

Paul A Wender

List of Publications by Year in Descending Order

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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.

232
papers

18,451
citations

77
h-index

128
g-index

262
ext. papers

19,768
ext. citations

10.7
avg, IF

6.84
L-index

#	Paper	IF	Citations
232	Latency reversal plus natural killer cells diminish HIV reservoir in vivo.. <i>Nature Communications</i> , 2022 , 13, 121	17.4	3
231	An mRNA SARS-CoV-2 vaccine employing Charge-Altering Releasable Transporters with a TLR-9 agonist induces neutralizing antibodies and T cell memory 2021 ,		3
230	Designed PKC-targeting bryostatin analogs modulate innate immunity and neuroinflammation. <i>Cell Chemical Biology</i> , 2021 , 28, 537-545.e4	8.2	2
229	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	16.8	8
228	Targeting of Escherichia coli with Vancomycin-Arginine. <i>Antimicrobial Agents and Chemotherapy</i> , 2021 , 65,	5.9	6
227	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	11.5	13
226	In Situ Detection of Endogenous HIV Activation by Dynamic Nuclear Polarization NMR and Flow Cytometry. <i>International Journal of Molecular Sciences</i> , 2020 , 21,	6.3	8
225	Synthesis and mechanistic investigations of pH-responsive cationic poly(aminoester)s. <i>Chemical Science</i> , 2020 , 11, 2951-2966	9.4	8
224	Tracking HIV Rebound following Latency Reversal Using Barcoded HIV. <i>Cell Reports Medicine</i> , 2020 , 1, 100162	18	4
223	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	5.7	9
222	Function-Oriented Synthesis: Design, Synthesis, and Evaluation of Highly Simplified Bryostatin Analogues. <i>Journal of Organic Chemistry</i> , 2020 , 85, 15116-15128	4.2	5
221	Charge-altering releasable transporters enable phenotypic manipulation of natural killer cells for cancer immunotherapy. <i>Blood Advances</i> , 2020 , 4, 4244-4255	7.8	12
220	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	11.6	10
219	Synthesis and evaluation of designed PKC modulators for enhanced cancer immunotherapy. <i>Nature Communications</i> , 2020 , 11, 1879	17.4	14
218	Vancomycin-Arginine Conjugate Inhibits Growth of Carbapenem-Resistant and Targets Cell-Wall Synthesis. <i>ACS Chemical Biology</i> , 2019 , 14, 2065-2070	4.9	28
217	Synthesis of Modified Nucleoside Oligophosphates Simplified: Fast, Pure, and Protecting Group Free. <i>Journal of the American Chemical Society</i> , 2019 , 141, 15013-15017	16.4	14
216	Local Delivery of , , and mRNA Kindles Global Anticancer Immunity. <i>Cancer Research</i> , 2019 , 79, 1624-1634	10.1	50

215	A Phosphoramidite Analogue of Cyclotriphosphate Enables Iterative Polyphosphorylations. <i>Angewandte Chemie - International Edition</i> , 2019 , 58, 3928-3933	16.4	16
214	A Phosphoramidite Analogue of Cyclotriphosphate Enables Iterative Polyphosphorylations. <i>Angewandte Chemie</i> , 2019 , 131, 3968-3973	3.6	5
213	Oligo(serine ester) Charge-Altering Releasable Transporters: Organocatalytic Ring-Opening Polymerization and their Use for in Vitro and in Vivo mRNA Delivery. <i>Journal of the American Chemical Society</i> , 2019 , 141, 8416-8421	16.4	34
212	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	7	7
211	Reversible RNA acylation for control of CRISPR-Cas9 gene editing. <i>Chemical Science</i> , 2019 , 11, 1011-1016	9.4	22
210	REDOR NMR Reveals Multiple Conformers for a Protein Kinase C Ligand in a Membrane Environment. <i>ACS Central Science</i> , 2018 , 4, 89-96	16.8	24
209	Functional DNA Delivery Enabled by Lipid-Modified Charge-Altering Releasable Transporters (CARTs). <i>Biomacromolecules</i> , 2018 , 19, 2812-2824	6.9	19
208	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	7	3
207	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	11.5	101
206	Delivery of Inorganic Polyphosphate into Cells Using Amphipathic Oligocarbonate Transporters. <i>ACS Central Science</i> , 2018 , 4, 1394-1402	16.8	11
205	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	16.4	65
204	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	11.5	60
203	Characterization of designed, synthetically accessible bryostatin analog HIV latency reversing agents. <i>Virology</i> , 2018 , 520, 83-93	3.6	26
202	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	11.5	137
201	Molecular dynamics simulations reveal ligand-controlled positioning of a peripheral protein complex in membranes. <i>Nature Communications</i> , 2017 , 8, 6	17.4	70
200	Vault Nanoparticles: Chemical Modifications for Imaging and Enhanced Delivery. <i>ACS Nano</i> , 2017 , 11, 872-881	16.7	23
199	Scalable synthesis of bryostatin 1 and analogs, adjuvant leads against latent HIV. <i>Science</i> , 2017 , 358, 218-223	3.9	64
198	Ynol Ethers as Ketene Equivalents in Rhodium-Catalyzed Intermolecular [5 + 2] Cycloaddition Reactions. <i>Organic Letters</i> , 2017 , 19, 5810-5813	6.2	17

