

# Nuno Maulide

## List of Publications by Year in descending order

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Version: 2024-02-01

173  
papers

8,116  
citations

44042

48  
h-index

60583

81  
g-index

193  
all docs

193  
docs citations

193  
times ranked

4695  
citing authors

#	ARTICLE	IF	CITATIONS
1	HFIP Mediates a Direct C-C Coupling between Michael Acceptors and Eschenmoser's salt. <i>Angewandte Chemie - International Edition</i> , 2022, 61, .	7.2	10
2	Leveraging electron-deficient iminium intermediates in a general synthesis of valuable amines. <i>Angewandte Chemie - International Edition</i> , 2022, , .	7.2	5
3	Electrochemical Umpolung C-H Functionalization of Oxindoles. <i>Journal of Organic Chemistry</i> , 2022, 87, 606-612.	1.7	13
4	Electrochemical Rearrangement of 3-Hydroxyoxindoles into Benzoxazinones. <i>Organic Letters</i> , 2022, 24, 27-32.	2.4	6
5	SpottingScience – a digital learning environment to introduce Green Chemistry to secondary students and the public. <i>Chemistry Teacher International</i> , 2022, .	0.9	1
6	Direct Stereodivergent Olefination of Carbonyl Compounds with Sulfur Ylides. <i>Journal of the American Chemical Society</i> , 2022, 144, 12536-12543.	6.6	19
7	Deployment of Sulfinimines in Charge-Accelerated Sulfonium Rearrangement Enables a Surrogate Asymmetric Mannich Reaction. <i>Journal of the American Chemical Society</i> , 2022, 144, 13044-13049.	6.6	15
8	Formal Enone $\beta$ -Arylation via I(III)-Mediated Aryl Migration/Elimination. <i>Organic Letters</i> , 2021, 23, 2094-2098.	2.4	15
9	Straightforward Access to Thiocyanates via Dealkylative Cyanation of Sulfoxides. <i>Organic Letters</i> , 2021, 23, 2510-2513.	2.4	10
10	Redox-Neutral Selenocatalysed Isomerisation of $\alpha$ -Hydroxamic Acids to $\alpha$ -Aminophenols. <i>Angewandte Chemie</i> , 2021, 133, 13896-13901.	1.6	3
11	Redox-Neutral Selenium-Catalysed Isomerisation of $\alpha$ -Hydroxamic Acids into $\alpha$ -Aminophenols. <i>Angewandte Chemie - International Edition</i> , 2021, 60, 13778-13782.	7.2	15
12	Frontispiz: Redox-Neutral Selenocatalysierte Isomerisierung von $\alpha$ -Hydroxamsäuren zu $\alpha$ -Aminophenolen. <i>Angewandte Chemie</i> , 2021, 133, .	1.6	0
13	Frontispiece: Redox-Neutral Selenium-Catalysed Isomerisation of $\alpha$ -Hydroxamic Acids into $\alpha$ -Aminophenols. <i>Angewandte Chemie - International Edition</i> , 2021, 60, .	7.2	0
14	Harnessingynamide activation to access deuterated carbonyls. <i>Tetrahedron</i> , 2021, 90, 132211.	1.0	2
15	Chemoselective $\alpha$ -Oxidation of $\beta,\beta$ -Unsaturated Amides with TEMPO. <i>Angewandte Chemie - International Edition</i> , 2021, 60, 19123-19127.	7.2	20
16	Direct Synthesis of Enamides via Electrophilic Activation of Amides. <i>Journal of the American Chemical Society</i> , 2021, 143, 10524-10529.	6.6	52
17	Chemoselektive $\alpha$ -Oxidation von $\beta,\beta$ -ungesättigten Amiden mit TEMPO. <i>Angewandte Chemie</i> , 2021, 133, 19271-19275.	1.6	7
18	As Similar As Possible, As Different As Necessary – On-Site Laboratory Teaching during the COVID-19 Pandemic. <i>Journal of Chemical Education</i> , 2021, 98, 3143-3152.	1.1	6

