Daniel P Becker

List of Publications by Year in descending order

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361296 434063 1,169 59 20 31 citations h-index g-index papers 62 62 62 1227 all docs docs citations times ranked citing authors

#	Article	IF	CITATIONS
1	Iron dysregulation in COVIDâ€19 and reciprocal evolution of SARSâ€CoVâ€2: Natura nihil frustra facit. Journal of Cellular Biochemistry, 2022, 123, 601-619.	1.2	21
2	Bisindolylmaleimide IX: A novel anti-SARS-CoV2 agent targeting viral main protease 3CLpro demonstrated by virtual screening pipeline and in-vitro validation assays. Methods, 2021, 195, 57-71.	1.9	29
3	Atomic-Resolution 1.3 Ã Crystal Structure, Inhibition by Sulfate, and Molecular Dynamics of the Bacterial Enzyme DapE. Biochemistry, 2021, 60, 908-917.	1.2	6
4	Gcn5-Related N-Acetyltransferases (GNATs) With a Catalytic Serine Residue Can Play Ping-Pong Too. Frontiers in Molecular Biosciences, 2021, 8, 646046.	1.6	8
5	Cyclobutanone Inhibitor of Cobalt-Functionalized Metallo- \hat{l}^3 -Lactonase AiiA with Cyclobutanone Ring Opening in the Active Site. ACS Omega, 2021, 6, 13567-13578.	1.6	3
6	Indoline-6-Sulfonamide Inhibitors of the Bacterial Enzyme DapE. Antibiotics, 2020, 9, 595.	1.5	9
7	(S)â€4―Amino â€5â€phenoxypentanoate designed as a potential selective agonist of the bacterial transcription factor GabR. Protein Science, 2020, 29, 1816-1828.	3.1	3
8	Carboraneâ€Containing Matrix Metalloprotease (MMP) Ligands as Candidates for Boron Neutronâ€Capture Therapy (BNCT). ChemMedChem, 2020, 15, 1897-1908.	1.6	12
9	A Review of the Preclinical and Clinical Efficacy of Remdesivir, Hydroxychloroquine, and Lopinavir-Ritonavir Treatments against COVID-19. SLAS Discovery, 2020, 25, 1108-1122.	1.4	25
10	Synthesis of a protected 2-aminocyclobutanone as a modular transition state synthon for medicinal chemistry. Tetrahedron Letters, 2020, 61, 151632.	0.7	5
11	Structural Evidence of a Major Conformational Change Triggered by Substrate Binding in DapE Enzymes: Impact on the Catalytic Mechanism. Biochemistry, 2018, 57, 574-584.	1.2	16
12	Iron(II) complexes of dimethyltriazacyclophane. Acta Crystallographica Section C, Structural Chemistry, 2018, 74, 1641-1649.	0.2	0
13	A Gcn5-RelatedN-Acetyltransferase (GNAT) Capable of Acetylating Polymyxin B and Colistin Antibioticsin Vitro. Biochemistry, 2018, 57, 7011-7020.	1.2	11
14	Attempted Resolution and Racemization of Beckmannâ€Derived CTV‣actam and the Use of Chirabiteâ€AR® to Determine the Optical Purity of the Supramolecular Scaffold. European Journal of Organic Chemistry, 2018, 2018, 4639-4645.	1.2	2
15	Practical spectrophotometric assay for the dapE-encoded N-succinyl-L,L-diaminopimelic acid desuccinylase, a potential antibiotic target. PLoS ONE, 2018, 13, e0196010.	1.1	11
16	Generating enzyme and radicalâ€mediated bisubstrates as tools for investigating Gcn5â€related <i>Nâ€</i> acetyltransferases. FEBS Letters, 2017, 591, 2348-2361.	1.3	5
17	An analytical method for detecting toxic metal cations using cyclotriveratrylene derivative capped gold nanoparticles. Tetrahedron Letters, 2015, 56, 5419-5423.	0.7	7
18	Metal-free tandem Beckmann–electrophilic aromatic substitution cascade affording diaryl imines, ketones, amines, and quinazolines. Tetrahedron Letters, 2015, 56, 5390-5392.	0.7	5

