.3	4
26	Inhibitory Effect of Lithospermic Acid on the HIV-1 Nucleocapsid Protein. Molecules, 2020, 25, 5434.	1.7	5
27	Targeting the RdRp of Emerging RNA Viruses: The Structure-Based Drug Design Challenge. Molecules, 2020, 25, 5695.	1.7	64
28	ent-Beyerane Diterpenes as a Key Platform for the Development of ArnT-Mediated Colistin Resistance Inhibitors. Journal of Organic Chemistry, 2020, 85, 10891-10901.	1.7	16
29	Sempervirine inhibits RNA polymerase I transcription independently from p53 in tumor cells. Cell Death Discovery, 2020, 6, 111.	2.0	10
30	14-3-3 binding creates a memory of kinase action by stabilizing the modified state of phospholamban. Science Signaling, 2020, 13, .	1.6	19
31	What Makes Thienoguanosine an Outstanding Fluorescent DNA Probe?. Journal of the American Chemical Society, 2020, 142, 16999-17014.	6.6	27
32	Experimental and Computational Druggability Exploration of the 14-3-3Î¶/SOS1pS1161 PPI Interface. Journal of Chemical Information and Modeling, 2020, 60, 6555-6565.	2.5	5
33	In Memory of Maurizio Botta: His Vision of Medicinal Chemistry. ACS Medicinal Chemistry Letters, 2020, 11, 611-611.	1.3	0
34	A novel colistin adjuvant identified by virtual screening for ArnT inhibitors. Journal of Antimicrobial Chemotherapy, 2020, 75, 2564-2572.	1.3	15
35	Identification of Phosphate-Containing Compounds as New Inhibitors of 14-3-3/c-Abl Proteinâ€“Protein Interaction. ACS Chemical Biology, 2020, 15, 1026-1035.	1.6	9
36	5,6-Dihydroxypyrimidine Scaffold to Target HIV-1 Nucleocapsid Protein. ACS Medicinal Chemistry Letters, 2020, 11, 766-772.	1.3	5

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37	A chalcone derivative binds a putative allosteric site of YopH: Inhibition of a virulence factor of Yersinia. Bioorganic and Medicinal Chemistry Letters, 2020, 30, 127350.	1.0	5
38	Alvaxanthone, a Thymidylate Synthase Inhibitor with Nematocidal and Tumoricidal Activities. Molecules, 2020, 25, 2894.	1.7	2
39	(Thia)calixarenephosphonic Acids as Potent Inhibitors of the Nucleic Acid Chaperone Activity of the HIV-1 Nucleocapsid Protein with a New Binding Mode and Multitarget Antiviral Activity. ACS Infectious Diseases, 2020, 6, 687-702.	1.8	9
40	Hedgehog signaling pathway inhibitors: an updated patent review (2015â€“present). Expert Opinion on Therapeutic Patents, 2020, 30, 235-250.	2.4	37
41	Discovery of small molecule inhibitors of Leishmania braziliensis Hsp90 chaperone. Journal of Enzyme Inhibition and Medicinal Chemistry, 2020, 35, 639-649.	2.5	13
42	Evaluation of sofosbuvir activity and resistance profile against West Nile virus in vitro. Antiviral Research, 2020, 175, 104708.	1.9	30
43	Structural Elucidation and Antimicrobial Characterization of Novel Diterpenoids from Fabiana densa var. ramulosa. ACS Medicinal Chemistry Letters, 2020, 11, 760-765.	1.3	14
44	A deadly spillover: SARS-CoV-2 outbreak. Expert Opinion on Therapeutic Patents, 2020, 30, 481-485.	2.4	29
45	Imidazo[1,2-a]pyridine Derivatives as Aldehyde Dehydrogenase Inhibitors: Novel Chemotypes to Target Glioblastoma Stem Cells. Journal of Medicinal Chemistry, 2020, 63, 4603-4616.	2.9	38
46	Dual SMO/BRAF Inhibition by Flavonolignans from Silybum marianum. Antioxidants, 2020, 9, 384.	2.2	13
47	Natural Products as an Important Source in Drug Discovery. Current Pharmaceutical Design, 2020, 26, 2805-2806.	0.9	2
48	Identification of a new family of pyrazolo[3,4-d]pyrimidine derivatives as multitarget Fyn-Blk-Lyn inhibitors active on B- and T-lymphoma cell lines. European Journal of Medicinal Chemistry, 2019, 181, 111545.	2.6	13