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19	Stable and easily available sulfide surrogates allow a stereoselective activation of alcohols. <i>Chemical Science</i> , 2021, 12, 7770-7774.	3.7	13
20	Recent discoveries on the structure of iodine( <sup>III</sup> ) reagents and their use in cross-nucleophile coupling. <i>Chemical Science</i> , 2021, 12, 853-864.	3.7	42
21	Aminoylation of Thioalkynes through Radical-Polar Crossover. <i>Synlett</i> , 2020, 31, 592-594.	1.0	1
22	Toward a Structural View of hERG Activation by the Small-Molecule Activator ICA-105574. <i>Journal of Chemical Information and Modeling</i> , 2020, 60, 360-371.	2.5	12
23	Chemoselective Alpha-Deuteration of Amides via Retroene Reaction. <i>Chemistry - A European Journal</i> , 2020, 26, 15509-15512.	1.7	24
24	Enantioselective Functionalisation of Ketones durch metallfreie elektrophile Aktivierung. <i>Angewandte Chemie</i> , 2020, 132, 21121-21125.	1.6	5
25	The unusual migratory aptitude in a case of alpha-carbonyl cation-driven 1,2-migration. <i>Tetrahedron</i> , 2020, 76, 131460.	1.0	4
26	Unconventional Macrocyclizations in Natural Product Synthesis. <i>ACS Central Science</i> , 2020, 6, 1869-1889.	5.3	27
27	Enantioselective Functionalisation of Ketones Through Metal-Free Electrophilic Activation. <i>Angewandte Chemie - International Edition</i> , 2020, 59, 20935-20939.	7.2	29
28	An Enantioselective Cyclopropanation of Carbonyl Derivatives by Oxidative Umpolung. <i>Angewandte Chemie</i> , 2020, 132, 18365-18369.	1.6	6
29	An Enantioselective Cyclopropanation of Carbonyl Derivatives by Oxidative Umpolung. <i>Angewandte Chemie - International Edition</i> , 2020, 59, 18208-18212.	7.2	17
30	A Novel Class of 7-Membered Heterocyclic Compounds. <i>European Journal of Organic Chemistry</i> , 2020, 2020, 3971-3974.	1.2	5
31	Synthesis of Novel Heterocycles by Amide Activation and Umpolung Cyclization. <i>Organic Letters</i> , 2020, 22, 2376-2380.	2.4	18
32	Gold-Catalyzed Cycloisomerization of Sulfur Ylides to Dihydrobenzothiepinines. <i>Chemistry - A European Journal</i> , 2020, 26, 10972-10975.	1.7	11
33	Reductive Iodonium: Teaching an Old Claisen New Tricks. <i>Trends in Chemistry</i> , 2020, 2, 589-592.	4.4	3
34	A Mild Synthesis of Bicyclic Alkoxyoxazolium Salts from Proline and Pipcolic Acid Derivatives. <i>European Journal of Organic Chemistry</i> , 2019, 2019, 5230-5233.	1.2	4
35	A Domino 10-Step Total Synthesis of FR252921 and Its Analogues, Complex Macrocyclic Immunosuppressants. <i>Journal of the American Chemical Society</i> , 2019, 141, 13772-13777.	6.6	18
36	A Chemoselective Enantioselective Oxytriflation Enables the Direct Asymmetric Arylation of Amides. <i>CheM</i> , 2019, 5, 1883-1891.	5.8	35