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19	Chapter 12. New Analytical Approaches for the Detection of Micropollutants in Natural Waters: Identification of 3,5-Dichloro-4-Hydroxybenzene Sulfonic Acid as an Unknown Persistent Pollutant. Comprehensive Series in Photochemical and Photobiological Sciences, 2015, , 239-256.	0.3	О
20	Lysine biosynthesis in bacteria: a metallodesuccinylase as a potential antimicrobial target. Journal of Biological Inorganic Chemistry, 2013, 18, 155-163.	1.1	57
21	Rearrangement of Cyclotriveratrylene (CTV) Diketone: 9,10-Diarylanthracenes with OLED Applications. Journal of Organic Chemistry, 2013, 78, 2051-2058.	1.7	18
22	Apparent Alkyl Transfer and Phenazine Formation via an Aryne Intermediate. Journal of Organic Chemistry, 2013, 78, 3532-3540.	1.7	11
23	Large Scale Structural Rearrangement of a Serine Hydrolase from Francisella tularensis Facilitates Catalysis. Journal of Biological Chemistry, 2013, 288, 10522-10535.	1.6	28
24	Synthesis, crystal structure and rearrangements of orthocyclophane cyclotetraveratrylene (CTTV) tetraketone. Supramolecular Chemistry, 2012, 24, 803-809.	1.5	6
25	Asymmetric α-hydroxy ketone synthesis by direct ketone oxidation using a bimetallic palladium(II) complex. Tetrahedron Letters, 2012, 53, 2699-2701.	0.7	26
26	MMP-13 selective alpha-sulfone hydroxamates: Identification of selective P1â \in 2 amides. Bioorganic and Medicinal Chemistry Letters, 2011, 21, 2823-2825.	1.0	11
27	MMP-13 selective α-sulfone hydroxamates: A survey of P1′ heterocyclic amide isosteres. Bioorganic and Medicinal Chemistry Letters, 2011, 21, 2820-2822.	1.0	11
28	Direct patterning of a cyclotriveratrylene derivative for directed self-assembly of C ₆₀ . Nanotechnology, 2011, 22, 275611.	1.3	8
29	Synthesis of an <i>ortho</i> -Triazacyclophane: <i>N</i> , <i>N</i> ,6≥3,6≥3,6≥3,6≥3,6≥3,6≥3,6≥3,6≥3,1 Chemistry, 2010, 75, 7887-7892.	1.7	22
30	Palladium(II)-catalyzed dicarboxymethylation of chiral allylic alcohols: chirality transfer affording optically active diesters containing three contiguous chiral centers. Tetrahedron Letters, 2010, 51, 3514-3517.	0.7	12
31	MMP-13 selective isonipecotamide α-sulfone hydroxamates. Bioorganic and Medicinal Chemistry Letters, 2010, 20, 3561-3564.	1.0	29
32	Orally bioavailable dual MMP-1/MMP-14 sparing, MMP-13 selective α-sulfone hydroxamates. Bioorganic and Medicinal Chemistry Letters, 2010, 20, 3557-3560.	1.0	24
33	Orally Active MMP-1 Sparing α-Tetrahydropyranyl and α-Piperidinyl Sulfone Matrix Metalloproteinase (MMP) Inhibitors with Efficacy in Cancer, Arthritis, and Cardiovascular Disease. Journal of Medicinal Chemistry, 2010, 53, 6653-6680.	2.9	89
34	An aza-cyclophane stacked in racemic columnar assemblies: whole-molecule disorder in a two-dimensional solid solution. Acta Crystallographica Section B: Structural Science, 2009, 65, 223-229.	1.8	3
35	Inhibitors of bacterial N-succinyl-I,I-diaminopimelic acid desuccinylase (DapE) and demonstration of in vitro antimicrobial activity. Bioorganic and Medicinal Chemistry Letters, 2009, 19, 6350-6352.	1.0	39
36	A Thermodynamic and Kinetic Characterization of the Solvent Dependence of the Saddleâ^'Crown Equilibrium of Cyclotriveratrylene Oxime. Journal of Physical Chemistry A, 2009, 113, 8258-8267.	1.1	11

