

Paul A Wender

List of Publications by Year in descending order

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Version: 2024-02-01

231
papers

21,122
citations

5891

81
h-index

11303

136
g-index

262
all docs

262
docs citations

262
times ranked

14674
citing authors

#	ARTICLE	IF	CITATIONS
1	Engineering circular RNA for enhanced protein production. <i>Nature Biotechnology</i> , 2023, 41, 262-272.	9.4	83
2	Latency reversal plus natural killer cells diminish HIV reservoir in vivo. <i>Nature Communications</i> , 2022, 13, 121.	5.8	36
3	Fingolimod-Conjugated Charge-Altering Releasable Transporters Efficiently and Specifically Deliver mRNA to Lymphocytes In Vivo and In Vitro. <i>Biomacromolecules</i> , 2022, 23, 2976-2988.	2.6	5
4	In Vivo Targeting of <i>Escherichia coli</i> with Vancomycin-Arginine. <i>Antimicrobial Agents and Chemotherapy</i> , 2021, 65, .	1.4	14
5	Designed PKC-targeting bryostatin analogs modulate innate immunity and neuroinflammation. <i>Cell Chemical Biology</i> , 2021, 28, 537-545.e4.	2.5	7
6	An mRNA SARS-CoV-2 Vaccine Employing Charge-Altering Releasable Transporters with a TLR-9 Agonist Induces Neutralizing Antibodies and T Cell Memory. <i>ACS Central Science</i> , 2021, 7, 1191-1204.	5.3	34
7	Clinical Correlates of Human Immunodeficiency Virus-1 (HIV-1) DNA and Inducible HIV-1 RNA Reservoirs in Peripheral Blood in Children With Perinatally Acquired HIV-1 Infection With Sustained Virologic Suppression for at Least 5 Years. <i>Clinical Infectious Diseases</i> , 2020, 70, 859-866.	2.9	20
8	Reversible RNA acylation for control of CRISPR-Cas9 gene editing. <i>Chemical Science</i> , 2020, 11, 1011-1016.	3.7	37
9	Function-Oriented Synthesis: Design, Synthesis, and Evaluation of Highly Simplified Bryostatin Analogues. <i>Journal of Organic Chemistry</i> , 2020, 85, 15116-15128.	1.7	7
10	Charge-altering releasable transporters enable phenotypic manipulation of natural killer cells for cancer immunotherapy. <i>Blood Advances</i> , 2020, 4, 4244-4255.	2.5	32
11	Prodrugs of PKC modulators show enhanced HIV latency reversal and an expanded therapeutic window. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2020, 117, 10688-10698.	3.3	34
12	In Situ Detection of Endogenous HIV Activation by Dynamic Nuclear Polarization NMR and Flow Cytometry. <i>International Journal of Molecular Sciences</i> , 2020, 21, 4649.	1.8	13
13	Synthesis and mechanistic investigations of pH-responsive cationic poly(aminoester)s. <i>Chemical Science</i> , 2020, 11, 2951-2966.	3.7	26
14	Synthesis and evaluation of designed PKC modulators for enhanced cancer immunotherapy. <i>Nature Communications</i> , 2020, 11, 1879.	5.8	29
15	Bryostatin 1 Promotes Synaptogenesis and Reduces Dendritic Spine Density in Cortical Cultures through a PKC-Dependent Mechanism. <i>ACS Chemical Neuroscience</i> , 2020, 11, 1545-1554.	1.7	16
16	Tracking HIV Rebound following Latency Reversal Using Barcoded HIV. <i>Cell Reports Medicine</i> , 2020, 1, 100162.	3.3	11
17	Vancomycin-Arginine Conjugate Inhibits Growth of Carbapenem-Resistant <i>E. coli</i> and Targets Cell-Wall Synthesis. <i>ACS Chemical Biology</i> , 2019, 14, 2065-2070.	1.6	67
18	Synthesis of Modified Nucleoside Oligophosphates Simplified: Fast, Pure, and Protecting Group Free. <i>Journal of the American Chemical Society</i> , 2019, 141, 15013-15017.	6.6	29