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37	Eine allgemeine Methode zur Hydroaminomethylierung von Alkenen und Alkinen. <i>Angewandte Chemie</i> , 2019, 131, 14781-14785.	1.6	12
38	Unified Approach to the Chemoselective $\alpha$ -Functionalization of Amides with Heteroatom Nucleophiles. <i>Journal of the American Chemical Society</i> , 2019, 141, 18437-18443.	6.6	65
39	A General Acid-Mediated Hydroaminomethylation of Unactivated Alkenes and Alkynes. <i>Angewandte Chemie - International Edition</i> , 2019, 58, 14639-14643.	7.2	35
40	Vinyl Cation Stabilization by Silicon Enables a Formal Metal-Free $\alpha$ -Arylation of Alkyl Ketones. <i>Angewandte Chemie - International Edition</i> , 2019, 58, 17303-17306.	7.2	27
41	Chemoselective formal $\alpha$ -functionalization of substituted aliphatic amides enabled by a facile stereoselective oxidation event. <i>Chemical Science</i> , 2019, 10, 9836-9840.	3.7	25
42	Towards a Scalable Synthesis of $\alpha$ -Oxabicyclo[2.2.0]hexanone Using Flow Photochemistry. <i>ChemPhotoChem</i> , 2019, 3, 229-232.	1.5	15
43	A redox-neutral synthesis of ketones by coupling of alkenes and amides. <i>Nature Communications</i> , 2019, 10, 2327.	5.8	30
44	Bond-Forming and -Breaking Reactions at Sulfur(IV): Sulfoxides, Sulfonium Salts, Sulfur Ylides, and Sulfinate Salts. <i>Chemical Reviews</i> , 2019, 119, 8701-8780.	23.0	533
45	$\alpha$ -Arylierung von Carbonylverbindungen mittels oxidativer C-C-Bindungsaktivierung. <i>Angewandte Chemie</i> , 2019, 131, 9921-9924.	1.6	12
46	$\alpha$ -Arylation of Carbonyl Compounds through Oxidative C-C Bond Activation. <i>Angewandte Chemie - International Edition</i> , 2019, 58, 9816-9819.	7.2	42
47	Eine flexible Kupplung von Aldehyden und Alkenen ermöglicht die diastereo- und enantioselective Herstellung von Stereotriaden. <i>Angewandte Chemie</i> , 2019, 131, 5947-5950.	1.6	1
48	$\alpha$ -Fluorination of carbonyls with nucleophilic fluorine. <i>Nature Chemistry</i> , 2019, 11, 329-334.	6.6	84
49	Diastereo- and Enantioselective Access to Stereotriads through a Flexible Coupling of Substituted Aldehydes and Alkenes. <i>Angewandte Chemie - International Edition</i> , 2019, 58, 5887-5890.	7.2	12
50	Die Stabilisierung von Vinylkationen durch Silicium erlaubt eine formale, metallfreie $\alpha$ -Arylierung von Alkylketonen. <i>Angewandte Chemie</i> , 2019, 131, 17463-17467.	1.6	9
51	Enantioselective Redox-Neutral Coupling of Aldehydes and Alkenes by an Iron-Catalyzed $\alpha$ -Catch-Release-Tethering Approach. <i>Journal of the American Chemical Society</i> , 2019, 141, 143-147.	6.6	22
52	On the formation of seven-membered rings by arene-ynamide cyclization. <i>Monatshefte für Chemie</i> , 2019, 150, 3-10.	0.9	11
53	On the Stability of Disubstituted Cyclobutenes – A Computational Study. <i>European Journal of Organic Chemistry</i> , 2019, 2019, 338-341.	1.2	7
54	A Short, Efficient, and Stereoselective Synthesis of Piperine and its Analogues. <i>Synlett</i> , 2019, 30, 413-416.	1.0	7

