

Dina Scarpi

List of Publications by Year in descending order

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papers

1,193
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304743

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#	ARTICLE	IF	CITATIONS
1	Gold(I)-Catalyzed Cycloisomerization/Hetero-Diels-Alder Reaction/Ring Opening Cascade to Functionalized Cyclopentadienes. <i>Journal of Organic Chemistry</i> , 2022, 87, 6038-6051.	3.2	3
2	Synthesis of (±)-epi-ungianol by the Gold(I)-Catalyzed Propargyl Claisen Rearrangement/Hydroarylation Cascade Reaction of Propargyl Vinyl Ethers. <i>European Journal of Organic Chemistry</i> , 2021, 2021, 1266-1273.	2.4	6
3	Gold(I)-Catalysed Hydroarylation of Lactam-Derived Enynes as an Entry to Tetrahydrobenzo[<i>g</i>]quinolines. <i>European Journal of Organic Chemistry</i> , 2020, 2020, 646-653.	2.4	6
4	Enantioselective Synthesis of <i>cis</i> and <i>trans</i> 4-Aminopiperic Acids as ³ Amino Acids for the Construction of Cyclic RGD-Containing Peptidomimetics Antagonists of $V\alpha 3$ Integrin. <i>European Journal of Organic Chemistry</i> , 2020, 2020, 4371-4383.	2.4	1
5	One-Pot Access to 1,7a-Dihydro-1,3a-ethano-indene and 1,8a-Dihydro-1,3a-ethano-azulene Skeletons by a Sequential Gold(I)-Catalyzed Propargyl Claisen Rearrangement/Nazarov Cyclization/[4+2] Cycloaddition Reaction. <i>Journal of Organic Chemistry</i> , 2020, 85, 5078-5086.	3.2	5
6	Pentannulation of N-heterocycles by a tandem gold-catalyzed [3,3]-rearrangement/Nazarov reaction of propargyl ester derivatives: a computational study on the crucial role of the nitrogen atom. <i>Beilstein Journal of Organic Chemistry</i> , 2020, 16, 3059-3068.	2.2	2
7	Recent Advances in the Synthesis of Indenes. <i>European Journal of Organic Chemistry</i> , 2019, 2019, 7401-7419.	2.4	39
8	Synthesis of Indenes by Tandem Gold(I)-Catalyzed Claisen Rearrangement/Hydroarylation Reaction of Propargyl Vinyl Ethers. <i>Journal of Organic Chemistry</i> , 2019, 84, 6298-6311.	3.2	14
9	Stereodivergent synthesis of 5-aminopiperic acids and application in the preparation of a cyclic RGD peptidomimetic as a nanomolar $V\alpha 3$ integrin ligand. <i>Organic and Biomolecular Chemistry</i> , 2018, 16, 3402-3414.	2.8	4
10	Short synthesis of racemic 5-hydroxy-6-hydroxymethylpiperidin-2-one. <i>Chemical Data Collections</i> , 2018, 13-14, 11-16.	2.3	0
11	Pentannulation Reaction by Tandem Gold(I)-Catalyzed Propargyl Claisen Rearrangement/Nazarov Cyclization of Enynyl Vinyl Ethers. <i>Organic Letters</i> , 2018, 20, 4713-4717.	4.6	19
12	A Gold(I)-Catalyzed Oxidative Rearrangement of Heterocycle-Derived 1,3-Enynes Provides an Efficient and Selective Route to Divinyl Ketones. <i>European Journal of Organic Chemistry</i> , 2017, 2017, 6228-6238.	2.4	12
13	Total Synthesis of Bruceolline I. <i>Journal of Natural Products</i> , 2017, 80, 2384-2388.	3.0	23
14	Synthesis and conformational analysis of peptides embodying 2,3-methanopiperic acids. <i>Organic and Biomolecular Chemistry</i> , 2017, 15, 6826-6836.	2.8	14
15	Construction of Cyclopenta[<i>b</i>]indol-1-ones by a Tandem Gold(I)-Catalyzed Rearrangement/Nazarov Reaction and Application to the Synthesis of Bruceolline H. <i>Organic Letters</i> , 2016, 18, 3922-3925.	4.6	33
16	Cyclic RGD peptidomimetics containing 4- and 5-amino-cyclopropane piperic acid (CPA) templates as dual $V\alpha 3$ and $V\alpha 1$ integrin ligands. <i>Bioorganic and Medicinal Chemistry</i> , 2016, 24, 703-711.	3.0	14
17	Annulated N-Heterocycles by Tandem Gold(I)-Catalyzed [3,3]-Rearrangement/Nazarov Reaction of Propargylic Ester Derivatives: an Experimental and Computational Study. <i>European Journal of Organic Chemistry</i> , 2015, 2015, 3943-3956.	2.4	32
18	Gold-Catalysed Synthesis of Exocyclic Vinylogous Amides and ² Amino Ketones: A Detailed Study on the <i>exo</i> / <i>endo</i> - <i>dig</i> Selectivity, Methodology and Scope. <i>European Journal of Organic Chemistry</i> , 2015, 2015, 3251-3265.	2.4	23