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19	Local Delivery of $Ox40$, $Cd80$, and $Cd86$ mRNA Kindles Global Anticancer Immunity. <i>Cancer Research</i> , 2019, 79, 1624-1634.	0.4	85
20	A Phosphoramidite Analogue of Cyclotriphosphate Enables Iterative Polyphosphorylations. <i>Angewandte Chemie - International Edition</i> , 2019, 58, 3928-3933.	7.2	23
21	A Phosphoramidite Analogue of Cyclotriphosphate Enables Iterative Polyphosphorylations. <i>Angewandte Chemie</i> , 2019, 131, 3968-3973.	1.6	8
22	Oligo(serine ester) Charge-Altering Releasable Transporters: Organocatalytic Ring-Opening Polymerization and their Use for <i>in Vitro</i> and <i>in Vivo</i> mRNA Delivery. <i>Journal of the American Chemical Society</i> , 2019, 141, 8416-8421.	6.6	61
23	Impact of Treatment Interruption on HIV Reservoirs and Lymphocyte Subsets in Individuals Who Initiated Antiretroviral Therapy During the Early Phase of Infection. <i>Journal of Infectious Diseases</i> , 2019, 220, 270-274.	1.9	11
24	REDOR NMR Reveals Multiple Conformers for a Protein Kinase C Ligand in a Membrane Environment. <i>ACS Central Science</i> , 2018, 4, 89-96.	5.3	28
25	Functional DNA Delivery Enabled by Lipid-Modified Charge-Altering Releasable Transporters (CARTs). <i>Biomacromolecules</i> , 2018, 19, 2812-2824.	2.6	29
26	Bryostatin and its synthetic analog, picolog rescue dermal fibroblasts from prolonged stress and contribute to survival and rejuvenation of human skin equivalents. <i>Journal of Cellular Physiology</i> , 2018, 233, 1523-1534.	2.0	4
27	Delivery of Inorganic Polyphosphate into Cells Using Amphipathic Oligocarbonate Transporters. <i>ACS Central Science</i> , 2018, 4, 1394-1402.	5.3	15
28	A Dual-Function Antibiotic-Transporter Conjugate Exhibits Superior Activity in Sterilizing MRSA Biofilms and Killing Persister Cells. <i>Journal of the American Chemical Society</i> , 2018, 140, 16140-16151.	6.6	109
29	mRNA vaccination with charge-altering releasable transporters elicits human T cell responses and cures established tumors in mice. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2018, 115, E9153-E9161.	3.3	92
30	Characterization of designed, synthetically accessible bryostatin analog HIV latency reversing agents. <i>Virology</i> , 2018, 520, 83-93.	1.1	33
31	Enhanced mRNA delivery into lymphocytes enabled by lipid-varied libraries of charge-altering releasable transporters. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2018, 115, E5859-E5866.	3.3	162
32	Charge-altering releasable transporters (CARTs) for the delivery and release of mRNA in living animals. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2017, 114, E448-E456.	3.3	207
33	Molecular dynamics simulations reveal ligand-controlled positioning of a peripheral protein complex in membranes. <i>Nature Communications</i> , 2017, 8, 6.	5.8	103
34	Vault Nanoparticles: Chemical Modifications for Imaging and Enhanced Delivery. <i>ACS Nano</i> , 2017, 11, 872-881.	7.3	30
35	Scalable synthesis of bryostatin 1 and analogs, adjuvant leads against latent HIV. <i>Science</i> , 2017, 358, 218-223.	6.0	86
36	Ynol Ethers as Ketene Equivalents in Rhodium-Catalyzed Intermolecular [5 + 2] Cycloaddition Reactions. <i>Organic Letters</i> , 2017, 19, 5810-5813.	2.4	18