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55	Chemoselective $\alpha,\beta$ -Dehydrogenation of Saturated Amides. <i>Angewandte Chemie - International Edition</i> , 2019, 58, 447-451.	7.2	52
56	Electrophilic Activation of Amides for the Preparation of Poly-Substituted Pyrimidines. <i>Synthesis</i> , 2019, 51, 194-202.	1.2	10
57	A Stereoselective Reductive Hosomi-Sakurai Reaction. <i>Organic Letters</i> , 2018, 20, 1461-1464.	2.4	25
58	Sulfur-Based Ylides in Transition-Metal-Catalysed Processes. <i>Topics in Current Chemistry</i> , 2018, 376, 15.	3.0	113
59	Unusual mechanisms in Claisen rearrangements: an ionic fragmentation leading to a <i>meta</i> -selective rearrangement. <i>Chemical Science</i> , 2018, 9, 4124-4131.	3.7	28
60	(3+2) Cycloadditions of Thiouronium Ylides: A Room-Temperature, One-Pot Approach to Dihydrothiophenes. <i>Journal of Organic Chemistry</i> , 2018, 83, 2479-2485.	1.7	8
61	Expeditious synthesis of polyacetylenic water hemlock toxins and their effects on the major GABA <sub>A</sub> receptor isoform. <i>Chemical Communications</i> , 2018, 54, 2008-2011.	2.2	9
62	Hydrative Aminooxylation of Ynamides: One Reaction, Two Mechanisms. <i>Chemistry - A European Journal</i> , 2018, 24, 2515-2519.	1.7	24
63	Machine Learning for Organic Synthesis: Are Robots Replacing Chemists?. <i>Angewandte Chemie - International Edition</i> , 2018, 57, 6978-6980.	7.2	56
64	Regioselective synthesis of pyridines by redox alkylation of pyridine N-oxides with malonates. <i>Monatshefte für Chemie</i> , 2018, 149, 715-719.	0.9	11
65	A Catalytic Cross-Olefination of Diazo Compounds with Sulfoxonium Ylides. <i>Angewandte Chemie - International Edition</i> , 2018, 57, 16215-16218.	7.2	81
66	Cooperative Metal-Ligand Hydroamination Catalysis Supported by C-H Activation in Cyclam Zr(IV) Complexes. <i>Inorganic Chemistry</i> , 2018, 57, 13034-13045.	1.9	12
67	Eine katalytische Kreuz-Olefinierung von Diazoverbindungen mit Sulfoxonium-Yliden. <i>Angewandte Chemie</i> , 2018, 130, 16448-16452.	1.6	15
68	Chemoselektive $\alpha,\beta$ -Dehydrierung von gesättigten Amiden. <i>Angewandte Chemie</i> , 2018, 131, 456.	1.6	13
69	Development of a reductive Hosomi-Sakurai reaction. <i>Tetrahedron</i> , 2018, 74, 6883-6889.	1.0	9
70	Redox-Neutral Synthesis of Selenoesters by Oxyarylation of Selenoalkynes under Mild Conditions. <i>Organic Letters</i> , 2018, 20, 5881-5885.	2.4	25
71	C-H-Aktivierung ermöglicht eine kurze Totalsynthese von Chinin und Analoga mit erhöhter Antimalaria-Aktivität. <i>Angewandte Chemie</i> , 2018, 130, 10897-10901.	1.6	6
72	C-H Activation Enables a Concise Total Synthesis of Quinine and Analogues with Enhanced Antimalarial Activity. <i>Angewandte Chemie - International Edition</i> , 2018, 57, 10737-10741.	7.2	49

