

Marino Petrini

List of Publications by Year in descending order

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205
docs citations

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times ranked

4140
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#	ARTICLE	IF	CITATIONS
1	Sustainable and fast synthesis of functionalized quinoxalines promoted by natural deep eutectic solvents (NADESs). <i>Green Chemistry</i> , 2022, 24, 3629-3633.	9.0	11
2	Synthesis of Nitro Alcohols by Riboflavin Promoted Tandem Nef-Henry Reactions on Nitroalkanes. <i>Advanced Synthesis and Catalysis</i> , 2021, 363, 742-746.	4.3	5
3	Enantioselective Catalyzed Synthesis of Amino Derivatives Using Electrophilic Open-Chain α -N-Activated Ketimines. <i>Advanced Synthesis and Catalysis</i> , 2021, 363, 3655-3692.	4.3	13
4	A New and Effective One-Pot Synthesis of Polysubstituted Carbazoles Starting from α -Nitro- β -Unsaturated Ketones and Indoles. <i>Asian Journal of Organic Chemistry</i> , 2021, 10, 2334-2337.	2.7	3
5	New Perspectives in the Indole Ring Functionalization using α -Indolylmethanols. <i>Advanced Synthesis and Catalysis</i> , 2020, 362, 1214-1232.	4.3	49
6	3-Alkylated indoles by reduction of sulfonyl indoles under flow chemical conditions. <i>Arkivoc</i> , 2020, 2019, 69-79.	0.5	4
7	Synthesis and practical applications of 2-(2-nitroalkyl)pyrroles. <i>Organic and Biomolecular Chemistry</i> , 2020, 18, 4533-4546.	2.8	13
8	Synthesis of Unsymmetrical Bisindolylmethanes by Reaction of Indolylmagnesium Bromides with Sulfonyl Indoles. <i>Advanced Synthesis and Catalysis</i> , 2020, 362, 1509-1513.	4.3	5
9	Recent synthetic applications of α -amido sulfones as precursors of N-acylimino derivatives. <i>Organic Chemistry Frontiers</i> , 2019, 6, 2142-2182.	4.5	36
10	Tryptophol and derivatives: natural occurrence and applications to the synthesis of bioactive compounds. <i>Natural Product Reports</i> , 2019, 36, 490-530.	10.3	41
11	Recent Advances in the Synthesis of Unsymmetrical Bisindolylmethane Derivatives. <i>Synthesis</i> , 2019, 51, 829-841.	2.3	35
12	β -Regioselective Functionalization of α -Alkenylindoles via 1,6-Addition to Extended Alkylideneindolenine Intermediates. <i>Advanced Synthesis and Catalysis</i> , 2018, 360, 1296-1302.	4.3	10
13	Oxidative Conversion of Sulfonyl Indoles into 3-Alkylidene-2-oxindoles under Flow Chemical Conditions. <i>Synthesis</i> , 2018, 50, 371-376.	2.3	6
14	Synthetic Approach to the Preparation of (2-Acetoxy)allyl Nitro Compounds. <i>Journal of Organic Chemistry</i> , 2018, 83, 12855-12862.	3.2	1
15	Novel antitumor copper(II) complexes designed to act through synergistic mechanisms of action, due to the presence of an NMDA receptor ligand and copper in the same chemical entity. <i>New Journal of Chemistry</i> , 2018, 42, 11878-11887.	2.8	16
16	Regioselective Direct α -Alkylation of Indoles. <i>Chemistry - A European Journal</i> , 2017, 23, 16115-16151.	3.3	88
17	Frontispiece: Regioselective Direct α -Alkylation of Indoles. <i>Chemistry - A European Journal</i> , 2017, 23, .	3.3	1
18	Sulfonyl Azoles in the Synthesis of 3-Functionalized Azole Derivatives. <i>Chemical Record</i> , 2016, 16, 1353-1379.	5.8	27