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37	Retrosynthetic Reaction Prediction Using Neural Sequence-to-Sequence Models. ACS Central Science, 2017, 3, 1103-1113.	5.3	308
38	Combinations of isoform-targeted histone deacetylase inhibitors and bryostatin analogues display remarkable potency to activate latent HIV without global T-cell activation. Scientific Reports, 2017, 7, 7456.	1.6	32
39	In vivo activation of latent HIV with a synthetic bryostatin analog effects both latent cell "kick" and "kill" in strategy for virus eradication. PLoS Pathogens, 2017, 13, e1006575.	2.1	73
40	Comparative analysis of the anti-chikungunya virus activity of novel bryostatin analogs confirms the existence of a PKC-independent mechanism. Biochemical Pharmacology, 2016, 120, 15-21.	2.0	11
41	Cellular delivery and photochemical release of a caged inositol-pyrophosphate induces PH-domain translocation in cellulo. Nature Communications, 2016, 7, 10622.	5.8	77
42	Simplified Bryostatin Analogues Protect Cells from Chikungunya Virus-Induced Cell Death. Journal of Natural Products, 2016, 79, 675-679.	1.5	16
43	Inhibition of Chikungunya Virus-Induced Cell Death by Salicylate-Derived Bryostatin Analogues Provides Additional Evidence for a PKC-Independent Pathway. Journal of Natural Products, 2016, 79, 680-684.	1.5	28
44	Cell-Penetrating, Guanidinium-Rich Oligophosphoesters: Effective and Versatile Molecular Transporters for Drug and Probe Delivery. Journal of the American Chemical Society, 2016, 138, 3510-3517.	6.6	96
45	Bioorthogonal Catalysis: A General Method To Evaluate Metal-Catalyzed Reactions in Real Time in Living Systems Using a Cellular Luciferase Reporter System. Bioconjugate Chemistry, 2016, 27, 376-382.	1.8	58
46	Function through Synthesis-Informed Design. Accounts of Chemical Research, 2015, 48, 752-760.	7.6	61
47	Studies on the regio- and diastereo-selective epoxidation of daphnanes and tiglanes. Tetrahedron Letters, 2015, 56, 3423-3427.	0.7	16
48	Guanidinium-Rich, Glycerol-Derived Oligocarbonates: A New Class of Cell-Penetrating Molecular Transporters That Complex, Deliver, and Release siRNA. Molecular Pharmaceutics, 2015, 12, 742-750.	2.3	21
49	Catalytic Efficiency Is a Function of How Rhodium(I) (5 + 2) Catalysts Accommodate a Conserved Substrate Transition State Geometry: Induced Fit Model for Explaining Transition Metal Catalysis. ACS Catalysis, 2015, 5, 1758-1763.	5.5	30
50	Toward a Biorelevant Structure of Protein Kinase C Bound Modulators: Design, Synthesis, and Evaluation of Labeled Bryostatin Analogues for Analysis with Rotational Echo Double Resonance NMR Spectroscopy. Journal of the American Chemical Society, 2015, 137, 3678-3685.	6.6	24
51	Tetramethyleneethane Equivalents: Recursive Reagents for Serialized Cycloadditions. Journal of the American Chemical Society, 2015, 137, 9088-9093.	6.6	32
52	Function through bio-inspired, synthesis-informed design: step-economical syntheses of designed kinase inhibitors. Organic Chemistry Frontiers, 2014, 1, 1166-1171.	2.3	5
53	Reactivity and Chemoselectivity of Allenes in Rh(I)-Catalyzed Intermolecular (5 + 2) Cycloadditions with Vinylcyclopropanes: Allene-Mediated Rhodacycle Formation Can Poison Rh(I)-Catalyzed Cycloadditions. Journal of the American Chemical Society, 2014, 136, 17273-17283.	6.6	96
54	Structural complexity through multicomponent cycloaddition cascades enabled by dual-purpose, reactivity regenerating 1,2,3-triene equivalents. Nature Chemistry, 2014, 6, 448-452.	6.6	57