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73	Chemoselective Activation of Diethyl Phosphonates: Modular Synthesis of Biologically Relevant Phosphonylated Scaffolds. <i>Angewandte Chemie - International Edition</i> , 2018, 57, 13330-13334.	7.2	32
74	Catalyst-dependent selectivity in sulfonium ylide cycloisomerization reactions. <i>Chemical Science</i> , 2018, 9, 7091-7095.	3.7	19
75	Chemoselektive Aktivierung von Diethylphosphonaten: modulare Synthese von biologisch relevanten phosphonylierten GrundgerÄ¼sten. <i>Angewandte Chemie</i> , 2018, 130, 13514-13518.	1.6	2
76	Amide activation: an emerging tool for chemoselective synthesis. <i>Chemical Society Reviews</i> , 2018, 47, 7899-7925.	18.7	282
77	Total Synthesis, Stereochemical Assignment, and Divergent Enantioselective Enzymatic Recognition of Larreatricin. <i>Chemistry - A European Journal</i> , 2018, 24, 15756-15760.	1.7	16
78	Stereodivergent synthesis of 1,4-dicarbonyls by traceless chargeâ€“accelerated sulfonium rearrangement. <i>Science</i> , 2018, 361, 664-667.	6.0	176
79	Maschinelles Lernen fÄ¼r die organische Synthese: Ersetzen Roboter Chemiker?. <i>Angewandte Chemie</i> , 2018, 130, 7096-7098.	1.6	8
80	An Asymmetric Redox Arylation: Chirality Transfer from Sulfur to Carbon through a Sulfonium [3,3]â€“Sigmatropic Rearrangement. <i>Angewandte Chemie - International Edition</i> , 2017, 56, 2212-2215.	7.2	115
81	Asymmetrische Redoxarylierung: ChiralitÄ¼tstransfer von Schwefel zu Kohlenstoff durch sigmatrope Sulfoniumâ€“[3,3]â€“Umlagerung. <i>Angewandte Chemie</i> , 2017, 129, 2248-2252.	1.6	38
82	A new home for organic chemistry in Austria: the workgroup organic chemistry of the Austrian Chemical Society. <i>Monatshefte fÄ¼r Chemie</i> , 2017, 148, 1-1.	0.9	4
83	Front Cover Picture: Redoxâ€“Neutral Arylations of Vinyl Cation Intermediates ( <i>Adv. Synth. Catal.</i> 1/2017). <i>Advanced Synthesis and Catalysis</i> , 2017, 359, 1-1.	2.1	18
84	Metal-Free Redox Transformations for Câ€“C and Câ€“N Bond Construction. <i>Synlett</i> , 2017, 28, 2707-2713.	1.0	8
85	Metalâ€“Free Formal Oxidative Câ€“C Coupling by Inâ€“Situ Generation of an Enolonium Species. <i>Angewandte Chemie - International Edition</i> , 2017, 56, 5921-5925.	7.2	103
86	Direct Regioselective Synthesis of Tetrazolium Salts by Activation of Secondary Amides under Mild Conditions. <i>Organic Letters</i> , 2017, 19, 2662-2665.	2.4	42
87	Flexible and Chemoselective Oxidation of Amides to Î±-Keto Amides and Î±-Hydroxy Amides. <i>Journal of the American Chemical Society</i> , 2017, 139, 6578-6581.	6.6	115
88	Metallfreie formale oxidative Câ€“C Kupplung durch Inâ€“situâ€“Erzeugung einer elektrophilen Enoloniumspezies. <i>Angewandte Chemie</i> , 2017, 129, 6015-6019.	1.6	37
89	Reversing Polarity: Carbonyl Î±â€“Aminations with Nitrogen Nucleophiles. <i>Angewandte Chemie - International Edition</i> , 2017, 56, 12416-12423.	7.2	61
90	Umkehr der PolaritÄ¼t: Î±â€“Aminierungen von Carbonylverbindungen mit Stickstoffnucleophilen. <i>Angewandte Chemie</i> , 2017, 129, 12588-12596.	1.6	13