49	A Molecular Tool Targeting the Baseâ€Flipping Activity of Human UHRF1. Chemistry - A European Journal, 2019, 25, 13363-13375.	1.7	8
50	A Smo/Gli Multitarget Hedgehog Pathway Inhibitor Impairs Tumor Growth. Cancers, 2019, 11, 1518.	1.7	39
51	1H-NMR metabolomics reveals the Glabrescione B exacerbation of glycolytic metabolism beside the cell growth inhibitory effect in glioma. Cell Communication and Signaling, 2019, 17, 108.	2.7	30
52	Call for Papers: Special Issue in Honor of Dr. Maurizio Botta. ACS Medicinal Chemistry Letters, 2019, 10, 1241-1241.	1.3	0
53	Nigritanine as a New Potential Antimicrobial Alkaloid for the Treatment of Staphylococcus aureus-Induced Infections. Toxins, 2019, 11, 511.	1.5	37
54	Synthesis of distal and proximal fleximer base analogues and evaluation in the nucleocapsid protein of HIV-1. Bioorganic and Medicinal Chemistry, 2019, 27, 2883-2892.	1.4	10

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55	Excitedâ€State Dynamics of Thienoguanosine, an Isomorphic Highly Fluorescent Analogue of Guanosine. Chemistry - A European Journal, 2019, 25, 7375-7386.	1.7	11
56	Chemically stable inhibitors of 14-3-3 proteinâ€“protein interactions derived from BV02. Journal of Enzyme Inhibition and Medicinal Chemistry, 2019, 34, 657-664.	2.5	12
57	Chalcones and Chalcone-mimetic Derivatives as Notch Inhibitors in a Model of T-cell Acute Lymphoblastic Leukemia. ACS Medicinal Chemistry Letters, 2019, 10, 639-643.	1.3	23
58	In Memory of Maurizio Botta: His Contribution to the Development of Computer-Aided Drug Design. Journal of Chemical Information and Modeling, 2019, 59, 4961-4967.	2.5	3
59	Synthesis and Evaluation of Bifunctional Aminothiazoles as Antiretrovirals Targeting the HIV-1 Nucleocapsid Protein. ACS Medicinal Chemistry Letters, 2019, 10, 463-468.	1.3	9
60	P300/CBPâ€associated factor regulates transcription and function of isocitrate dehydrogenase 2 during muscle differentiation. FASEB Journal, 2019, 33, 4107-4123.	0.2	11
61	Natural Products Inspired Modulators of Cancer Stem Cells-specific Signaling Pathways Notch and Hedgehog. Current Pharmaceutical Design, 2019, 24, 4251-4269.	0.9	21
62	Itch/Î²-arrestin2-dependent non-proteolytic ubiquitylation of SuFu controls Hedgehog signalling and medulloblastoma tumorigenesis. Nature Communications, 2018, 9, 976.	5.8	53
63	Design, synthesis, SAR and biological investigation of 3-(carboxymethyl)rhodanine and aminothiazole inhibitors of Mycobacterium tuberculosis Zmp1. Bioorganic and Medicinal Chemistry Letters, 2018, 28, 637-641.	1.0	13
64	Chemical, computational and functional insights into the chemical stability of the Hedgehog pathway inhibitor GANT61. Journal of Enzyme Inhibition and Medicinal Chemistry, 2018, 33, 349-358.	2.5	45
65	Identification of novel 2-benzoxazolinone derivatives with specific inhibitory activity against the HIV-1 nucleocapsid protein. European Journal of Medicinal Chemistry, 2018, 145, 154-164.	2.6	10
66	Naturally occurring Diels-Alder-type adducts from Morus nigra as potent inhibitors of Mycobacterium tuberculosis protein tyrosine phosphatase B. European Journal of Medicinal Chemistry, 2018, 144, 277-288.	2.6	29
67	Structure-Based Identification of HIV-1 Nucleocapsid Protein Inhibitors Active against Wild-Type and Drug-Resistant HIV-1 Strains. ACS Chemical Biology, 2018, 13, 253-266.	1.6	13