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19	Recent Developments in the Stereoselective Synthesis of Nitrogen-Containing Heterocycles using α -Acylimines as Reactive Substrates. <i>Advanced Synthesis and Catalysis</i> , 2016, 358, 3657-3682.	4.3	62
20	The Nitro to Carbonyl Conversion (Nef Reaction): New Perspectives for a Classical Transformation. <i>Advanced Synthesis and Catalysis</i> , 2015, 357, 2371-2402.	4.3	111
21	Reaction of α -amido sulfones with functionalized nitrocompounds: a new two-step synthesis of N-alkoxycarbonyl-2,5-disubstituted pyrroles. <i>RSC Advances</i> , 2014, 4, 43258-43261.	3.6	6
22	α -Acryloylamidoalkyl Sulfones in a Synthetic Approach for the Preparation of α -Alkyltetrahydropyridinones. <i>European Journal of Organic Chemistry</i> , 2014, 2014, 5433-5441.	2.4	5
23	Synthetic Approaches to 3-(2-Nitroalkyl) Indoles and Their Use to Access Tryptamines and Related Bioactive Compounds. <i>Chemical Reviews</i> , 2014, 114, 7108-7149.	47.7	284
24	Synthesis of 3-(2-Nitroalkyl)pyrroles from Sulfonylpyrroles and their Conversion to α -Azaindole Derivatives. <i>Advanced Synthesis and Catalysis</i> , 2013, 355, 3285-3289.	4.3	12
25	A Photochemical Route to Benzo[<i>a</i>]carbazoles via Domino Elimination/Electrocyclization of α -Aryl- β -(α -tosylalkyl)indoles. <i>Advanced Synthesis and Catalysis</i> , 2013, 355, 643-646.	4.3	30
26	Synthesis and Functionalization of Unsymmetrical Arylsulfonyl Bisindoles and Bisbenzazoles. <i>Advanced Synthesis and Catalysis</i> , 2012, 354, 3539-3544.	4.3	24
27	Ketosulfonyl indoles in the regiodefined synthesis of tryptophols and related indole derivatives. <i>Organic and Biomolecular Chemistry</i> , 2012, 10, 3486.	2.8	18
28	Solvent-Free Non-Covalent Organocatalysis: Enantioselective Addition of Nitroalkanes to Alkylideneindolenines as a Flexible Gateway to Optically Active Tryptamine Derivatives. <i>Advanced Synthesis and Catalysis</i> , 2012, 354, 1373-1380.	4.3	43
29	Arylsulfonyl Group: Activating Properties and Recent Synthetic Applications. <i>Phosphorus, Sulfur and Silicon and the Related Elements</i> , 2011, 186, 1032-1045.	1.6	19
30	Regioselective Synthesis of 3-Substituted Pyrroles by Nucleophilic Addition of 3-(α -Arylsulfonylalkyl) Pyrroles Activated under Basic or Acid Conditions. <i>Chemistry - A European Journal</i> , 2011, 17, 7183-7187.	3.3	20
31	Nitroalkanes as Key Compounds for the Synthesis of Amino Derivatives. <i>Current Organic Chemistry</i> , 2011, 15, 1482-1506.	1.6	35
32	Metal-Free Synthesis of Imido Derivatives by Direct Oxidation of α -Amido Sulfones. <i>European Journal of Organic Chemistry</i> , 2010, 2010, 5085-5089.	2.4	9
33	A Two-Step Synthesis of Unsymmetrical 1,4-Disubstituted Carbazoles from Sulfonylindoles Under Heterogeneous Catalysis. <i>Advanced Synthesis and Catalysis</i> , 2010, 352, 2459-2462.	4.3	29
34	Reaction of carbon nucleophiles with alkylideneindazolium and alkylideneindolium ions generated from their 3-(1-arylsulfonylalkyl) indazole and indole precursors. <i>Organic and Biomolecular Chemistry</i> , 2010, 8, 706-712.	2.8	24
35	Synthesis of 3-substituted indoles via reactive alkylideneindolenine intermediates. <i>Organic and Biomolecular Chemistry</i> , 2010, 8, 1259-1270.	2.8	178
36	A green procedure for the regio- and chemoselective hydrophosphonylation of unsaturated systems using CaO under solventless conditions. <i>Green Chemistry</i> , 2010, 12, 1171.	9.0	33

