Armen Zakarian

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.

75	2,391	28	47
papers	citations	h-index	g-index
80	2,718 ext. citations	10.8	5.29
ext. papers		avg, IF	L-index

#	Paper	IF	Citations
75	Scalable Total Synthesis, IP3R Inhibitory Activity of Desmethylxestospongin B, and Effect on Mitochondrial Function and Cancer Cell Survival. <i>Angewandte Chemie - International Edition</i> , 2021 , 60, 11278-11282	16.4	1
74	Scalable Total Synthesis, IP3R Inhibitory Activity of Desmethylxestospongin B, and Effect on Mitochondrial Function and Cancer Cell Survival. <i>Angewandte Chemie</i> , 2021 , 133, 11378-11382	3.6	0
73	First evidence that emerging pinnatoxin-G, a contaminant of shellfish, reaches the brain and crosses the placental barrier. <i>Science of the Total Environment</i> , 2021 , 790, 148125	10.2	1
7 ²	Stereoselective \oplus -Tertiary Alkylation of -(Arylacetyl)oxazolidinones. <i>Synlett</i> , 2020 , 31, 683-686	2.2	2
71	Concise Synthesis of (+)-[13C4]-Anatoxin-a by Dynamic Kinetic Resolution of a Cyclic Iminium Ion. <i>Angewandte Chemie</i> , 2020 , 132, 11460-11464	3.6	
7°	Cancer cells with defective oxidative phosphorylation require endoplasmic reticulum-to-mitochondria Ca transfer for survival. <i>Science Signaling</i> , 2020 , 13,	8.8	23
69	Concise Synthesis of (+)-[C]-Anatoxin-a by Dynamic Kinetic Resolution of a Cyclic Iminium Ion. <i>Angewandte Chemie - International Edition</i> , 2020 , 59, 11364-11368	16.4	2
68	Enantioselective Alkylation of 2-Alkylpyridines Controlled by Organolithium Aggregation. <i>Journal of the American Chemical Society</i> , 2019 , 141, 15024-15028	16.4	10
67	Synthetic Pinnatoxins A and G Reversibly Block Mouse Skeletal Neuromuscular Transmission In Vivo and In Vitro. <i>Marine Drugs</i> , 2019 , 17,	6	8
66	Direct Enantioselective and Regioselective Alkylation of IDInsaturated Carboxylic Acids with Chiral Lithium Amides as Traceless Auxiliaries. <i>Organic Letters</i> , 2019 , 21, 1930-1934	6.2	5
65	Total Synthesis of Covalent Cysteine Protease Inhibitor N-Desmethyl Thalassospiramide C and Crystallographic Evidence for Its Mode of Action. <i>Organic Letters</i> , 2019 , 21, 508-512	6.2	3
64	Short Total Synthesis of [N]-Cylindrospermopsins from NHCl Enables Precise Quantification of Freshwater Cyanobacterial Contamination. <i>Journal of the American Chemical Society</i> , 2018 , 140, 6027-6	03 ^{6.4}	19
63	Stereodivergence in the Ireland-Claisen Rearrangement of ⊞-Alkoxy Esters. <i>Organic Letters</i> , 2018 , 20, 4867-4870	6.2	10
62	Total Synthesis of (+)-Guadinomic Acid via Hydroxyl-Directed Guanidylation. <i>Journal of Organic Chemistry</i> , 2018 , 83, 9492-9496	4.2	6
61	Canvass: A Crowd-Sourced, Natural-Product Screening Library for Exploring Biological Space. <i>ACS Central Science</i> , 2018 , 4, 1727-1741	16.8	26
60	Direkte asymmetrische Alkylierung von Ketonen: noch immer ein unerreichtes Ziel. <i>Angewandte Chemie</i> , 2017 , 129, 9406-9418	3.6	20
59	Direct Asymmetric Alkylation of Ketones: Still Unconquered. <i>Angewandte Chemie - International Edition</i> , 2017 , 56, 9278-9290	16.4	52

(2015-2017)