197	In vivo activation of latent HIV with a synthetic bryostatin analog effects both latent cell "kick" and "kill" in strategy for virus eradication. <i>PLoS Pathogens</i> , 2017 , 13, e1006575	7.6	55
196	Retrosynthetic Reaction Prediction Using Neural Sequence-to-Sequence Models. <i>ACS Central Science</i> , 2017 , 3, 1103-1113	16.8	173
195	Combinations of isoform-targeted histone deacetylase inhibitors and bryostatin analogues display remarkable potency to activate latent HIV without global T-cell activation. <i>Scientific Reports</i> , 2017 , 7, 7456	4.9	23
194	Cellular delivery and photochemical release of a caged inositol-pyrophosphate induces PH-domain translocation in cellulose. <i>Nature Communications</i> , 2016 , 7, 10622	17.4	62
193	Simplified Bryostatin Analogues Protect Cells from Chikungunya Virus-Induced Cell Death. <i>Journal of Natural Products</i> , 2016 , 79, 675-9	4.9	16
192	Inhibition of Chikungunya Virus-Induced Cell Death by Salicylate-Derived Bryostatin Analogues Provides Additional Evidence for a PKC-Independent Pathway. <i>Journal of Natural Products</i> , 2016 , 79, 680-4	4.9	26
191	Cell-Penetrating, Guanidinium-Rich Oligophosphoesters: Effective and Versatile Molecular Transporters for Drug and Probe Delivery. <i>Journal of the American Chemical Society</i> , 2016 , 138, 3510-7	16.4	82
190	Bioorthogonal Catalysis: A General Method To Evaluate Metal-Catalyzed Reactions in Real Time in Living Systems Using a Cellular Luciferase Reporter System. <i>Bioconjugate Chemistry</i> , 2016 , 27, 376-82	6.3	49
189	Comparative analysis of the anti-chikungunya virus activity of novel bryostatin analogs confirms the existence of a PKC-independent mechanism. <i>Biochemical Pharmacology</i> , 2016 , 120, 15-21	6	11
188	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. <i>Journal of the American Chemical Society</i> , 2015 , 137, 3678-85	16.4	24
187	Tetramethyleneethane Equivalents: Recursive Reagents for Serialized Cycloadditions. <i>Journal of the American Chemical Society</i> , 2015 , 137, 9088-93	16.4	30
186	Carbonyl(chloro)bis(triphenylphosphine)rhodium(I) 2015 , 1-22		0
185	Function through synthesis-informed design. <i>Accounts of Chemical Research</i> , 2015 , 48, 752-60	24.3	54
184	Studies on the regio- and diastereo-selective epoxidation of daphnanes and tiglanes. <i>Tetrahedron Letters</i> , 2015 , 56, 3423-3427	2	9
183	Guanidinium-rich, glycerol-derived oligocarbonates: a new class of cell-penetrating molecular transporters that complex, deliver, and release siRNA. <i>Molecular Pharmaceutics</i> , 2015 , 12, 742-50	5.6	17
182	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. <i>ACS Catalysis</i> , 2015 , 5, 1758-1763	13.1	29
181	Rethinking the Role of Natural Products: Function-Oriented Synthesis, Bryostatin, and Bryologs. <i>Methods and Principles in Medicinal Chemistry</i> , 2014 , 473-544	0.4	8
180	Structural complexity through multicomponent cycloaddition cascades enabled by dual-purpose, reactivity regenerating 1,2,3-triene equivalents. <i>Nature Chemistry</i> , 2014 , 6, 448-52	17.6	53