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37	A "Click"™ Approach to the Synthesis of 3-[2-(1-Alkyltriazol-4-yl)ethyl]indoles. <i>Synthesis</i> , 2009, 2009, 3143-3149.	2.3	15
38	Synthesis of 3-(Tosylalkyl)indazoles and their Desulfonylation Reactions – A New Entry to 3-Substituted Indazoles by an Unprecedented Friedel–Crafts Process. <i>European Journal of Organic Chemistry</i> , 2009, 2009, 3184-3188.	2.4	18
39	Synthesis of indolylalkylphosphonates and 3-(1-diphenylphosphinoalkyl) indoles by reaction of 3-(1-arylsulfonylalkyl) indoles with phosphorus derivatives. <i>Tetrahedron Letters</i> , 2008, 49, 5645-5648.	1.4	18
40	Proline-Catalyzed Asymmetric Formal α -Alkylation of Aldehydes via Vinylogous Iminium Ion Intermediates Generated from Arylsulfonyl Indoles. <i>Angewandte Chemie - International Edition</i> , 2008, 47, 8707-8710.	13.8	187
41	Reaction of 3-(1-Arylsulfonylalkyl)indoles with Easily Enolisable Derivatives Promoted by Potassium Fluoride on Basic Alumina. <i>Advanced Synthesis and Catalysis</i> , 2008, 350, 129-134.	4.3	59
42	Improved preparation of alkyl 2-(3-indolyl)-3-nitroalkanoates under fully heterogeneous conditions: stereoselective synthesis of alkyl (E)-2-(3-indolyl)-2-alkenoates. <i>Tetrahedron</i> , 2008, 64, 5435-5441.	1.9	24
43	Double Functionalization of <i>N</i> -Boc-3-(Tosylmethyl)indole Exploiting the Activating Properties of the Tosyl Group. <i>Synlett</i> , 2008, 2008, 1845-1851.	1.8	6
44	Nitroalkanes as Central Reagents in the Synthesis of Spiroketal. <i>Molecules</i> , 2008, 13, 319-330.	3.8	34
45	Nitroalkanes as key building blocks for the synthesis of heterocyclic derivatives. <i>Arkivoc</i> , 2008, 2009, 195-223.	0.5	15
46	Recent Advances in Stereoselective Syntheses Using <i>N</i> -Acylimines. <i>Synthesis</i> , 2007, 2007, 159-186.	2.3	100
47	An Efficient Diastereoselective Route to Differentially Protected anti-4-Amino-1-alken-3-ols. <i>Journal of Organic Chemistry</i> , 2007, 72, 1834-1837.	3.2	15
48	Simplified Synthesis of 3-(1-Arylsulfonylalkyl) Indoles and Their Reaction with Reformatsky Reagents. <i>Journal of Organic Chemistry</i> , 2007, 72, 1863-1866.	3.2	61
49	Synthesis of 3-(2-nitroalkyl) indoles by reaction of 3-(1-arylsulfonylalkyl) indoles with nitroalkanes. <i>Tetrahedron Letters</i> , 2007, 48, 5653-5656.	1.4	20
50	Stereoselective synthesis of vicinal aminodiols, diamines and diaminiols by the use of enantiopure aldehydes in the three-component aromatic Mannich-type reaction. <i>Tetrahedron: Asymmetry</i> , 2007, 18, 1022-1029.	1.8	19
51	Solventless Clay-Promoted Friedel–Crafts Reaction of Indoles with α -Amido Sulfones: Unexpected Synthesis of 3-(1-Arylsulfonylalkyl) Indoles. <i>Organic Letters</i> , 2006, 8, 4093-4096.	4.6	100
52	α -Amido sulfones from natural α -amino acids and their reaction with carbon nucleophiles. <i>Tetrahedron</i> , 2006, 62, 960-967.	1.9	11
53	Aza-Henry reaction of substituted nitroalkanes using α -formamidoaryl sulfones as <i>N</i> -acylimino equivalents. <i>Tetrahedron Letters</i> , 2006, 47, 3501-3503.	1.4	19
54	Conjugate Addition of Indoles to Nitroalkenes Promoted by Basic Alumina in Solventless Conditions. <i>Advanced Synthesis and Catalysis</i> , 2006, 348, 191-196.	4.3	54

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55	Nitrocompounds as useful reagents for the synthesis of dicarbonyl derivatives. <i>Arkivoc</i> , 2006, 2006, 127-152.	0.5	12
