

Santos Fustero

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.

214
papers

8,938
citations

39
h-index

88
g-index

299
ext. papers

10,148
ext. citations

5.7
avg, IF

6.11
L-index

#	Paper	IF	Citations
214	ASYMMETRIC CARBON-HALOGEN BOND FORMING REACTIONS (EXCLUDING C-H ACTIVATION PROCESSES) 2022 , 491-529		
213	Recent Advances on the Halo- and Cyano-Trifluoromethylation of Alkenes and Alkynes. <i>Molecules</i> , 2021 , 26,	4.8	1
212	Asymmetric Methods for Carbon-Fluorine Bond Formation. <i>European Journal of Organic Chemistry</i> , 2021 , 2021, 5946	3.2	3
211	Chemical Aspects of Human and Environmental Overload with Fluorine. <i>Chemical Reviews</i> , 2021 , 121, 4678-4742	68.1	49
210	Pentafluoroethylation of Carbonyl Compounds by HFC-125 the Encapsulation of the K Cation with Glymes. <i>Journal of Organic Chemistry</i> , 2021 , 86, 5883-5893	4.2	3
209	Diastereoselectivity of the Addition of Propargylic Magnesium Reagents to Fluorinated Aromatic Sulfinyl Imines. <i>Organic Letters</i> , 2021 , 23, 3691-3695	6.2	1
208	The Self-Disproportionation of Enantiomers (SDE): Fluorine as an SDE-Phoric Substituent 2021 , 281-306		
207	Selective Transformation of Norbornadiene into Functionalized Azaheterocycles and β -Amino Esters with Stereo- and Regiocontrol. <i>Chemistry - an Asian Journal</i> , 2021 , 16, 3873-3881	4.5	1
206	Tailor-Made Amino Acids and Fluorinated Motifs as Prominent Traits in Modern Pharmaceuticals. <i>Chemistry - A European Journal</i> , 2020 , 26, 11349-11390	4.8	50
205	Diastereoselective Synthesis of Enantioenriched Trifluoromethylated Ethylenediamines and Isoindolines Containing Two Stereogenic Carbon Centers by Nucleophilic Trifluoromethylation Using HFC-23. <i>Journal of Organic Chemistry</i> , 2020 , 85, 7976-7985	4.2	10
204	The Ruthenium-Catalyzed Domino Cross Enyne Metathesis/Ring-Closing Metathesis in the Synthesis of Enantioenriched Nitrogen-Containing Heterocycles. <i>European Journal of Organic Chemistry</i> , 2020 , 2020, 4193-4207	3.2	4
203	Nucleic acid recognition and antiviral activity of 1,4-substituted terphenyl compounds mimicking all faces of the HIV-1 Rev protein positively-charged β -helix. <i>Scientific Reports</i> , 2020 , 10, 7190	4.9	2
202	Fluorine-containing drugs approved by the FDA in 2019. <i>Chinese Chemical Letters</i> , 2020 , 31, 2401-2413	8.1	75
201	Asymmetric Synthesis of Fluorinated Monoterpenic Alkaloid Derivatives from Chiral Fluoroalkyl Aldimines via the Pauson-Khand Reaction. <i>Advanced Synthesis and Catalysis</i> , 2020 , 362, 1378-1384	5.6	6
200	Pauson-Khand reaction of fluorinated compounds. <i>Beilstein Journal of Organic Chemistry</i> , 2020 , 16, 1662-1682	15.82	7
199	The Fluoro-Pauson-Khand Reaction in the Synthesis of Enantioenriched Nitrogenated Bicycles Bearing a Quaternary C-F Stereogenic Center. <i>Organic Letters</i> , 2019 , 21, 7294-7297	6.2	5
198	Chemistry of detrifluoroacetylately in situ generated fluoro-enolates. <i>Organic and Biomolecular Chemistry</i> , 2019 , 17, 762-775	3.9	20