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55	Improved Protein Kinase C Affinity through Final Step Diversification of a Simplified Salicylate-Derived Bryostatin Analog Scaffold. <i>Organic Letters</i> , 2014, 16, 5140-5143.	2.4	20
56	Cell-Penetrating, Guanidinium-Rich Molecular Transporters for Overcoming Efflux-Mediated Multidrug Resistance. <i>Molecular Pharmaceutics</i> , 2014, 11, 2553-2565.	2.3	53
57	Computer-Guided Design, Synthesis, and Protein Kinase C Affinity of a New Salicylate-Based Class of Bryostatin Analogs. <i>Organic Letters</i> , 2014, 16, 5136-5139.	2.4	31
58	Bioengineered Vaults: Self-Assembling Protein Shell Lipophilic Core Nanoparticles for Drug Delivery. <i>ACS Nano</i> , 2014, 8, 7723-7732.	7.3	54
59	Toward the ideal synthesis and molecular function through synthesis-informed design. <i>Natural Product Reports</i> , 2014, 31, 433-440.	5.2	185
60	Propargyltrimethylsilanes as Allene Equivalents in Transition Metal-Catalyzed [5 + 2] Cycloadditions. <i>Organic Letters</i> , 2014, 16, 2923-2925.	2.4	25
61	Highly potent, synthetically accessible prostratin analogs induce latent HIV expression in vitro and ex vivo. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2013, 110, 11698-11703.	3.3	130
62	Toward the ideal synthesis and transformative therapies: the roles of step economy and function oriented synthesis. <i>Tetrahedron</i> , 2013, 69, 7529-7550.	1.0	101
63	Fifteen Years of Cell-Penetrating, Guanidinium-Rich Molecular Transporters: Basic Science, Research Tools, and Clinical Applications. <i>Accounts of Chemical Research</i> , 2013, 46, 2944-2954.	7.6	270
64	Mechanistic and Computational Studies of Exocyclic Stereocontrol in the Synthesis of Bryostatin-like <i>Cis</i> -2,6-Disubstituted 4-Alkylidene tetrahydropyrans by Prins Cyclization. <i>Journal of Organic Chemistry</i> , 2013, 78, 104-115.	1.7	12
65	Lead Diversification through a Prins-Driven Macrocyclization Strategy: Application to C13-Diversified Bryostatin Analogues. <i>Synthesis</i> , 2013, 45, 1815-1824.	1.2	8
66	Effect of Histone Deacetylase Inhibitors on HIV Production in Latently Infected, Resting CD4+ T Cells From Infected Individuals Receiving Effective Antiretroviral Therapy. <i>Journal of Infectious Diseases</i> , 2012, 206, 765-769.	1.9	83
67	A molecular method for the delivery of small molecules and proteins across the cell wall of algae using molecular transporters. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2012, 109, 13225-13230.	3.3	52
68	Designed guanidinium-rich amphipathic oligocarbonate molecular transporters complex, deliver and release siRNA in cells. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2012, 109, 13171-13176.	3.3	107
69	Bryostatin analogue-induced apoptosis in mantle cell lymphoma cell lines. <i>Experimental Hematology</i> , 2012, 40, 646-656.e2.	0.2	5
70	Ligand Effects on Rates and Regioselectivities of Rh(I)-Catalyzed (5 + 2) Cycloadditions: A Computational Study of Cyclooctadiene and Dinaphthocyclooctatetraene as Ligands. <i>Journal of the American Chemical Society</i> , 2012, 134, 11012-11025.	6.6	110
71	Beyond cell penetrating peptides: designed molecular transporters. <i>Drug Discovery Today: Technologies</i> , 2012, 9, e49-e55.	4.0	47
72	Designed, synthetically accessible bryostatin analogues potently induce activation of latent HIV reservoirs in vitro. <i>Nature Chemistry</i> , 2012, 4, 705-710.	6.6	152