56	TiCl ₄ -promoted addition of nucleophiles to open chain $\hat{\pm}$ -amidoalkylphenyl sulfones. <i>Tetrahedron Letters</i> , 2005, 46, 5999-6003.	1.4	30
57	Conjugate Additions of Nitroalkanes to Electron-Poor Alkenes: Recent Results. <i>Chemical Reviews</i> , 2005, 105, 933-972.	47.7	465
58	Conjugate Additions of Nitroalkanes to Electron-Poor Alkenes: Recent Results. <i>ChemInform</i> , 2005, 36, no.	0.0	1
59	TiCl ₄ -Promoted Addition of Nucleophiles to Open Chain $\hat{\pm}$ -Amidoalkylphenyl Sulfones.. <i>ChemInform</i> , 2005, 36, no.	0.0	0
60	$\hat{\pm}$ -Amido Sulfones as Stable Precursors of Reactive N-Acylimino Derivatives. <i>Chemical Reviews</i> , 2005, 105, 3949-3977.	47.7	221
61	Michael Addition of Nitroalkanes to Optically Active Acrylates Mediated by Cetyltrimethylammonium Hydroxide (CTAOH). <i>Letters in Organic Chemistry</i> , 2004, 1, 335-339.	0.5	4
62	Recent synthetic developments in the nitro to carbonyl conversion (Nef reaction). <i>Tetrahedron</i> , 2004, 60, 1017-1047.	1.9	416
63	Highly Diastereoselective Addition of Nitromethane Anion to Chiral $\hat{\pm}$ -Amidoalkylphenyl Sulfones. Synthesis of Optically Active $\hat{\pm}$ -Amino Acid Derivatives.. <i>ChemInform</i> , 2004, 35, no.	0.0	0
64	Recent Synthetic Developments in the Nitro to Carbonyl Conversion (Nef Reaction). <i>ChemInform</i> , 2004, 35, no.	0.0	0
65	Investigation into the Allylation Reactions of Aldehydes Promoted by the CeCl ₃ ·7H ₂ O~NaI System as a Lewis Acid.. <i>ChemInform</i> , 2004, 35, no.	0.0	0
66	Synthesis of advanced intermediates for the preparation of aza-analogues of podophyllotoxin. <i>Tetrahedron Letters</i> , 2004, 45, 2133-2136.	1.4	29
67	Investigation into the Allylation Reactions of Aldehydes Promoted by the CeCl ₃ ·7H ₂ O~NaI System as a Lewis Acid. <i>Journal of Organic Chemistry</i> , 2004, 69, 1290-1297.	3.2	45
68	Reactivity of Chiral $\hat{\pm}$ -Amidoalkylphenyl Sulfones with Stabilized Carbanions. Stereoselective Synthesis of Optically Active 1-Aminopyrrolizidine. <i>Journal of Organic Chemistry</i> , 2004, 69, 7303-7308.	3.2	37
69	Reactivity of Chiral Exocyclic N-Acyliminium Ions with Aromatic Derivatives.. <i>ChemInform</i> , 2003, 34, no.	0.0	0
70	Conjugate Addition of Nitroalkanes to N-Substituted Maleimides. Synthesis of 3-Alkylsuccinimides and Pyrrolidines.. <i>ChemInform</i> , 2003, 34, no.	0.0	0
71	Conjugate Addition of Nitroalkanes to Dimethyl Maleate. Regioselective Formation of Both Monoesters of 2-Alkylsuccinic Acids.. <i>ChemInform</i> , 2003, 34, no.	0.0	0
72	Conjugate addition of nitroalkanes to N-substituted maleimides. Synthesis of 3-alkylsuccinimides and pyrrolidines. <i>Tetrahedron</i> , 2003, 59, 3603-3608.	1.9	50

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73	Conjugate addition of nitroalkanes to dimethyl maleate. Regioselective formation of both monoesters of 2-alkylsuccinic acids. <i>Tetrahedron</i> , 2003, 59, 7283-7289.	1.9	13
74	Reactivity of chiral exocyclic N-acyliminium ions with aromatic derivatives. <i>Tetrahedron: Asymmetry</i> , 2003, 14, 1171-1178.	1.8	24
75	Highly diastereoselective addition of nitromethane anion to chiral $\hat{1}\pm$ -amidoalkylphenyl sulfones. Synthesis of optically active $\hat{1}\pm$ -amino acid derivatives. <i>Organic and Biomolecular Chemistry</i> , 2003, 1, 4275-4281.	2.8	82