179	Improved protein kinase C affinity through final step diversification of a simplified salicylate-derived bryostatin analog scaffold. <i>Organic Letters</i> , 2014 , 16, 5140-3	6.2	20
178	Cell-penetrating, guanidinium-rich molecular transporters for overcoming efflux-mediated multidrug resistance. <i>Molecular Pharmaceutics</i> , 2014 , 11, 2553-65	5.6	44
177	Computer-guided design, synthesis, and protein kinase C affinity of a new salicylate-based class of bryostatin analogs. <i>Organic Letters</i> , 2014 , 16, 5136-9	6.2	30
176	Bioengineered vaults: self-assembling protein shell-lipophilic core nanoparticles for drug delivery. <i>ACS Nano</i> , 2014 , 8, 7723-32	16.7	49
175	Toward the ideal synthesis and molecular function through synthesis-informed design. <i>Natural Product Reports</i> , 2014 , 31, 433-40	15.1	157
174	Propargyltrimethylsilanes as allene equivalents in transition metal-catalyzed [5 + 2] cycloadditions. <i>Organic Letters</i> , 2014 , 16, 2923-5	6.2	21
173	Function through bio-inspired, synthesis-informed design: step-economical syntheses of designed kinase inhibitors. Dedicated to Max Malacria, a friend and scholar whose science and creative contributions to step-economical synthesis have inspired us all and moved the field closer to the ideal.	5.2	5
172	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. <i>Journal of the American Chemical Society</i> , 2014 , 136, 17273-83	16.4	88
171	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, 11698-703	11.5	109
170	Toward the Ideal Synthesis and Transformative Therapies: The Roles of Step Economy and Function Oriented Synthesis. <i>Tetrahedron</i> , 2013 , 69, 7529-7550	2.4	87
169	Fifteen years of cell-penetrating, guanidinium-rich molecular transporters: basic science, research tools, and clinical applications. <i>Accounts of Chemical Research</i> , 2013 , 46, 2944-54	24.3	230
168	Mechanistic and computational studies of exocyclic stereocontrol in the synthesis of bryostatin-like cis-2,6-disubstituted 4-alkylidenetetrahydropyrans by Prins cyclization. <i>Journal of Organic Chemistry</i> , 2013 , 78, 104-15	4.2	12
167	Lead Diversification through a Prins-Driven Macrocyclization Strategy: Application to C13-Diversified Bryostatin Analogues. <i>Synthesis</i> , 2013 , 45, 1815-1824	2.9	8
166	Taxol-oligoarginine conjugates overcome drug resistance in-vitro in human ovarian carcinoma. <i>Gynecologic Oncology</i> , 2012 , 126, 118-23	4.9	23
165	Bryostatin analogue-induced apoptosis in mantle cell lymphoma cell lines. <i>Experimental Hematology</i> , 2012 , 40, 646-56.e2	3.1	5
164	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, 11012-25	16.4	106
163	Beyond Cell Penetrating Peptides: Designed Molecular Transporters. <i>Drug Discovery Today: Technologies</i> , 2012 , 9, e49-e55	7.1	40
162	Designed, synthetically accessible bryostatin analogues potently induce activation of latent HIV reservoirs in vitro. <i>Nature Chemistry</i> , 2012 , 4, 705-10	17.6	131

161	Rhodium Dinaphthocyclooctatetraene Complexes: Synthesis, Characterization and Catalytic Activity in [5+2] Cycloadditions. <i>Angewandte Chemie</i> , 2012 , 124, 2790-2794	3.6	17
160	Rhodium dinaphthocyclooctatetraene complexes: synthesis, characterization and catalytic activity in [5+2] cycloadditions. <i>Angewandte Chemie - International Edition</i> , 2012 , 51, 2736-40	16.4	64
159	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, 765-9	7	71
158	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, 13225-30	11.5	45
157	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, 13171-6	11.5	99
156	"Picolog," a synthetically-available bryostatin analog, inhibits growth of MYC-induced lymphoma in vivo. <i>Oncotarget</i> , 2012 , 3, 58-66	3.3	33
155	Function Oriented Synthesis: Preparation and Initial Biological Evaluation of New A-Ring-Modified Bryologs. <i>Tetrahedron</i> , 2011 , 67, 9998-10005	2.4	16
154	Translating Nature [®] Library: The Bryostatins and Function-Oriented Synthesis. <i>Israel Journal of Chemistry</i> , 2011 , 51, 453-472	3.4	47
153	Gateway synthesis of daphnane congeners and their protein kinase C affinities and cell-growth activities. <i>Nature Chemistry</i> , 2011 , 3, 615-9	17.6	68
152	Total synthesis of bryostatin 9. <i>Journal of the American Chemical Society</i> , 2011 , 133, 9228-31	16.4	112
151	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, 6721-6	11.5	92
150	The Preparation of Cyclohept-4-Enones by Rhodium-Catalyzed Intermolecular [5+2] Cycloaddition 2011 , 109-120		4
149	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, 1604-7	6.2	48
148	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, 10127-35	16.4	120
147	A metal-catalyzed intermolecular [5+2] cycloaddition/Nazarov cyclization sequence and cascade. <i>Journal of the American Chemical Society</i> , 2010 , 132, 2532-3	16.4	103
146	The Diene Effect: The Design, Development, and Mechanistic Investigation of Metal-Catalyzed Diene-yne, Diene-ene, and Diene-allene [2+2+1] Cycloaddition Reactions. <i>European Journal of Organic Chemistry</i> , 2010 , 2010, 19-32	3.2	56
145	A cellular model of Alzheimer [®] disease therapeutic efficacy: PKC activation reverses Abeta-induced biomarker abnormality on cultured fibroblasts. <i>Neurobiology of Disease</i> , 2009 , 34, 332-9	7.5	58
144	A proapoptotic signaling pathway involving RasGRP, Erk, and Bim in B cells. <i>Experimental Hematology</i> , 2009 , 37, 122-134	3.1	36