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73	Rhodium Dinaphthocyclooctatetraene Complexes: Synthesis, Characterization and Catalytic Activity in [5+2] Cycloadditions. <i>Angewandte Chemie - International Edition</i> , 2012, 51, 2736-2740.	7.2	72
74	Taxol-oligoarginine conjugates overcome drug resistance in-vitro in human ovarian carcinoma. <i>Gynecologic Oncology</i> , 2012, 126, 118-123.	0.6	29
75	â€œPicolog,â€œa Synthetically-Available Bryostatin Analog, Inhibits Growth of MYC-Induced Lymphoma<i>In Vivo</i>. <i>Oncotarget</i> , 2012, 3, 58-66.	0.8	37
76	Gateway synthesis of daphnane congeners and their protein kinase C affinities and cell-growth activities. <i>Nature Chemistry</i> , 2011, 3, 615-619.	6.6	77
77	Total Synthesis of Bryostatin 9. <i>Journal of the American Chemical Society</i> , 2011, 133, 9228-9231.	6.6	117
78	Function oriented synthesis: preparation and initial biological evaluation of new A-ring-modified bryologs. <i>Tetrahedron</i> , 2011, 67, 9998-10005.	1.0	17
79	Translating Natureâ€™s Library: The Bryostatins and Functionâ€œOriented Synthesis. <i>Israel Journal of Chemistry</i> , 2011, 51, 453-472.	1.0	48
80	Editorial: To Eun Lee on His 65th Birthday. <i>Chemistry - an Asian Journal</i> , 2011, 6, 1900-1900.	1.7	0
81	Design, synthesis, and evaluation of potent bryostatin analogs that modulate PKC translocation selectivity. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2011, 108, 6721-6726.	3.3	96
82	The Diene Effect: The Design, Development, and Mechanistic Investigation of Metalâ€œCatalyzed Dieneâ€œyne, Dieneâ€œene, and Dieneâ€œallene [2+2+1] Cycloaddition Reactions. <i>European Journal of Organic Chemistry</i> , 2010, 2010, 19-32.	1.2	61
83	Highly Efficient, Facile, Room Temperature Intermolecular [5 + 2] Cycloadditions Catalyzed by Cationic Rhodium(I): One Step to Cycloheptenes and Their Libraries. <i>Organic Letters</i> , 2010, 12, 1604-1607.	2.4	50
84	Electronic and Steric Control of Regioselectivities in Rh(I)-Catalyzed (5 + 2) Cycloadditions: Experiment and Theory. <i>Journal of the American Chemical Society</i> , 2010, 132, 10127-10135.	6.6	128
85	A Metal-Catalyzed Intermolecular [5+2] Cycloaddition/Nazarov Cyclization Sequence and Cascade. <i>Journal of the American Chemical Society</i> , 2010, 132, 2532-2533.	6.6	109
86	A cellular model of Alzheimer's disease therapeutic efficacy: PKC activation reverses A β -induced biomarker abnormality on cultured fibroblasts. <i>Neurobiology of Disease</i> , 2009, 34, 332-339.	2.1	64
87	A proapoptotic signaling pathway involving RasGRP, Erk, and Bim in B cells. <i>Experimental Hematology</i> , 2009, 37, 122-134.e2.	0.2	48
88	The Synthesis of Highly Substituted Cyclooctatetraene Scaffolds by Metalâ€œCatalyzed [2+2+2+2] Cycloadditions: Studies on Regioselectivity, Dynamic Properties, and Metal Chelation. <i>Angewandte Chemie - International Edition</i> , 2009, 48, 7687-7690.	7.2	53
89	Synthesis at the molecular frontier. <i>Nature</i> , 2009, 460, 197-201.	13.7	489
90	Cyclocarboamination of Alkynes with Aziridines: Synthesis of 2,3-Dihydropyrroles by a Catalyzed Formal [3 + 2] Cycloaddition. <i>Journal of the American Chemical Society</i> , 2009, 131, 7528-7529.	6.6	138