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37	Beckmann rearrangement of cyclotriveratrylene (CTV) oxime: tandem Beckmann-electrophilic aromatic addition. Tetrahedron Letters, 2008, 49, 5003-5005.	0.7	21
38	Synthesis of Substituted 2â€Aminoâ€cyclobutanones. Synthetic Communications, 2008, 38, 1679-1687.	1.1	12
39	Isolation of the saddle and crown conformers of cyclotriveratrylene (CTV) oxime. Tetrahedron Letters, 2007, 48, 6368-6371.	0.7	16
40	2,3,8,12,13-Pentamethoxy-5 <i>H</i> -dibenzo[<i>c</i> , <i>n</i>]acridin-7(6 <i>H</i>)-one toluene solvate. Acta Crystallographica Section E: Structure Reports Online, 2007, 63, o3857-o3858.	0.2	2
41	2,3,5′,6,6′,7-Hexamethoxy-3′H,10H-spiro[anthracene-9,1′-isobenzofuran]-3′,10-dione. Acta Crystallo Section E: Structure Reports Online, 2007, 63, o4390-o4391.	ographica 6.2	3
42	Pyrrolizidine Esters and Amides as 5-HT4Receptor Agonists and Antagonists. Journal of Medicinal Chemistry, 2006, 49, 1125-1139.	2.9	40
43	Synthesis and Structureâ 'Activity Relationships of \hat{I}^2 - and $\hat{I}\pm$ -Piperidine Sulfone Hydroxamic Acid Matrix Metalloproteinase Inhibitors with Oral Antitumor Efficacy. Journal of Medicinal Chemistry, 2005, 48, 6713-6730.	2.9	64
44	Bridgehead-methyl analog of SC-53116 as a 5-HT4 agonist. Bioorganic and Medicinal Chemistry Letters, 2004, 14, 3073-3075.	1.0	15
45	Azaadamantane benzamide 5-HT4 agonists: gastrointestinal prokinetic SC-54750. Bioorganic and Medicinal Chemistry Letters, 2004, 14, 5509-5512.	1.0	11
46	Selective, orally active MMP inhibitors with an aryl backbone. Bioorganic and Medicinal Chemistry Letters, 2001, 11, 2481-2483.	1.0	17
47	\hat{l}_{\pm} -Amino- \hat{l}^2 -sulphone hydroxamates as potent MMP-13 inhibitors that spare MMP-1. Bioorganic and Medicinal Chemistry Letters, 2001, 11, 2719-2722.	1.0	40
48	α-Alkyl-α-amino-β-sulphone hydroxamates as potent MMP inhibitors that spare MMP-1. Bioorganic and Medicinal Chemistry Letters, 2001, 11, 2723-2725.	1.0	29
49	Synthesis and activity of selective MMP inhibitors with an aryl backbone. Bioorganic and Medicinal Chemistry Letters, 2000, 10, 2815-2817.	1.0	30
50	Enantioselective synthesis of dual serotonergic azanoradamantane SC-52491. Tetrahedron, 1999, 55, 11787-11802.	1.0	14
51	Serotonin 5-HT4 agonist activity of a series of meso-azanoradamantane benzamides. Bioorganic and Medicinal Chemistry Letters, 1997, 7, 2149-2154.	1.0	6
52	One-Pot Preparation of 1,3-Dihydro-1-(trifluoromethyl)isobenzofuran-1-ol Derivatives from 1,2-Dibromobenzene. Synthetic Communications, 1996, 26, 3127-3135.	1.1	7
53	Synthesis of N-BOC-3-azabicyclo[3.3.0]octan-7-one via reductive pauson-khand cyclization and subsequent conversion to a novel diazatricyclic ring system. Tetrahedron, 1993, 49, 5047-5054.	1.0	35
54	Studies of the solid-phase Pauson-Khand reaction: Selective in-situ enone reduction to 3-azabicyclo[3.3.0]octanones. Tetrahedron Letters, 1993, 34, 2087-2090.	0.7	24

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55	SC-53116: the first selective agonist at the newly identified serotonin 5-HT4 receptor subtype. Journal of Medicinal Chemistry, 1992, 35, 1486-1489.	2.9	84
56	Use of atom-transfer radical cyclizations as an efficient entry into a new "serotonergic― azanoradamantane. Tetrahedron Letters, 1992, 33, 7283-7286.	0.7	18
57	1,3,4-trisubstituted pyrrolidinones as scaffolds for construction of peptidomimetic cholecystokinin antagonists. Bioorganic and Medicinal Chemistry Letters, 1992, 2, 1251-1256.	1.0	15
58	New aza(nor)adamantanes are agonists at the newly identified serotonin 5-HT4 receptor and antagonists at the 5-HT3 receptor. Bioorganic and Medicinal Chemistry Letters, 1992, 2, 1613-1618.	1.0	17
59	Stereospecific dicobalt octacarbonyl-mediated enyne cyclization for the enantiospecific synthesis of a 6a-carbocycline analog. Journal of the American Chemical Society, 1987, 109, 7495-7498.	6.6	66