76	Reaction of Allylzinc Reagents and Zinc Enolates of Ketones with $\hat{1}\pm$ -Amidoalkylphenyl Sulfones. <i>Journal of Organic Chemistry</i> , 2002, 67, 4530-4535.	3.2	44
77	Allylation of Exocyclic N-Acyliminium Ions Generated from Chiral N-[1-(Phenylsulfonyl)alkyl]oxazolidin-2-ones. <i>Journal of Organic Chemistry</i> , 2002, 67, 2989-2994.	3.2	25
78	Unprecedented, selective Nef reaction of secondary nitroalkanes promoted by DBU under basic homogeneous conditions. <i>Tetrahedron Letters</i> , 2002, 43, 5233-5235.	1.4	55
79	Reaction of $\hat{1}\pm$ -Amidoalkylphenyl Sulfones with Lithiated Nitriles: $\hat{1}\pm$ -Syn-Selective Synthesis of $\hat{1}\pm$ -Amino Nitriles. <i>Journal of Organic Chemistry</i> , 2001, 66, 8264-8267.	3.2	20
80	Stereoselective Synthesis of (E)-4-Alkylidenecyclopent-2-en-1-ones by a Tandem Ring Closure $\hat{1}\pm$ -Michael Addition $\hat{1}\pm$ -Elimination. <i>Organic Letters</i> , 2001, 3, 1265-1267.	4.6	40
81	Synthesis of functionalized nitrocyclohexene derivatives from 2-nitrocycloalkanones, via anionic domino reactions. <i>Tetrahedron</i> , 2001, 57, 6079-6081.	1.9	7
82	Claisen-Johnson Orthoester Rearrangement of $\hat{1}\pm$ -Hydroxy $\hat{1}\pm$, $\hat{1}\pm$ -Unsaturated Ketones and Nitriles. <i>European Journal of Organic Chemistry</i> , 2001, 2001, 713-718.	2.4	15
83	Conjugate Addition of Amines to $\hat{1}\pm$, $\hat{1}\pm$ -Enones Promoted by $\text{CeCl}_3 \cdot 7\text{H}_2\text{O} / \text{NaI}$ System Supported in Silica Gel. <i>Journal of Organic Chemistry</i> , 2001, 66, 9052-9055.	3.2	166
84	Synthesis of (E)-3-Alkylidenepyrrolidines by Nucleophilic Ring Closure of (E)-2-Alkylidene-1,4-diol Derivatives. <i>European Journal of Organic Chemistry</i> , 2000, 2000, 2927-2931.	2.4	14
85	Reaction of $\hat{1}\pm$ -amidoalkylphenyl sulfones with Reformatsky reagents. A new entry to $\hat{1}\pm$ -amino esters. <i>Tetrahedron Letters</i> , 2000, 41, 2709-2712.	1.4	29
86	2,5-Dialkylfurans and Nitroalkanes as Source of 2,3,5-Trialkylpyrroles. <i>Synlett</i> , 2000, 2000, 391-393.	1.8	6
87	A Novel Route to the Vinyl Sulfide Nine-Membered Macrocyclic Moiety of Griseoviridin. <i>Journal of Organic Chemistry</i> , 2000, 65, 4553-4559.	3.2	98
88	Acyclic Stereoselection in the Reaction of Nucleophilic Reagents with Chiral N-Acyliminium Ions Generated from N-[1-(Phenylsulfonyl)alkyl]imidazolidin-2-ones. <i>Journal of Organic Chemistry</i> , 2000, 65, 8277-8282.	3.2	26
89	An Efficient Procedure for the Diastereoselective Dehydration of $\hat{1}\pm$ -Hydroxy Carbonyl Compounds by $\text{CeCl}_3 \cdot 7\text{H}_2\text{O} / \text{NaI}$ System. <i>Organic Letters</i> , 2000, 2, 1791-1793.	4.6	28
90	Base assisted substitution of $\hat{1}\pm$ -amidoalkyl sulfones by nitromethane anion. A new entry to functionalized $\hat{1}\pm$ -amino acids. <i>Tetrahedron Letters</i> , 1999, 40, 4449-4452.	1.4	37

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91	Synthesis of Allylic and Propargylic Primary Amines by Reaction of Organometallic Reagents with β -Amidoalkyl Sulfones. <i>Journal of Organic Chemistry</i> , 1999, 64, 8970-8972.	3.2	101
92	Claisen rearrangement of β -hydroxyvinyl sulfones via ketene acetal derivatives. A new entry to functionalized (2E,4E)-alkadienoic esters. <i>Tetrahedron Letters</i> , 1998, 39, 5827-5830.	1.4	20