68	A promising natural product, pristimerin, results in cytotoxicity against breast cancer stem cells in vitro and xenografts in vivo through apoptosis and an incomplete autopaghy in breast cancer. Pharmacological Research, 2018, 129, 500-514.	3.1	62
69	Stable Oxidative Cytosine Modifications Accumulate in Cardiac Mesenchymal Cells From Type2 Diabetes Patients. Circulation Research, 2018, 122, 31-46.	2.0	33
70	Synthetic thiosemicarbazones as a new class of Mycobacterium tuberculosis protein tyrosine phosphatase A inhibitors. Bioorganic and Medicinal Chemistry, 2018, 26, 5742-5750.	1.4	21
71	Current trends in Hedgehog signaling pathway inhibition by small molecules. Bioorganic and Medicinal Chemistry Letters, 2018, 28, 3131-3140.	1.0	30
72	A High-throughput Screening of a Chemical Compound Library in Ovarian Cancer Stem Cells. Combinatorial Chemistry and High Throughput Screening, 2018, 21, 50-56.	0.6	3

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73	Potent and Selective Carboxylic Acid Inhibitors of Tumor-Associated Carbonic Anhydrases IX and XII. Molecules, 2018, 23, 17.	1.7	14
74	Synergistic inhibition of the Hedgehog pathway by newly designed Smo and Gli antagonists bearing the isoflavone scaffold. European Journal of Medicinal Chemistry, 2018, 156, 554-562.	2.6	29
75	Structure, Function, Involvement in Diseases and Targeting of 14-3-3 Proteins: An Update. Current Medicinal Chemistry, 2018, 25, 5-21.	1.2	56
76	Oregonin from Alnus incana bark affects DNA methyltransferases expression and mitochondrial DNA copies in mouse embryonic fibroblasts. Journal of Enzyme Inhibition and Medicinal Chemistry, 2018, 33, 1055-1063.	2.5	7
77	Design, Palladium-Catalyzed Synthesis, and Biological Investigation of 2-Substituted 3-Aroylquinolin-4(1H)-ones as Inhibitors of the Hedgehog Signaling Pathway. Journal of Medicinal Chemistry, 2017, 60, 1469-1477.	2.9	26
78	Novel coumarin- and quinolinone-based polycycles as cell division cycle 25-A and -C phosphatases inhibitors induce proliferation arrest and apoptosis in cancer cells. European Journal of Medicinal Chemistry, 2017, 134, 316-333.	2.6	24
79	Identification of a novel chalcone derivative that inhibits Notch signaling in T-cell acute lymphoblastic leukemia. Scientific Reports, 2017, 7, 2213.	1.6	42
80	Highâ€Affinity â€œClickâ€•RGD Peptidomimetics as Radiolabeled Probes for Imaging Î±_vÎ²₃ Integrin. ChemMedChem, 2017, 12, 1142-1151.	1.6	13
81	Differentially activated Src kinase in chemoâ€naÃ¯ve human primary osteosarcoma cells and effects of a Src kinase inhibitor. BioFactors, 2017, 43, 801-811.	2.6	8
82	Natural modulators of nonalcoholic fatty liver disease: Mode of action analysis and in silico ADME-Tox prediction. Toxicology and Applied Pharmacology, 2017, 337, 45-66.	1.3	14
83	Synthesis, biological evaluation and molecular modeling studies on novel quinonoid inhibitors of CDC25 phosphatases. Journal of Enzyme Inhibition and Medicinal Chemistry, 2017, 32, 113-118.	2.5	11
84	Novel Sulfamide-Containing Compounds as Selective Carbonic Anhydrase I Inhibitors. Molecules, 2017, 22, 1049.	1.7	24
85	Green Routes for the Production of Enantiopure Benzylisoquinoline Alkaloids. International Journal of Molecular Sciences, 2017, 18, 2464.	1.8	12
86	Editorial (Thematic Issue: Challenging Organic Syntheses and Pharmacological Applications of) Tj ETQq0 0 0 rgBT /Overlock 10 Tf 50 222	0.9	4
87	Tautomers of a Fluorescent G Surrogate and Their Distinct Photophysics Provide Additional Information Channels. Angewandte Chemie - International Edition, 2016, 55, 7974-7978.	7.2	36