197	Asymmetric Vinylogous Mukaiyama-Mannich Reactions of Heterocyclic Siloxy Dienes with Ellman's Fluorinated Aldimines. <i>Advanced Synthesis and Catalysis</i> , 2019 , 361, 3860-3867	5.6	4
196	Fluorine-Containing Drugs Approved by the FDA in 2018. <i>Chemistry - A European Journal</i> , 2019 , 25, 11797-11812	4.8	13
195	Copper-Catalyzed Regioselective Synthesis of (E)-Fluorovinyl Sulfones. <i>Molecules</i> , 2019 , 24,	4.8	2
194	Vinyl Fluorides: Competent Olefinic Counterparts in the Intramolecular Pauson-Khand Reaction. <i>Organic Letters</i> , 2019 , 21, 2569-2573	6.2	7
193	Synthesis of polyfluoroalkyl sp-iminosugar glycolipids and evaluation of their immunomodulatory properties towards anti-tumor, anti-leishmanial and anti-inflammatory therapies. <i>European Journal of Medicinal Chemistry</i> , 2019 , 182, 111604	6.8	10
192	Tandem Organocatalytic Cycloaromatization/Intramolecular Friedel-Crafts Alkylation Sequence for the Synthesis of Indolizinones and Pyrrolo-azepinone Derivatives. <i>Journal of Organic Chemistry</i> , 2019 , 84, 10785-10795	4.2	5
191	Asymmetric synthesis of polycyclic 3-fluoroalkylproline derivatives by intramolecular azomethine ylide cycloaddition. <i>Organic Chemistry Frontiers</i> , 2019 , 6, 2916-2923	5.2	3
190	Biorelevant fluorine-containing N-heterocycles 2019 , 575-606		3
189	Metal-Free and User-Friendly Regioselective Hydroxyfluorination of Olefins. <i>Organic Letters</i> , 2018 , 20, 2338-2341	6.2	14
188	Fluorine-Containing Functionalized Cyclopentene Scaffolds Through Ring Contraction and Deoxyfluorination of Various Substituted Cyclohexenes. <i>European Journal of Organic Chemistry</i> , 2018 , 2018, 3735-3742	3.2	4
187	A metal-free and regioselective approach to (Z)-E-fluorovinyl sulfones and their chemoselective hydrogenation to E-fluoroalkyl sulfones. <i>Journal of Fluorine Chemistry</i> , 2018 , 206, 108-116	2.1	6
186	Recent advances in the synthesis of functionalised monofluorinated compounds. <i>Chemical Communications</i> , 2018 , 54, 9706-9725	5.8	46
185	Asymmetric Vinylogous Mannich-Type Addition of β -Dicyanoalkenes to E-fluoroalkyl Sulfinyl Imines. <i>Advanced Synthesis and Catalysis</i> , 2018 , 360, 366-373	5.6	10
184	Recent progress in the application of fluorinated chiral sulfinimine reagents. <i>Journal of Fluorine Chemistry</i> , 2018 , 216, 57-70	2.1	15
183	Ring-opening metathesis of some strained bicyclic systems; stereocontrolled access to diolefinated saturated heterocycles with multiple stereogenic centers. <i>Beilstein Journal of Organic Chemistry</i> , 2018 , 14, 2698-2707	2.5	5
182	Synthesis of fluorinated amino acid derivatives through late-stage deoxyfluorinations. <i>Tetrahedron</i> , 2018 , 74, 6367-6418	2.4	17
181	Dual Role of Vinyl Sulfonamides as N-Nucleophiles and Michael Acceptors in the Enantioselective Synthesis of Bicyclic E-sultams. <i>Advanced Synthesis and Catalysis</i> , 2018 , 360, 2885-2893	5.6	14
180	Sildenafil reduces neuroinflammation in cerebellum, restores GABAergic tone, and improves motor in-coordination in rats with hepatic encephalopathy. <i>CNS Neuroscience and Therapeutics</i> , 2017 , 23, 386-394	6.8	31