58	Determination of the Absolute Configuration of Echiral Primary Alcohols Using the Competing Enantioselective Conversion Method. <i>Organic Letters</i> , 2017 , 19, 2953-2956	6.2	14
57	Antidiabetic Disruptors of the Glucokinase-Glucokinase Regulatory Protein Complex Reorganize a Coulombic Interface. <i>Biochemistry</i> , 2017 , 56, 3150-3157	3.2	5
56	Cyclic imine toxins from dinoflagellates: a growing family of potent antagonists of the nicotinic acetylcholine receptors. <i>Journal of Neurochemistry</i> , 2017 , 142 Suppl 2, 41-51	6	39
55	Lithium Enolates in the Enantioselective Construction of Tetrasubstituted Carbon Centers with Chiral Lithium Amides as Noncovalent Stereodirecting Auxiliaries. <i>Journal of the American Chemical Society</i> , 2017 , 139, 527-533	16.4	32
54	Total Synthesis of Unsymmetrically Oxidized Nuphar Thioalkaloids via Copper-Catalyzed Thiolane Assembly. <i>Journal of the American Chemical Society</i> , 2017 , 139, 13272-13275	16.4	24
53	Synthesis of Functionalized Dihydrobenzofurans by Direct Aryl CD Bond Formation under Mild Conditions. <i>Angewandte Chemie</i> , 2016 , 128, 11797-11800	3.6	7
52	Synthesis of Functionalized Dihydrobenzofurans by Direct Aryl C-O Bond Formation under Mild Conditions. <i>Angewandte Chemie - International Edition</i> , 2016 , 55, 11625-11628	16.4	23
51	Mixed Aggregates of the Dilithiated Koga Tetraamine: NMR Spectroscopic and Computational Studies. <i>Angewandte Chemie - International Edition</i> , 2016 , 55, 10093-7	16.4	6
50	Stereoselective Synthesis of Cyclic Guanidines by Directed Diamination of Unactivated Alkenes. <i>Organic Letters</i> , 2016 , 18, 5532-5535	6.2	15
49	Mixed Aggregates of the Dilithiated Koga Tetraamine: NMR Spectroscopic and Computational Studies. <i>Angewandte Chemie</i> , 2016 , 128, 10247-10251	3.6	3
49		3.6	
	Studies. <i>Angewandte Chemie</i> , 2016 , 128, 10247-10251 Cutting-Edge and Time-Honored Strategies for Stereoselective Construction of C-N Bonds in Total		
48	Studies. <i>Angewandte Chemie</i> , 2016 , 128, 10247-10251 Cutting-Edge and Time-Honored Strategies for Stereoselective Construction of C-N Bonds in Total Synthesis. <i>Chemical Reviews</i> , 2016 , 116, 4441-557		107
48	Studies. <i>Angewandte Chemie</i> , 2016 , 128, 10247-10251 Cutting-Edge and Time-Honored Strategies for Stereoselective Construction of C-N Bonds in Total Synthesis. <i>Chemical Reviews</i> , 2016 , 116, 4441-557 Spirolides and Cyclic Imines: Toxicological Profile 2016 , 193-217 Marine Macrocyclic Imines, Pinnatoxins A and G: Structural Determinants and Functional Properties	68.1	107