143	The synthesis of highly substituted cyclooctatetraene scaffolds by metal-catalyzed [2+2+2+2] cycloadditions: studies on regioselectivity, dynamic properties, and metal chelation. <i>Angewandte Chemie - International Edition</i> , 2009 , 48, 7687-90	16.4	49
142	Synthesis at the molecular frontier. <i>Nature</i> , 2009 , 460, 197-201	50.4	424
141	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, 7528-9	16.4	125
140	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	3.8	19
139	An approach to the site-selective diversification of apoptolidin A with peptide-based catalysts. <i>Journal of Natural Products</i> , 2009 , 72, 1864-9	4.9	61
138	Oligocarbonate molecular transporters: oligomerization-based syntheses and cell-penetrating studies. <i>Journal of the American Chemical Society</i> , 2009 , 131, 16401-3	16.4	108
137	Apoptolidins E and F, new glycosylated macrolactones isolated from <i>Nocardiosis</i> sp. <i>Organic Letters</i> , 2009 , 11, 5474-7	6.2	24
136	The design, synthesis, and evaluation of C7 diversified bryostatin analogs reveals a hot spot for PKC affinity. <i>Organic Letters</i> , 2008 , 10, 3331-4	6.2	52
135	Practical synthesis of prostratin, DPP, and their analogs, adjuvant leads against latent HIV. <i>Science</i> , 2008 , 320, 649-52	33.3	151
134	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, 12128-33	11.5	196
133	Function-oriented synthesis: biological evaluation of laulimalide analogues derived from a last step cross metathesis diversification strategy. <i>Molecular Pharmaceutics</i> , 2008 , 5, 829-38	5.6	24
132	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, 6658-9	16.4	134
131	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, 2378-9	16.4	142
130	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, 3939-41	16.4	101
129	Function-oriented synthesis, step economy, and drug design. <i>Accounts of Chemical Research</i> , 2008 , 41, 40-9	24.3	931
128	The design of guanidinium-rich transporters and their internalization mechanisms. <i>Advanced Drug Delivery Reviews</i> , 2008 , 60, 452-72	18.5	344
127	N-Alkoxyimidazolylidene Transition-Metal Complexes: Application to [5+2] and [4+2] Cycloaddition Reactions. <i>Organometallics</i> , 2007 , 26, 4541-4545	3.8	42
126	Isolation, structure determination, and anti-cancer activity of apoptolidin D. <i>Organic Letters</i> , 2007 , 9, 691-4	6.2	28