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91	Rhodium(I)-Catalyzed [2+2], [2+2+2], and [2+2+2+2] Cycloadditions of Dienes or Alkynes with a Bis-ene. <i>Organometallics</i> , 2009, 28, 5841-5844.	1.1	19
92	An Approach to the Site-Selective Diversification of Apoptolidin A with Peptide-Based Catalysts. <i>Journal of Natural Products</i> , 2009, 72, 1864-1869.	1.5	66
93	Oligocarbonate Molecular Transporters: Oligomerization-Based Syntheses and Cell-Penetrating Studies. <i>Journal of the American Chemical Society</i> , 2009, 131, 16401-16403.	6.6	112
94	Apoptolidins E and F, New Glycosylated Macrolactones Isolated from <i>Nocardiosis</i> sp.. <i>Organic Letters</i> , 2009, 11, 5474-5477.	2.4	27
95	Substituent Effects, Reactant Preorganization, and Ligand Exchange Control the Reactivity in Rh ^I -Catalyzed (5+2) Cycloadditions between Vinylcyclopropanes and Alkynes. <i>Angewandte Chemie - International Edition</i> , 2008, 47, 3939-3941.	7.2	105
96	Function-Oriented Synthesis, Step Economy, and Drug Design. <i>Accounts of Chemical Research</i> , 2008, 41, 40-49.	7.6	1,043
97	The design of guanidinium-rich transporters and their internalization mechanisms. <i>Advanced Drug Delivery Reviews</i> , 2008, 60, 452-472.	6.6	371
98	The Design, Synthesis, and Evaluation of C7 Diversified Bryostatin Analogs Reveals a Hot Spot for PKC Affinity. <i>Organic Letters</i> , 2008, 10, 3331-3334.	2.4	56
99	Practical Synthesis of Prostratin, DPP, and Their Analogs, Adjuvant Leads Against Latent HIV. <i>Science</i> , 2008, 320, 649-652.	6.0	176
100	Overcoming multidrug resistance of small-molecule therapeutics through conjugation with releasable octaarginine transporters. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2008, 105, 12128-12133.	3.3	220
101	Function-Oriented Synthesis: Biological Evaluation of Laulimalide Analogues Derived from a Last Step Cross Metathesis Diversification Strategy. <i>Molecular Pharmaceutics</i> , 2008, 5, 829-838.	2.3	24
102	Efficient Synthetic Access to a New Family of Highly Potent Bryostatin Analogues via a Prins-Driven Macrocyclization Strategy. <i>Journal of the American Chemical Society</i> , 2008, 130, 6658-6659.	6.6	137
103	Origins of Differences in Reactivities of Alkenes, Alkynes, and Allenes in [Rh(CO)2Cl] ₂ -Catalyzed (5 + 2) Cycloaddition Reactions with Vinylcyclopropanes. <i>Journal of the American Chemical Society</i> , 2008, 130, 2378-2379.	6.6	145
104	Real-time analysis of uptake and bioactivatable cleavage of luciferin-transporter conjugates in transgenic reporter mice. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2007, 104, 10340-10345.	3.3	82
105	Function-Oriented Synthesis: Studies Aimed at the Synthesis and Mode of Action of 1 [±] -Alkyldaphnane Analogues. <i>Organic Letters</i> , 2007, 9, 1829-1832.	2.4	43
106	Nickel(0)-Catalyzed [2 + 2 + 2 + 2] Cycloadditions of Terminal Diynes for the Synthesis of Substituted Cyclooctatetraenes. <i>Journal of the American Chemical Society</i> , 2007, 129, 13402-13403.	6.6	63
107	N-Alkoxyimidazolylidene Transition-Metal Complexes: Application to [5+2] and [4+2] Cycloaddition Reactions. <i>Organometallics</i> , 2007, 26, 4541-4545.	1.1	43
108	Isolation, Structure Determination, and Anti-Cancer Activity of Apoptolidin D. <i>Organic Letters</i> , 2007, 9, 691-694.	2.4	31