48 47 46	Cutting-Edge and Time-Honored Strategies for Stereoselective Construction of C-N Bonds in Total Synthesis. <i>Chemical Reviews</i> , 2016 , 116, 4441-557 Spirolides and Cyclic Imines: Toxicological Profile 2016 , 193-217 Marine Macrocyclic Imines, Pinnatoxins A and G: Structural Determinants and Functional Properties to Distinguish Neuronal \Box 7 from Muscle \Box 1(2) \Box 1 hAChRs. <i>Structure</i> , 2015 , 23, 1106-15 Toward the Synthesis of Nuphar Sesquiterpene Thioalkaloids: Stereodivergent Rhodium-Catalyzed	68.1 5.2	107 2 35
48 47 46 45	Cutting-Edge and Time-Honored Strategies for Stereoselective Construction of C-N Bonds in Total Synthesis. <i>Chemical Reviews</i> , 2016 , 116, 4441-557 Spirolides and Cyclic Imines: Toxicological Profile 2016 , 193-217 Marine Macrocyclic Imines, Pinnatoxins A and G: Structural Determinants and Functional Properties to Distinguish Neuronal #7 from Muscle #1(2)IhAChRs. <i>Structure</i> , 2015 , 23, 1106-15 Toward the Synthesis of Nuphar Sesquiterpene Thioalkaloids: Stereodivergent Rhodium-Catalyzed Synthesis of the Thiolane Subunit. <i>Journal of Organic Chemistry</i> , 2015 , 80, 7581-9 Total synthesis and structural revision of (+)-muironolide a. <i>Journal of the American Chemical</i>	68.1 5.2 4.2	107 2 35 25
48 47 46 45 44	Cutting-Edge and Time-Honored Strategies for Stereoselective Construction of C-N Bonds in Total Synthesis. <i>Chemical Reviews</i> , 2016 , 116, 4441-557 Spirolides and Cyclic Imines: Toxicological Profile 2016 , 193-217 Marine Macrocyclic Imines, Pinnatoxins A and G: Structural Determinants and Functional Properties to Distinguish Neuronal #7 from Muscle #1(2)IhAChRs. <i>Structure</i> , 2015 , 23, 1106-15 Toward the Synthesis of Nuphar Sesquiterpene Thioalkaloids: Stereodivergent Rhodium-Catalyzed Synthesis of the Thiolane Subunit. <i>Journal of Organic Chemistry</i> , 2015 , 80, 7581-9 Total synthesis and structural revision of (+)-muironolide a. <i>Journal of the American Chemical Society</i> , 2015 , 137, 5907-10 Toward the Synthesis of Muironolide A: Synthesis and Structure of Heteroleptic LanthanideTerpyridine Complexes with 2-Oxo Amides. <i>European Journal of Organic Chemistry</i> ,	68.1 5.2 4.2 16.4	107 2 35 25 27