93	Synthesis and Radical Cyclization of 2-Allylamino-3-chloropropylphenyl Sulfones to 2,4-Disubstituted Pyrrolidines. <i>Synlett</i> , 1998, 1998, 90-92.	1.8	9
94	Conjugate addition of allylic and prop-2-ynylic alcohols to 3-halogeno-1-phenylsulfonylprop-1-enes; synthesis and radical induced cyclization of 2-alkenyloxy-3-halogenopropylphenyl sulfones. <i>Chemical Communications</i> , 1997, , 1829.	4.1	5
95	Radical induced allylations of functionalized β -haloalkylphenyl sulfones. <i>Tetrahedron Letters</i> , 1997, 38, 1995-1998.	1.4	17
96	Ruthenium tetroxide catalyzed oxidations of 3-alkyl-4-(2-furyl)-4-oxobutanenitriles: Synthesis of methyl 2-alkyl-3-cyanopropanoates. <i>Tetrahedron Letters</i> , 1997, 38, 3781-3784.	1.4	16
97	Oxidative Ring Cleavage of 2-Nitrocycloalkanones: Synthesis and Radical-Induced Transformations of Methyl β -Dihalo- β -nitroalkanoates. <i>Journal of Organic Chemistry</i> , 1996, 61, 5652-5655.	3.2	18
98	A Stereoselective Synthesis of α,β -Unsaturated Ketones Involving the Reactions of Organocerium Reagents with Secondary β -Enamino Ketones. <i>Chemistry - A European Journal</i> , 1996, 2, 913-918.	3.3	33
99	LiClO ₄ Mediated Substitution of β -Phenylsulfonyl- β -oxo Arenebutanenitriles by Organomagnesium Reagents. <i>Synlett</i> , 1996, 1996, 1001-1003.	1.8	9
100	A new synthesis of α -phoracantholide, α -dihydroreifeolide, and α -muscone via β -nitro ketones. <i>Liebigs Annalen</i> , 1995, 1995, 1381-1383.	0.8	13
101	Oxidation of secondary amines to nitrones using urea-hydrogen peroxide complex (UHP) and metal catalysts. <i>Tetrahedron Letters</i> , 1995, 36, 3561-3562.	1.4	81
102	A New Procedure for the Desulfonylation of β -Keto Phenylsulfones Using Bu ₃ SnCl/NaBH ₃ CN Couple. <i>Synlett</i> , 1995, 1995, 973-974.	1.8	13
103	Stereoselective Total Synthesis of (+)-Lentiginosine Using a Chiral Nitrone Intermediate. <i>Journal of Organic Chemistry</i> , 1995, 60, 5706-5707.	3.2	70
104	Cerium chloride (III) promoted nucleophilic addition of organolithium reagents to β -diphenylphosphinoyl ketones. An efficient method for the synthesis of horner-wittig intermediates. <i>Tetrahedron Letters</i> , 1994, 35, 8453-8456.	1.4	28
105	Chemo- and Diastereoselective Reduction of β -Enamino Esters: A Convenient Synthesis of Both cis- and trans- γ -Amino Alcohols and β -Amino Esters. <i>Journal of Organic Chemistry</i> , 1994, 59, 5328-5335.	3.2	197
106	Highly stereoselective synthesis of α,β -unsaturated ketones by CeCl ₃ mediated addition of grignard reagents to β -enamino ketones. <i>Journal of the Chemical Society Chemical Communications</i> , 1994, , 715-716.	2.0	19
107	CeCl ₃ -Mediated Addition of Grignard Reagents to 1,3-Diketones. <i>Angewandte Chemie International Edition in English</i> , 1993, 32, 1061-1062.	4.4	38
108	Retro Claisen cleavage of β -nitrocycloalkanones using trimethylsilylmethylmagnesium chloride (Peterson reagent): Synthesis of functionalized β -keto-trimethylsilanes. <i>Tetrahedron Letters</i> , 1993, 34, 3301-3304.	1.4	19

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109	Cerium(III) chloride mediated addition of Grignard reagents to nitroalkanes: synthesis of N,N-disubstituted hydroxylamines. <i>Journal of the Chemical Society Chemical Communications</i> , 1993, , 1373.	2.0	26