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19	Stereochemical Assignment of Strigolactone Analogues Confirms Their Selective Biological Activity. <i>Journal of Natural Products</i> , 2015, 78, 2624-2633.	3.0	24
20	A Short, Chemoenzymatic Synthesis of Both Enantiomers of <i>trans</i> -3-Hydroxy- β -pipecolic Acid. <i>European Journal of Organic Chemistry</i> , 2014, 2014, 5448-5455.	2.4	14
21	Cyclopropane Pipecolic Acids as Templates for Linear and Cyclic Peptidomimetics: Application in the Synthesis of an Arg-Gly-Asp (RGD)-Containing Peptide as an α - β -Integrin Ligand. <i>Chemistry - A European Journal</i> , 2014, 20, 11187-11203.	3.3	17
22	Tailoring fluorescent strigolactones for in vivo investigations: a computational and experimental study. <i>Organic and Biomolecular Chemistry</i> , 2014, 12, 2960-2968.	2.8	28
23	Synthesis of Vinylogous Amides by Gold(I)-Catalyzed Cyclization of N-Boc-Protected 6-Alkynyl-3,4-dihydro-2H-pyridines. <i>Journal of Organic Chemistry</i> , 2013, 78, 11007-11016.	3.2	31
24	Complementary and Stereodivergent Approaches to the Synthesis of 5-Hydroxy- and 4,5-Dihydroxypipicolic Acids from Enantiopure Hydroxylated Lactams. <i>European Journal of Organic Chemistry</i> , 2013, 2013, 1306-1317.	2.4	23
25	Synthesis of Both Enantiomers of the Streptomyces Alkaloid 4-epi-SS20846A. <i>Synthesis</i> , 2012, 44, 3688-3692.	2.3	4
26	Low molecular weight, non-peptidic agonists of TrkA receptor with NGF-mimetic activity. <i>Cell Death and Disease</i> , 2012, 3, e339-e339.	6.3	48
27	Expeditious Racemic and Enantiodivergent Synthesis of 1-Deoxymannojirimycin and 1,4-Dideoxymannojirimycin. <i>European Journal of Organic Chemistry</i> , 2012, 2012, 2597-2605.	2.4	8
28	New Potent Fluorescent Analogues of Strigolactones: Synthesis and Biological Activity in Parasitic Weed Germination and Fungal Branching. <i>European Journal of Organic Chemistry</i> , 2011, 2011, 3781-3793.	2.4	69
29	Diastereodivergent Synthesis of 4-Hydroxy-2,3-methanopipicolic Acid Derivatives as Conformationally Constrained Homoserine Analogues. <i>European Journal of Organic Chemistry</i> , 2011, 2011, 6544-6552.	2.4	13
30	Enantiodivergent Chemoenzymatic Synthesis of 4-Hydroxypiperidine Alkaloids. <i>European Journal of Organic Chemistry</i> , 2010, 2010, 5831-5840.	2.4	33
31	Chemistry of Lactam-Derived Vinyl Phosphates: Stereoselective Synthesis of (+)-Fagomine. <i>Synlett</i> , 2010, 2010, 839-839.	1.8	0
32	A New, Practical and Efficient Method for Protecting Alcohols as tert-Butyl Ethers. <i>Synlett</i> , 2010, 2010, 812-816.	1.8	2
33	One-Pot Pictet-Spengler Reaction and Esterification for the Preparation of a Key Tadalafil Synthetic Intermediate. <i>Letters in Organic Chemistry</i> , 2010, 7, 311-313.	0.5	1
34	Carbonylative Palladium-Catalyzed Reactions of Lactam-, Lactone-, and Thiolactone-Derived Vinyl Triflates and Phosphates for the Synthesis of N-, O-, and S-Heterocycles. <i>Heterocycles</i> , 2010, 80, 697.	0.7	18
35	Chemistry of Lactam-Derived Vinyl Phosphates: Stereoselective Synthesis of (+)-Fagomine. <i>Synlett</i> , 2009, 2009, 913-916.	1.8	3
36	A Short and Convenient Synthesis of Enantiopure cis- and trans-4-Hydroxypipicolic Acid. <i>Synthesis</i> , 2009, 2009, 3611-3616.	2.3	17