40	10-Step Asymmetric Total Synthesis and Stereochemical Elucidation of (+)-Dragmacidin D. <i>Angewandte Chemie - International Edition</i> , 2015 , 54, 9971-5	16.4	20
39	10-Step Asymmetric Total Synthesis and Stereochemical Elucidation of (+)-Dragmacidin D. <i>Angewandte Chemie</i> , 2015 , 127, 10109-10113	3.6	8
38	Direct enantioselective conjugate addition of carboxylic acids with chiral lithium amides as traceless auxiliaries. <i>Journal of the American Chemical Society</i> , 2015 , 137, 656-9	16.4	33
37	Spirolides and Cyclic Imines: Toxicological Profile 2015 , 1-19		2
36	Asymmetric radical addition of TEMPO to titanium enolates. Organic Letters, 2014, 16, 516-9	6.2	32
35	Enantioselective synthesis of (-)-maoecrystal V by enantiodetermining C-H functionalization. <i>Journal of the American Chemical Society</i> , 2014 , 136, 17738-49	16.4	85
34	Stereoselective <code>\(\properature \) -fluorination of N-acyloxazolidinones at room temperature within 1 h. \(\textit{Journal of Organic Chemistry, } \) 2014, 79, 6206-20</code>	4.2	23
33	Total synthesis of maoecrystal V: early-stage C-H functionalization and lactone assembly by radical cyclization. <i>Journal of the American Chemical Society</i> , 2013 , 135, 14552-5	16.4	107
32	Pinnatoxin G is responsible for atypical toxicity in mussels (Mytilus galloprovincialis) and clams (Venerupis decussata) from Ingril, a French Mediterranean lagoon. <i>Toxicon</i> , 2013 , 75, 16-26	2.8	59
31	Radical Haloalkylation 2013 , 285-291		2
31	Radical Haloalkylation 2013, 285-291 Enantioselective synthesis of tatanans A-C and reinvestigation of their glucokinase-activating properties. <i>Nature Chemistry</i> , 2013, 5, 410-6	17.6	39
	Enantioselective synthesis of tatanans A-C and reinvestigation of their glucokinase-activating	17.6 6.2	
30	Enantioselective synthesis of tatanans A-C and reinvestigation of their glucokinase-activating properties. <i>Nature Chemistry</i> , 2013 , 5, 410-6 An efficient synthesis of the fully elaborated isoindolinone unit of muironolide A. <i>Organic Letters</i> ,	6.2	39 15
30	Enantioselective synthesis of tatanans A-C and reinvestigation of their glucokinase-activating properties. <i>Nature Chemistry</i> , 2013 , 5, 410-6 An efficient synthesis of the fully elaborated isoindolinone unit of muironolide A. <i>Organic Letters</i> , 2013 , 15, 3314-7 Enediolate-dilithium amide mixed aggregates in the enantioselective alkylation of arylacetic acids:	6.2	39 15
30 29 28	Enantioselective synthesis of tatanans A-C and reinvestigation of their glucokinase-activating properties. <i>Nature Chemistry</i> , 2013 , 5, 410-6 An efficient synthesis of the fully elaborated isoindolinone unit of muironolide A. <i>Organic Letters</i> , 2013 , 15, 3314-7 Enediolate-dilithium amide mixed aggregates in the enantioselective alkylation of arylacetic acids: structural studies and a stereochemical model. <i>Journal of the American Chemical Society</i> , 2013 , 135, 168 Stability of cyclic imine toxins: interconversion of pinnatoxin amino ketone and pinnatoxin A in	6.2 5 3 - 6 4	39 15 35
30 29 28 27	Enantioselective synthesis of tatanans A-C and reinvestigation of their glucokinase-activating properties. <i>Nature Chemistry</i> , 2013 , 5, 410-6 An efficient synthesis of the fully elaborated isoindolinone unit of muironolide A. <i>Organic Letters</i> , 2013 , 15, 3314-7 Enediolate-dilithium amide mixed aggregates in the enantioselective alkylation of arylacetic acids: structural studies and a stereochemical model. <i>Journal of the American Chemical Society</i> , 2013 , 135, 168 Stability of cyclic imine toxins: interconversion of pinnatoxin amino ketone and pinnatoxin A in aqueous media. <i>Journal of Organic Chemistry</i> , 2012 , 77, 10435-40 A simple method for asymmetric trifluoromethylation of N-acyl oxazolidinones via Ru-catalyzed	6.2 53-64 4.2	39 15 35 21
30 29 28 27 26	Enantioselective synthesis of tatanans A-C and reinvestigation of their glucokinase-activating properties. <i>Nature Chemistry</i> , 2013 , 5, 410-6 An efficient synthesis of the fully elaborated isoindolinone unit of muironolide A. <i>Organic Letters</i> , 2013 , 15, 3314-7 Enediolate-dilithium amide mixed aggregates in the enantioselective alkylation of arylacetic acids: structural studies and a stereochemical model. <i>Journal of the American Chemical Society</i> , 2013 , 135, 168 Stability of cyclic imine toxins: interconversion of pinnatoxin amino ketone and pinnatoxin A in aqueous media. <i>Journal of Organic Chemistry</i> , 2012 , 77, 10435-40 A simple method for asymmetric trifluoromethylation of N-acyl oxazolidinones via Ru-catalyzed radical addition to zirconium enolates. <i>Journal of the American Chemical Society</i> , 2012 , 134, 6976-9 Studies toward the synthesis of spirolide C: exploration into the formation of the 23-membered	6.2 53-64 4.2 16.4	39 15 35 21 119