110	Reactivity of .alpha.-nitro ketones toward organometallic reagents: straightforward synthesis of tertiary .beta.-nitroalkanols. <i>Journal of Organic Chemistry</i> , 1993, 58, 3368-3372.	3.2	14
111	Amberlyst A 21 as New and Efficient Surface Catalyst for the Cleavage of 2-Nitrocycloalkanones. <i>Synthesis</i> , 1992, 1992, 355-357.	2.3	25
112	1%-Nitroalcohols as Precursors of Aldehydes 1%-Functionalized: A New Synthesis of 7-Acetoxyheptanal, 7-(2-Tetrahydropyranloxy)Heptanal and 8-(2-Tetrahydropyranloxy)Octanal. <i>Synthetic Communications</i> , 1992, 22, 641-647.	2.1	1
113	Hydroxy-functionalized conjugated nitroolefins as immediate precursors of spiroketals. A new synthesis of 1,7-dioxaspiro[5.5]undecane and (E)-2-methyl-1,7-dioxaspiro[5.6]dodecane. <i>Journal of the Chemical Society Perkin Transactions 1</i> , 1992, , 3159.	0.9	17
114	Nitrones from addition of benzyl and allyl Grignard reagents to alkyl nitro compounds: chemo-, regio-, and stereoselectivity of the reaction. <i>Journal of Organic Chemistry</i> , 1992, 57, 5834-5840.	3.2	25
115	A nitron-based approach to the enantioselective total synthesis of (-)-anisomycin. <i>Journal of Organic Chemistry</i> , 1992, 57, 1316-1318.	3.2	103
116	Synthesis of functionalized nitroalkanes by oxidation of oximes with urea-hydrogen peroxide complex and trifluoroacetic anhydride. <i>Tetrahedron Letters</i> , 1992, 33, 4835-4838.	1.4	57
117	Chemoselective synthesis of functionalized conjugated nitroalkenes. <i>Journal of Organic Chemistry</i> , 1992, 57, 2160-2162.	3.2	94
118	Enantioselective synthesis of nitrogen derivatives by allyl Grignard addition on optically active nitroalkanes. <i>Journal of the Chemical Society Chemical Communications</i> , 1991, , 793.	2.0	9
119	An Improved and Simple Synthesis of Methyl or Ethyl 7-Oxoheptanoate and 7-Acetoxyheptanal. <i>Synthetic Communications</i> , 1991, 21, 1075-1081.	2.1	8
120	Mechanistic studies on the reaction of nitro- and nitrosoarenes with vinyl Grignard reagents. <i>Journal of the Chemical Society Perkin Transactions II</i> , 1991, , 657.	0.9	61
121	Enantioselective synthesis of the lactone moiety of the mevinic acids using D-xylose as a chiral precursor. <i>Journal of the Chemical Society Perkin Transactions 1</i> , 1991, , 490.	0.9	11
122	New and efficient synthesis of 1%-nitroalcohols and spiroketals by chemo- and regioselective reductive cleavage of 2-nitrocycloalkanones. <i>Tetrahedron</i> , 1990, 46, 7531-7538.	1.9	44
123	A new rearrangement of nitrones: acid promoted conversion of vlnitrones into N-(1 ³ -ketoalkyl)-N-phenylhydroxylamines. <i>Tetrahedron Letters</i> , 1990, 31, 6089-6092.	1.4	3
124	A new approach to the synthesis of 2-substituted indoles: reaction of dimetallated ortho-trimethylsilylmethylanilides with esters. <i>Tetrahedron</i> , 1990, 46, 1379-1384.	1.9	24
125	A New Procedure for Dethioacetalization via Equilibrium Exchange with Aqueous Acetone, Paraformaldehyde and Amberlyst 15 as Acidic Catalyst. <i>Synthesis</i> , 1990, 1990, 336-337.	2.3	18
126	Synthetic studies on the mevinic acids using the chiron approach: total synthesis of (+)-dihydromevinolin. <i>Journal of Organic Chemistry</i> , 1990, 55, 5766-5777.	3.2	55

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