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37	N-Substituent effects on the diethylzinc addition to benzaldehyde catalysed by bicyclic 1,4-amino alcohols. <i>Tetrahedron: Asymmetry</i> , 2009, 20, 340-350.	1.8	19
38	Structural diversity of bicyclic amino acids. <i>Amino Acids</i> , 2008, 34, 1-24.	2.7	67
39	Predicting Reactivity and Stereoselectivity in the Nazarov Reaction: A Combined Computational and Experimental Study. <i>Chemistry - A European Journal</i> , 2008, 14, 9292-9304.	3.3	27
40	Stereoselective Synthesis of (2 <i>S</i> ,4 <i>R</i>)-4-Hydroxypipicolinic Acid. <i>European Journal of Organic Chemistry</i> , 2008, 2008, 524-531.	2.4	16
41	3-Aza-8,10-dioxabicyclo[5.2.1]decane (9-exo BTKa) carboxylic acid as a new reverse turn inducer: synthesis and conformational analysis of a model peptide. <i>Tetrahedron</i> , 2006, 62, 1575-1582.	1.9	2
42	Synthesis of a new 1,4-aminoalcohol and its use as catalyst in the enantioselective addition of organozinc to aldehydes. <i>Tetrahedron: Asymmetry</i> , 2006, 17, 1409-1414.	1.8	16
43	Design, Synthesis, and Applications of 3-Aza-6,8-Dioxabicyclo[3.2.1]Octane-Based Scaffolds for Peptidomimetic Chemistry. <i>Synlett</i> , 2006, 2006, 0331-0353.	1.8	5
44	Selectivity of <i>Daucus carota</i> roots and baker's yeast in the enantioselective reduction of β -nitroketones. <i>Tetrahedron: Asymmetry</i> , 2005, 16, 1479-1483.	1.8	20
45	5 α -Reductase activity in <i>Lycopersicon esculentum</i> : Cloning and functional characterization of LeDET2 and evidence of the presence of two isoenzymes. <i>Journal of Steroid Biochemistry and Molecular Biology</i> , 2005, 96, 287-299.	2.5	17
46	Enantioselective addition of diethylzinc to aldehydes using 1,4-aminoalcohols as chiral ligands. <i>Tetrahedron: Asymmetry</i> , 2004, 15, 1319-1324.	1.8	26
47	Synthesis of New Molecular Scaffolds: 3-Aza-7,9-dioxabicyclo[4.2.1]nonane (8-exo BTKa) and 3-Aza-8,10-dioxabicyclo[5.2.1]decane (9-exo BTKa) Carboxylic Acids.. <i>ChemInform</i> , 2004, 35, no.	0.0	0
48	Enantioselective Addition of Diethylzinc to Aldehydes Using 1,4-Aminoalcohols as Chiral Ligands.. <i>ChemInform</i> , 2004, 35, no.	0.0	0
49	Inhibition of human β -tryptase by Bowman's Birk inhibitor derived peptides: creation of a new tri-functional inhibitor. <i>Bioorganic and Medicinal Chemistry</i> , 2004, 12, 6045-6052.	3.0	20
50	Synthesis of new molecular scaffolds: 3-aza-7,9-dioxabicyclo[4.2.1]nonane (8-exo BTKa) and 3-aza-8,10-dioxabicyclo[5.2.1]decane (9-exo BTKa) carboxylic acids. <i>Tetrahedron</i> , 2004, 60, 2583-2591.	1.9	8
51	Synthesis of 17 β -N-Substituted 19-Nor-10-azasteroids as Inhibitors of Human 5 α -Reductases I and II. <i>Bioorganic and Medicinal Chemistry</i> , 2002, 10, 3455-3461.	3.0	10
52	Synthesis of a new enantiopure bicyclic β/β' -amino acid (BTKa) derived from tartaric acid and β -amino acetophenone. <i>Tetrahedron</i> , 2002, 58, 9865-9870.	1.9	24
53	Introduction of the new dipeptide isostere 7-endo-BtA as reverse turn inducer in a Bowman-Birk proteinase inhibitor. <i>Bioorganic and Medicinal Chemistry</i> , 2001, 9, 1625-1632.	3.0	18
54	Effect of C-ring modifications in benzo[c]quinolizin-3-ones, new selective inhibitors of human 5 α -reductase 1. <i>Bioorganic and Medicinal Chemistry</i> , 2001, 9, 1385-1393.	3.0	22