88	Tautomers of a Fluorescent G Surrogate and Their Distinct Photophysics Provide Additional Information Channels. Angewandte Chemie, 2016, 128, 8106-8110.	1.6	11
89	Histone demethylating agents as potential S-adenosyl-<scp>l</scp>-methionine-competitors. MedChemComm, 2016, 7, 1245-1255.	3.5	5
90	Discovery of inÂvitro antitubercular agents through in silico ligand-based approaches. European Journal of Medicinal Chemistry, 2016, 121, 169-180.	2.6	22

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91	Yeast as a tool to select inhibitors of the cullin deneddylating enzyme Csn5. Journal of Enzyme Inhibition and Medicinal Chemistry, 2016, 31, 1632-1637.	2.5	15
92	Inhibition of Hedgehog-dependent tumors and cancer stem cells by a newly identified naturally occurring chemotype. Cell Death and Disease, 2016, 7, e2376-e2376.	2.7	49
93	The plant-derived triterpenoid tingenin B is a potent anticancer agent due to its cytotoxic activity on cancer stem cells of breast cancer inÂvitro. Chemico-Biological Interactions, 2016, 260, 248-255.	1.7	20
94	Mycobacterial carbonic anhydrase inhibition with phenolic acids and esters: kinetic and computational investigations. Organic and Biomolecular Chemistry, 2016, 14, 8322-8330.	1.5	29
95	Occurrence of Enantioselectivity in Nature: The Case of (S)â€Norcoclaurine. Chirality, 2016, 28, 169-180.	1.3	19
96	Molecular insights to the bioactive form of BV02 , a reference inhibitor of 14-3-3Ïƒ proteinâ€“protein interactions. Bioorganic and Medicinal Chemistry Letters, 2016, 26, 894-898.	1.0	10
97	One Hundred Faces of Cyclopamine. Current Pharmaceutical Design, 2016, 22, 1658-1681.	0.9	21
98	Mycobacterium tuberculosis-Secreted Tyrosine Phosphatases as Targets Against Tuberculosis: Exploring Natural Sources in Searching for New Drugs. Current Pharmaceutical Design, 2016, 22, 1561-1569.	0.9	20
99	Click Reaction as a Tool to Combine Pharmacophores: The Case of Vismodegib. ChemPlusChem, 2015, 80, 938-943.	1.3	19
100	Editorial (Thematic Issue: Challenging Organic Syntheses and Pharmacological Applications of) Tj ETQq0 0 0 rgBT /Overlock 10 Tf 50 382	0.9	8
101	Gli1/ <scp>DNA</scp> interaction is a druggable target for Hedgehogâ€dependent tumors. EMBO Journal, 2015, 34, 200-217.	3.5	147
102	Nucleocapsid Protein: A Desirable Target for Future Therapies Against HIV-1. Current Topics in Microbiology and Immunology, 2015, 389, 53-92.	0.7	56
103	Targeting GLI factors to inhibit the Hedgehog pathway. Trends in Pharmacological Sciences, 2015, 36, 547-558.	4.0	100
104	Hit Recycling: Discovery of a Potent Carbonic Anhydrase Inhibitor by in Silico Target Fishing. ACS Chemical Biology, 2015, 10, 1964-1969.	1.6	19
105	Exploring Oxidovanadium(IV) Complexes as YopH Inhibitors: Mechanism of Action and Modeling Studies. ACS Medicinal Chemistry Letters, 2015, 6, 1035-1040.	1.3	17
106	Molecular Dynamics Simulations and Structural Analysis of Giardia duodenalis 14-3-3 Proteinâ€“Protein Interactions. Journal of Chemical Information and Modeling, 2015, 55, 2611-2622.	2.5	23
107	Hydrolytic inhibition of Î±-chymotrypsin by 2,8,14,20-tetrakis(<scp>d</scp>-leucyl-<scp>d</scp>-valinamido)resorc[4]arenecarboxylic acid: a spectroscopic NMR and computational combined approach. Organic and Biomolecular Chemistry, 2015, 13, 916-924.	1.5	3
108	Synergistic Effects of Trace Amounts of Water in the Enantiodiscrimination Processes by Lipodex E: A Spectroscopic and Computational Investigation. Chirality, 2015, 27, 95-103.	1.3	8