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91	Direct Functionalization of C-H Bonds by Iron, Nickel, and Cobalt Catalysis. <i>Chemistry - A European Journal</i> , 2017, 23, 9206-9232.	1.7	177
92	Synthesis and antimicrobial evaluation of novel analogues of dehydroabiatic acid prepared by C-H-Activation. <i>European Journal of Medicinal Chemistry</i> , 2017, 126, 937-943.	2.6	25
93	Direct synthesis of $\hat{1}^3$ -pyrones by electrophilic condensation of $\hat{1}^2$ -ketoesters. <i>Organic and Biomolecular Chemistry</i> , 2017, 15, 680-683.	1.5	16
94	Ynamide Preactivation Allows a Regio- and Stereoselective Synthesis of $\hat{1}^{\pm}, \hat{1}^2$ -Disubstituted Enamides. <i>Angewandte Chemie - International Edition</i> , 2017, 56, 15723-15727.	7.2	49
95	Regio- und stereoselektive Synthese $\hat{1}^{\pm}, \hat{1}^2$ -disubstituierter Enamide durch Voraktivierung von Inamiden. <i>Angewandte Chemie</i> , 2017, 129, 15929-15933.	1.6	11
96	C2-Modified Sparteine Derivatives Are a New Class of Potentially Long-Acting Sodium Channel Blockers. <i>ChemMedChem</i> , 2017, 12, 1819-1822.	1.6	10
97	A three-membered ring approach to carbonyl olefination. <i>Nature Communications</i> , 2017, 8, 1091.	5.8	10
98	$\hat{1}^{\pm}$ -Carbonylkationen in Sulfoxid-ermittelten oxidativen Cyclisierungen. <i>Angewandte Chemie</i> , 2017, 129, 13454-13458.	1.6	9
99	$\hat{1}^{\pm}$ -Carbonyl Cations in Sulfoxide-Driven Oxidative Cyclizations. <i>Angewandte Chemie - International Edition</i> , 2017, 56, 13270-13274.	7.2	34
100	Frontispiece: Direct Functionalization of C-H Bonds by Iron, Nickel, and Cobalt Catalysis. <i>Chemistry - A European Journal</i> , 2017, 23, .	1.7	2
101	Hydroxams-uren als chemoselektive (<i>ortho</i>-Amino)arylierungsreagenzien durch sigmatrope Umlagerung. <i>Angewandte Chemie</i> , 2017, 129, 11078-11081.	1.6	12
102	Hydroxamic Acids as Chemoselective (<i>ortho</i>-Amino)arylation Reagents via Sigmatropic Rearrangement. <i>Angewandte Chemie - International Edition</i> , 2017, 56, 10938-10941.	7.2	40
103	Intramolecular hydrogen bonding in conformationally semi-rigid $\hat{1}^{\pm}$ -acylmethane derivatives: a theoretical NMR study. <i>Organic and Biomolecular Chemistry</i> , 2017, 15, 7572-7579.	1.5	2
104	Sulfur Ylides in Organic Synthesis and Transition Metal Catalysis. <i>Structure and Bonding</i> , 2017, , 73-115.	1.0	16
105	Chemoselective Intermolecular Cross-Enolate-Type Coupling of Amides. <i>Journal of the American Chemical Society</i> , 2017, 139, 16040-16043.	6.6	85
106	Mechanistic Pathways in Amide Activation: Flexible Synthesis of Oxazoles and Imidazoles. <i>Organic Letters</i> , 2017, 19, 3815-3818.	2.4	36
107	Synthesis of $\hat{1}^3$ -pyrones via decarboxylative condensation of $\hat{1}^2$ -ketoacids. <i>Monatshefte für Chemie</i> , 2017, 148, 57-62.	0.9	11
108	Redox-Neutral Arylations of Vinyl Cation Intermediates. <i>Advanced Synthesis and Catalysis</i> , 2017, 359, 64-77.	2.1	57