125	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, 10060-1	16.4	172
124	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, 10340-5	11.5	77
123	Function-oriented synthesis: studies aimed at the synthesis and mode of action of 1 α -alkyldaphnane analogues. <i>Organic Letters</i> , 2007 , 9, 1829-32	6.2	39
122	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, 13402-3	16.4	57
121	Molecular transporters: synthesis of oligoguanidinium transporters and their application to drug delivery and real-time imaging. <i>ChemBioChem</i> , 2006 , 7, 1497-515	3.8	128
120	Metal-catalyzed [2+2+1] cycloadditions of 1,3-dienes, allenes, and CO. <i>Angewandte Chemie - International Edition</i> , 2006 , 45, 2459-62	16.4	62
119	RhI-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, 3957-60	16.4	76
118	Metal-Catalyzed [2+2+1] Cycloadditions of 1,3-Dienes, Allenes, and CO. <i>Angewandte Chemie</i> , 2006 , 118, 2519-2522	3.6	13
117	RhI-Catalyzed C-C Bond Activation: Seven-Membered Ring Synthesis by a [6+1] Carbonylative Ring-Expansion Reaction of Allenylcyclobutanes. <i>Angewandte Chemie</i> , 2006 , 118, 4061-4064	3.6	27
116	Molecular Understanding of Cellular Uptake by Arginine-Rich Cell Penetrating Peptides. <i>ACS Symposium Series</i> , 2006 , 166-181	0.4	1
115	Intracellular cargo delivery by an octaarginine transporter adapted to target prostate cancer cells through cell surface protease activation. <i>Bioconjugate Chemistry</i> , 2006 , 17, 787-96	6.3	61
114	Asymmetric catalysis of the [5 + 2] cycloaddition reaction of vinylcyclopropanes and pi-systems. <i>Journal of the American Chemical Society</i> , 2006 , 128, 6302-3	16.4	173
113	Pharmacophore mapping in the laulimalide series: total synthesis of a vinylogue for a late-stage metathesis diversification strategy. <i>Organic Letters</i> , 2006 , 8, 4105-8	6.2	27
112	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, 5354-5	16.4	53
111	Correlation of F0F1-ATPase inhibition and antiproliferative activity of apoptolidin analogues. <i>Organic Letters</i> , 2006 , 8, 589-92	6.2	27
110	Total synthesis and biological evaluation of 11-desmethyllaulimalide, a highly potent simplified laulimalide analogue. <i>Organic Letters</i> , 2006 , 8, 1507-10	6.2	27
109	Laulimalide and synthetic laulimalide analogues are synergistic with paclitaxel and 2-methoxyestradiol. <i>Molecular Pharmaceutics</i> , 2006 , 3, 457-67	5.6	40
108	Design, synthesis, and biological evaluation of a potent, PKC selective, B-ring analog of bryostatin. <i>Organic Letters</i> , 2006 , 8, 1893-6	6.2	35

107	Total synthesis and initial biological evaluation of new B-ring-modified bryostatin analogs. <i>Organic Letters</i> , 2006 , 8, 5299-302	6.2	42
106	Cyclopentadienone synthesis by rhodium(I)-catalyzed [3 + 2] cycloaddition reactions of cyclopropenones and alkynes. <i>Journal of the American Chemical Society</i> , 2006 , 128, 14814-5	16.4	115
105	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, 4581-4	6.2	35
104	Studies on oxidopyrylium [5+2] cycloadditions: toward a general synthetic route to the C12-hydroxydaphnetoxins. <i>Organic Letters</i> , 2006 , 8, 5373-6	6.2	57
103	New reactions and step economy: the total synthesis of (±)-salsolene oxide based on the type II transition metal-catalyzed intramolecular [4+4] cycloaddition. <i>Tetrahedron</i> , 2006 , 62, 7505-7511	2.4	177
102	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, 6526-7	16.4	127
101	Rh(I)-catalyzed cleavage of unactivated 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	0.9	5
100	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, 1995-8	6.2	24
99	Transition metal-catalyzed intermolecular [5+2] and [5+2+1] cycloadditions of allenes and vinylcyclopropanes. <i>Journal of the American Chemical Society</i> , 2005 , 127, 6530-1	16.4	149
98	Identification of a tunable site in bryostatin analogs: C20 Bryologs through late stage diversification. <i>Organic Letters</i> , 2005 , 7, 1177-80	6.2	24
97	Apoptolidins B and C: isolation, structure determination, and biological activity. <i>Organic Letters</i> , 2005 , 7, 3025-8	6.2	40
96	Dendrimeric molecular transporters: synthesis and evaluation of tunable polyguanidino dendrimers that facilitate cellular uptake. <i>Organic Letters</i> , 2005 , 7, 4815-8	6.2	58
95	Late-stage intermolecular CH activation for lead diversification: a highly chemoselective oxyfunctionalization of the C-9 position of potent bryostatin analogues. <i>Organic Letters</i> , 2005 , 7, 79-82	6.2	92
94	Multicomponent cycloadditions: the four-component [5+1+2+1] cycloaddition of vinylcyclopropanes, alkynes, and CO. <i>Journal of the American Chemical Society</i> , 2005 , 127, 2836-7	16.4	136
93	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	5	20
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