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109	Discovery of 14â€3â€3 Proteinâ€“Protein Interaction Inhibitors that Sensitize Multidrugâ€Resistant Cancer Cells to Doxorubicin and the Akt Inhibitor GSK690693. ChemMedChem, 2014, 9, 973-983.	1.6	30
110	Functional and Structural Characterization of 2-Amino-4-phenylthiazole Inhibitors of the HIV-1 Nucleocapsid Protein with Antiviral Activity. ACS Chemical Biology, 2014, 9, 1950-1955.	1.6	25
111	Discovery of the first potent and selective Mycobacterium tuberculosis Zmp1 inhibitor. Bioorganic and Medicinal Chemistry Letters, 2014, 24, 2508-2511.	1.0	22
112	Free Energy Profile and Kinetics Studies of Paclitaxel Internalization from the Outer to the Inner Wall of Microtubules. Journal of Chemical Theory and Computation, 2013, 9, 698-706.	2.3	9
113	Small molecules modulation of 14-3-3 proteinâ€“protein interactions. Drug Discovery Today: Technologies, 2013, 10, e541-e547.	4.0	24
114	Discovery of a New Class of Potent MMP Inhibitors by Structure-Based Optimization of the Arylsulfonamide Scaffold. ACS Medicinal Chemistry Letters, 2013, 4, 565-569.	1.3	18
115	A Combination Strategy to Inhibit Pimâ€1: Synergism between Noncompetitive and ATPâ€Competitive Inhibitors. ChemMedChem, 2013, 8, 484-496.	1.6	13
116	Discovery of Mycobacterium tuberculosis Protein Tyrosine Phosphatase B (PtpB) Inhibitors from Natural Products. PLoS ONE, 2013, 8, e77081.	1.1	46
117	Structure Prediction and Validation of the ERK8 Kinase Domain. PLoS ONE, 2013, 8, e52011.	1.1	10
118	Use of virtual screening for discovering antiretroviral compounds interacting with the HIV-1 nucleocapsid protein. Virus Research, 2012, 169, 377-387.	1.1	25
119	A Highly Soluble Matrix Metalloproteinaseâ€9 Inhibitor for Potential Treatment of Dry Eye Syndrome. Basic and Clinical Pharmacology and Toxicology, 2012, 111, 289-295.	1.2	14
120	Predicting the Binding Mode of Known NCp7 Inhibitors To Facilitate the Design of Novel Modulators. Journal of Chemical Information and Modeling, 2011, 51, 446-454.	2.5	34
121	Computational techniques are valuable tools for the discovery of proteinâ€“protein interaction inhibitors: The 14-3-3ĩ*f* case. Bioorganic and Medicinal Chemistry Letters, 2011, 21, 6867-6871.	1.0	31
122	Targeting Protein-Protein and Protein-Nucleic Acid Interactions for Anti-HIV Therapy. Current Pharmaceutical Design, 2011, 17, 3713-3728.	0.9	22
123	Probing the Pore Drug Binding Site of Microtubules with Fluorescent Taxanes: Evidence of Two Binding Poses. Chemistry and Biology, 2010, 17, 243-253.	6.2	21
124	2-Hydroxypropyl-Î²-cyclodextrin strongly improves water solubility and anti-proliferative activity of pyrazolo[3,4-d]pyrimidines Src-Abl dual inhibitors. European Journal of Medicinal Chemistry, 2010, 45, 5958-5964.	2.6	36
125	Molecular Dynamics and DFT Study on HIV-1 Nucleocapsid Protein-7 in Complex with Viral Genome. Journal of Chemical Information and Modeling, 2010, 50, 638-650.	2.5	41
126	Fragment Docking to S100 Proteins Reveals a Wide Diversity of Weak Interaction Sites. ChemMedChem, 2007, 2, 1648-1654.	1.6	14

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127	Theoretical study on binding of S100B protein. Journal of Molecular Modeling, 2007, 13, 1123-1131.	0.8	7
128	Hetero Dielsâ€“Alder reactions (HDAR) of Î±,Î±â€²-dioxothiones on solid support. Tetrahedron, 2005, 61, 5005-5010.	1.0	12
129	Inverse Electron Demand Hetero Dielsâ€“Alder Reactions of Solid Supported Î±-Acilthiones. Phosphorus, Sulfur and Silicon and the Related Elements, 2005, 180, 1327-1331.	0.8	2