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109	Synthesis and Photocatalytic Reactivity of Vinylsulfonium Ylides. <i>Journal of Organic Chemistry</i> , 2016, 81, 7201-7210.	1.7	19
110	Temporary Generation of a Cyclopropyl Oxocarbenium Ion Enables Highly Diastereoselective Donor-acceptor Cyclopropane Cycloaddition. <i>Angewandte Chemie - International Edition</i> , 2016, 55, 6780-6783.	7.2	91
111	Brønsted Acid-Mediated Hydrative Arylation of Unactivated Alkynes. <i>Chemistry - A European Journal</i> , 2016, 22, 4727-4732.	1.7	83
112	Metal-Free Synthesis of Highly Substituted Pyridines by Formal [2+2+2] Cycloaddition under Mild Conditions. <i>Angewandte Chemie</i> , 2016, 128, 13056-13059.	1.6	29
113	Making the Least Reactive Electrophile the First in Class: Domino Electrophilic Activation of Amides. <i>Journal of Organic Chemistry</i> , 2016, 81, 4421-4428.	1.7	182
114	A Gold(I)-Catalyzed Domino Coupling of Alcohols with Allenes Enables the Synthesis of Highly Substituted Indenes. <i>Chemistry - A European Journal</i> , 2016, 22, 14471-14474.	1.7	11
115	Metal-Free Synthesis of Highly Substituted Pyridines by Formal [2+2+2] Cycloaddition under Mild Conditions. <i>Angewandte Chemie - International Edition</i> , 2016, 55, 12864-12867.	7.2	87
116	Bridging C-H Activation: Mild and Versatile Cleavage of the 8-Aminoquinoline Directing Group. <i>Chemistry - A European Journal</i> , 2016, 22, 16805-16808.	1.7	53
117	Metallfreie meta-selektive Oxyarylierung von Alkinen mit Pyridin-N-Oxiden: schnelle Synthese von Metyrapon-Analoga. <i>Angewandte Chemie</i> , 2016, 128, 15650-15654.	1.6	8
118	Metal-Free meta-Selective Alkyne Oxyarylation with Pyridine N-Oxides: Rapid Assembly of Metyrapone Analogues. <i>Angewandte Chemie - International Edition</i> , 2016, 55, 15424-15428.	7.2	33
119	Metal-free intermolecular formal cycloadditions enable an orthogonal access to nitrogen heterocycles. <i>Nature Communications</i> , 2016, 7, 10914.	5.8	96
120	Cyclobutenes: At a Crossroad between Diastereoselective Syntheses of Dienes and Unique Palladium-Catalyzed Asymmetric Allylic Substitutions. <i>Accounts of Chemical Research</i> , 2016, 49, 2444-2458.	7.6	114
121	Divergent ynamide reactivity in the presence of azides – an experimental and computational study. <i>Chemical Science</i> , 2016, 7, 6032-6040.	3.7	32
122	Chemo- and Stereoselective Transition-Metal-Free Amination of Amides with Azides. <i>Journal of the American Chemical Society</i> , 2016, 138, 8348-8351.	6.6	109
123	Temporäre Bildung eines Cyclopropyl-Oxocarbeniumions ermöglicht eine außergewöhnlich diastereoselektive Cycloaddition von Donor-akzeptor-Cyclopropanen. <i>Angewandte Chemie</i> , 2016, 128, 6892-6895.	1.6	39
124	Enantioconvergent Fukuyama Cross-Coupling of Racemic Benzylic Organozinc Reagents. <i>Angewandte Chemie - International Edition</i> , 2016, 55, 4587-4590.	7.2	48
125	Redox-Neutral $\hat{\pm}$ -Amino C-H Functionalization: When the Catalyst Is Also the Nucleophile. <i>Organic Letters</i> , 2016, 18, 345-347.	2.4	23
126	Visible-Light, Metal-Free $\hat{\pm}$ -Amino C(sp <sup>3</sup> )-H Activation through 1,5-Hydrogen Migration: A Concise Method for the Preparation of Bis(indolyl)alkanes. <i>European Journal of Organic Chemistry</i> , 2015, 2015, 7643-7647.	1.2	12



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127	Dimeric TADDOL Phosphoramidites in Asymmetric Catalysis: Domino Deracemization and Cyclopropanation of Sulfonium Ylides. <i>Angewandte Chemie - International Edition</i> , 2015, 54, 10365-10369.	7.2	71
128	Strong Bonds Made Weak: Towards the General Utility of Amides as Synthetic Modules. <i>Angewandte Chemie - International Edition</i> , 2015, 54, 13856-13858.	7.2	141
129	Stereoselective Gold(I) Domino Catalysis of Allylic Isomerization and Olefin Cyclopropanation: Mechanistic Studies. <i>Journal of Organic Chemistry</i> , 2015, 80, 5719-5729.	1.7	26
130	Dynamic behaviour of monohaptoallylpalladium species: internal coordination as a driving force in allylic alkylation chemistry. <i>Chemical Science</i> , 2015, 6, 5734-5739.	3.7	8
131	A family of low molecular-weight, organic catalysts for reductive C=C bond formation. <i>Chemical Communications</i> , 2015, 51, 13902-13905.	2.2	62
132	Investigation of cationic Claisen-type electrophilic rearrangements of amides. <i>Tetrahedron</i> , 2015, 71, 5994-6005.	1.0	14
133	From Stereodefined Cyclobutenes to Dienes: Total Syntheses of leodomycin D and the Southern Fragment of Macrolactin A. <i>Organic Letters</i> , 2015, 17, 4486-4489.	2.4	43
134	A Brønsted Acid Catalyzed Redox Arylation. <i>Angewandte Chemie - International Edition</i> , 2014, 53, 8718-8721.	7.2	159
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