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109	A Computationally Designed Rh(I)-Catalyzed Two-Component [5+2+1] Cycloaddition of Ene-vinylcyclopropanes and CO for the Synthesis of Cyclooctenones. <i>Journal of the American Chemical Society</i> , 2007, 129, 10060-10061.	6.6	184
110	Intracellular Cargo Delivery by an Octaarginine Transporter Adapted to Target Prostate Cancer Cells through Cell Surface Protease Activation. <i>Bioconjugate Chemistry</i> , 2006, 17, 787-796.	1.8	65
111	Asymmetric Catalysis of the [5 + 2] Cycloaddition Reaction of Vinylcyclopropanes and $\bar{\text{I}}\text{C}$ -Systems. <i>Journal of the American Chemical Society</i> , 2006, 128, 6302-6303.	6.6	180
112	Pharmacophore Mapping in the Laulimalide Series: Total Synthesis of a Vinylogue for a Late-Stage Metathesis Diversification Strategy. <i>Organic Letters</i> , 2006, 8, 4105-4108.	2.4	29
113	Rhodium(I)-Catalyzed [4+2+2] Cycloadditions of 1,3-Dienes, Alkenes, and Alkynes for the Synthesis of Cyclooctadienes. <i>Journal of the American Chemical Society</i> , 2006, 128, 5354-5355.	6.6	55
114	Correlation of FOF1-ATPase Inhibition and Antiproliferative Activity of Apoptolidin Analogues. <i>Organic Letters</i> , 2006, 8, 589-592.	2.4	31
115	Total Synthesis and Biological Evaluation of 11-Desmethyllaulimalide, a Highly Potent Simplified Laulimalide Analogue. <i>Organic Letters</i> , 2006, 8, 1507-1510.	2.4	27
116	Laulimalide and Synthetic Laulimalide Analogues Are Synergistic with Paclitaxel and 2-Methoxyestradiol. <i>Molecular Pharmaceutics</i> , 2006, 3, 457-467.	2.3	41
117	Design, Synthesis, and Biological Evaluation of a Potent, PKC Selective, B-Ring Analog of Bryostatin. <i>Organic Letters</i> , 2006, 8, 1893-1896.	2.4	35
118	Total Synthesis and Initial Biological Evaluation of New B-Ring-Modified Bryostatin Analogs. <i>Organic Letters</i> , 2006, 8, 5299-5302.	2.4	43
119	Cyclopentadienone Synthesis by Rhodium(I)-Catalyzed [3 + 2] Cycloaddition Reactions of Cyclopropanones and Alkynes. <i>Journal of the American Chemical Society</i> , 2006, 128, 14814-14815.	6.6	137
120	Synthesis and PKC Binding of a New Class of A-Ring Diversifiable Bryostatin Analogues Utilizing a Double Asymmetric Hydrogenation and Cross-Coupling Strategy. <i>Organic Letters</i> , 2006, 8, 4581-4584.	2.4	36
121	Studies on Oxidopyrylium [5 + 2] Cycloadditions: Toward a General Synthetic Route to the C12-Hydroxy Daphnetoxins. <i>Organic Letters</i> , 2006, 8, 5373-5376.	2.4	58
122	New reactions and step economy: the total synthesis of ($\hat{\text{A}}\pm$)-salsolene oxide based on the type II transition metal-catalyzed intramolecular [4+4] cycloaddition. <i>Tetrahedron</i> , 2006, 62, 7505-7511.	1.0	194
123	Releasable Luciferin Transporter Conjugates: Tools for the Real-Time Analysis of Cellular Uptake and Release. <i>Journal of the American Chemical Society</i> , 2006, 128, 6526-6527.	6.6	136
124	Dendrimeric Molecular Transporters: Synthesis and Evaluation of Tunable Polyguanidino Dendrimers that Facilitate Cellular Uptake. <i>ChemInform</i> , 2006, 37, no.	0.1	0
125	Molecular Transporters: Synthesis of Oligoguanidinium Transporters and Their Application to Drug Delivery and Real-Time Imaging. <i>ChemBioChem</i> , 2006, 7, 1497-1515.	1.3	133
126	Metal-Catalyzed [2+2+1] Cycloadditions of 1,3-Dienes, Allenes, and CO. <i>Angewandte Chemie - International Edition</i> , 2006, 45, 2459-2462.	7.2	66

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127	Rh(I)-Catalyzed C=C Bond Activation: Seven-Membered Ring Synthesis by a [6+1] Carbonylative Ring-Expansion Reaction of Allenylcyclobutanes. <i>Angewandte Chemie - International Edition</i> , 2006, 45, 3957-3960.	7.2	82
128	Molecular Understanding of Cellular Uptake by Arginine-Rich Cell Penetrating Peptides. <i>ACS Symposium Series</i> , 2006, , 166-181.	0.5	1
129	Effect of Serum and Antioxidants on the Immunogenicity of Protein Kinase C-Activated Chronic Lymphocytic Leukemia Cells. <i>Journal of Immunotherapy</i> , 2005, 28, 28-39.	1.2	20
130	Adaptive translocation: the role of hydrogen bonding and membrane potential in the uptake of guanidinium-rich transporters into cells. <i>Advanced Drug Delivery Reviews</i> , 2005, 57, 495-504.	6.6	259
131	Rhodium(I)-Catalyzed [5+2], [6+2], and [5+2+1] Cycloadditions: New Reactions for Organic Synthesis. , 2005, , 263-299.		53
132	Multicomponent Cycloadditions: The Four-Component [5 + 1 + 2 + 1] Cycloaddition of Vinylcyclopropanes, Alkynes, and CO.. <i>ChemInform</i> , 2005, 36, no.	0.1	0
133	Rh(I)-catalyzed cleavage of unactivated C=C—O bonds — Carbonylative rearrangement reactions of allenyl ethers to 2-carboalkoxy-1,3-dienes. <i>Canadian Journal of Chemistry</i> , 2005, 83, 838-842.	0.6	8
134	Role of the A-Ring of Bryostatin Analogues in PKC Binding: — Synthesis and Initial Biological Evaluation of New A-Ring-Modified Bryologs. <i>Organic Letters</i> , 2005, 7, 1995-1998.	2.4	24
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