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55	Stereoselective Meisenheimer rearrangement using BTAA's as chiral auxiliaries. <i>Tetrahedron: Asymmetry</i> , 2000, 11, 4227-4238.	1.8	23
56	Synthesis of 8-chloro-benzo[c]quinolizin-3-ones as potent and selective inhibitors of human steroid 5 β -reductase 1. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2000, 10, 353-356.	2.2	19
57	Modification of the Aza-Robinson Annulation for the Synthesis of 4-Methyl-Benzo[c]quinolizin-3-ones, Potent Inhibitors of Steroid 5 β -Reductase 1. <i>Journal of Organic Chemistry</i> , 2000, 65, 8093-8095.	3.2	27
58	Benzo[c]quinolizin-3-ones: A Novel Class of Potent and Selective Nonsteroidal Inhibitors of Human Steroid 5 β -Reductase 1. <i>Journal of Medicinal Chemistry</i> , 2000, 43, 3718-3735.	6.4	31
59	A Short and Efficient Route to Enantiopure 3,5-Diarylpyrrolizidines. <i>Journal of Organic Chemistry</i> , 1999, 64, 1727-1732.	3.2	11
60	Synthesis and Reactivity of Bicycles Derived from Tartaric Acid and β -Amino Acids: A Novel Class of Conformationally Constrained Dipeptide Isosteres Based upon Enantiopure 3-Aza-6,8-dioxabicyclo[3.2.1]octane-7-carboxylic Acid. <i>Journal of Organic Chemistry</i> , 1999, 64, 7347-7364.	3.2	43
61	Stereoselectivity in the TiCl ₄ -catalysed reaction of Danishefsky's diene with a N-(acyloxy)iminium ion: Synthesis of 5 β versus 5 α 1(2)-19-Nor-10-azasteroids. 4. <i>Tetrahedron</i> , 1998, 54, 11589-11596.	1.9	10
62	Synthesis of benzo[c]quinolizin-3-ones: Selective non-steroidal inhibitors of steroid 5 β -reductase 1. <i>Bioorganic and Medicinal Chemistry Letters</i> , 1998, 8, 2871-2876.	2.2	21
63	A Concise Route to 19-Nor-10-azasteroids, a New Class of Steroid 5 β -Reductase Inhibitors. 3.1 Synthesis of (+)-19-Nor-10-azatestosterone and (+)-17 β -(Acetyloxy)-(5 β)-10-azaestr-1-en-3-one. <i>Journal of Organic Chemistry</i> , 1998, 63, 4111-4115.	3.2	24
64	Asymmetric hydrogenation of prochiral β -nitroketones by ruthenium complexes. <i>Journal of Molecular Catalysis A</i> , 1996, 110, 129-134.	4.8	3
65	Synthesis of enantiopure 2,7-diaryl-1,6-dioxaspiro[4.4]nonanes via enantioselective reduction of prochiral β -nitroketones by diisopinocampheylchloroborane (DIP-Cl ₂ , ϕ). <i>Tetrahedron: Asymmetry</i> , 1996, 7, 1929-1942.	1.8	9
66	Baker's yeast reduction of prochiral β -nitroketones. II.1 straightforward enantioselective synthesis of 2,7-dimethyl-1,6-dioxaspiro[4.4]nonanes. <i>Tetrahedron: Asymmetry</i> , 1995, 6, 2971-2976.	1.8	27
67	Baker's yeast reduction of prochiral β -nitroketones: Enantioselective synthesis of (S)-4-nitroalcohols. <i>Tetrahedron</i> , 1995, 51, 1775-1788.	1.9	24