(2006-2011)

22	Dual Ti R u Catalysis in the Direct Radical Haloalkylation of N-Acyl Oxazolidinones. <i>Angewandte Chemie</i> , 2011 , 123, 7274-7277	3.6	19
21	Dual Ti-Ru catalysis in the direct radical haloalkylation of N-acyl oxazolidinones. <i>Angewandte Chemie - International Edition</i> , 2011 , 50, 7136-9	16.4	61
20	Highly enantioselective direct alkylation of arylacetic acids with chiral lithium amides as traceless auxiliaries. <i>Journal of the American Chemical Society</i> , 2011 , 133, 11936-9	16.4	74
19	Valence tautomerism in titanium enolates: catalytic radical haloalkylation and application in the total synthesis of neodysidenin. <i>Journal of the American Chemical Society</i> , 2010 , 132, 1482-3	16.4	94
18	Marine Toxins with Spiroimine Rings: Total Synthesis of Pinnatoxin A. <i>European Journal of Organic Chemistry</i> , 2010 , 2010, 5743-5765	3.2	28
17	Concise Total Synthesis of Sintokamides A, B, and E by a Unified, Protecting-Group-Free Strategy. <i>Angewandte Chemie</i> , 2010 , 122, 9896-9899	3.6	22
16	Concise total synthesis of sintokamides A, B, and E by a unified, protecting-group-free strategy. <i>Angewandte Chemie - International Edition</i> , 2010 , 49, 9702-5	16.4	82
15	Consecutive sigmatropic rearrangements in the enantioselective total synthesis of (Pjoubertinamine and (Pmesembrine. <i>Tetrahedron</i> , 2009 , 65, 3261-3269	2.4	34
14	Studies toward the synthesis of spirolides: assembly of the elaborated E-ring fragment. <i>Organic Letters</i> , 2009 , 11, 839-42	6.2	28
13	[3,3]-Sigmatropic rearrangements: recent applications in the total synthesis of natural products. <i>Chemical Society Reviews</i> , 2009 , 38, 3133-48	58.5	183
12	Total synthesis of (+)-pinnatoxin A. Journal of the American Chemical Society, 2008, 130, 3774-6	16.4	115
11	Total synthesis of (+/-)trichodermamide B and of a putative biosynthetic precursor to aspergillazine a using an oxaza-cope rearrangement. <i>Angewandte Chemie - International Edition</i> , 2008 , 47, 6829-31	16.4	34
10	Total Synthesis of (⊞)-Trichodermamide B and of a Putative Biosynthetic Precursor to Aspergillazine A Using an Oxaza-Cope Rearrangement. <i>Angewandte Chemie</i> , 2008 , 120, 6935-6937	3.6	10
9	Synthesis of (2R,3R)-2,3-Dimethyl-1,4-Butanediol by Oxidative Homocoupling of (4S)-Isopropyl-3-Propionyl-2-Oxazolidinone 2008 , 158-171		1
8	Studies toward the synthesis of pinnatoxins: the B,C,D-dispiroketal fragment. <i>Organic Letters</i> , 2007 , 9, 3161-3	6.2	32
7	Acyclic stereocontrol in the Ireland-Claisen rearrangement of alpha-branched esters. <i>Angewandte Chemie - International Edition</i> , 2007 , 46, 7466-9	16.4	80
6	Studies toward the synthesis of pinnatoxins: the spiroimine fragment. <i>Tetrahedron Letters</i> , 2007 , 48, 6845-6848	2	18
5	Development of the 1,2-oxaza-Cope rearrangement. <i>Journal of the American Chemical Society</i> , 2006 , 128, 5356-7	16.4	24

4	ClaisenMislowEvans rearrangement. <i>Tetrahedron Letters</i> , 2006 , 47, 7519-7523	4	2/
3	An approach to the imine ring system of pinnatoxins. <i>Organic Letters</i> , 2005 , 7, 1629-31	6.2	45
2	Chemistry of pinnatoxins49-68		
1	A feed-forward Ca2+-dependent mechanism boosting glycolysis and OXPHOS by activating Aralar-malate-aspartate shuttle, upon neuronal stimulation.		1

Synthesis of the A,G-spiroimine of pinnatoxins by a microwave-assisted tandem