179	Fluorinated Chaperone- β -Cyclodextrin Formulations for β -Glucocerebrosidase Activity Enhancement in Neuronopathic Gaucher Disease. <i>Journal of Medicinal Chemistry</i> , 2017 , 60, 1829-1842	8.3	23
178	Olefin-Bond Chemodifferentiation through Cross-Metathesis Reactions: A Stereocontrolled Approach to Functionalized α,β -Amino Acid Derivatives. <i>European Journal of Organic Chemistry</i> , 2017 , 2017, 1894-1901	3.2	13
177	Intramolecular Nitrono Cycloaddition of α -(Trifluoromethyl)styrenes. Role of the CF Group in the Regioselectivity. <i>Journal of Organic Chemistry</i> , 2017 , 82, 2505-2514	4.2	18
176	Regio-specific synthesis of new 1-(tert-butyl)-1H-pyrazolecarboxamide derivatives. <i>Tetrahedron Letters</i> , 2017 , 58, 2441-2444	2	2
175	An Insight into Substrate-Dependent Fluorination of some Highly Substituted Alicyclic Scaffolds. <i>ChemistrySelect</i> , 2017 , 2, 3049-3052	1.8	3
174	Organocatalytic Enantioselective Synthesis of Trifluoromethyl-Containing Tetralin Derivatives by Sequential (Hetero)Michael Reaction β Intramolecular Nitrono Cycloaddition. <i>Advanced Synthesis and Catalysis</i> , 2017 , 359, 3752-3764	5.6	10
173	Homoallylic o-halobenzylamines: asymmetric diversity-oriented synthesis of benzo-fused cyclic amines. <i>Structural Chemistry</i> , 2017 , 28, 445-452	1.8	6
172	Fluorination of some highly functionalized cycloalkanes: chemoselectivity and substrate dependence. <i>Beilstein Journal of Organic Chemistry</i> , 2017 , 13, 2364-2371	2.5	6
171	Chemoselective, Substrate-directed Fluorination of Functionalized Cyclopentane β -Amino Acids. <i>Chemistry - an Asian Journal</i> , 2016 , 11, 3376-3381	4.5	10
170	Asymmetric Allylation/RCM-Mediated Synthesis of Fluorinated Benzo-Fused Bicyclic Homoallylic Amines As Dihydronaphthalene Derivatives. <i>Journal of Organic Chemistry</i> , 2016 , 81, 8876-8887	4.2	10
169	8-Iodonaphthalene-1-carbaldehyde: A Versatile Building Block for Diversity-Oriented Synthesis. <i>Organic Letters</i> , 2016 , 18, 4722-5	6.2	13
168	Recent Developments in the Chiral Brønsted Acid-catalyzed Allylboration Reaction with Polyfunctionalized Substrates. <i>Chemical Record</i> , 2016 , 16, 2046-60	6.6	16
167	Substrate-dependent fluorinations of highly functionalized cycloalkanes. <i>Tetrahedron</i> , 2016 , 72, 781-787	2.4	10
166	Asymmetric Synthesis of Monofluorinated 1-Amino-1,2-dihydronaphthalene and 1,3-Amino Alcohol Derivatives. <i>Organic Letters</i> , 2016 , 18, 948-51	6.2	23
165	Chiral Brønsted Acid-Catalyzed Asymmetric Allyl(propargyl)boration Reaction of ortho-Alkynyl Benzaldehydes: Synthetic Applications and Factors Governing the Enantioselectivity. <i>ACS Catalysis</i> , 2016 , 6, 2506-2514	13.1	39
164	Synthesis of 3-substituted isoindolin-1-ones via a tandem desilylation, cross-coupling, hydroamidation sequence under aqueous phase-transfer conditions. <i>Organic and Biomolecular Chemistry</i> , 2016 , 14, 85-92	3.9	30
163	A Stereocontrolled Protocol to Highly Functionalized Fluorinated Scaffolds through a Fluoride Opening of Oxiranes. <i>Molecules</i> , 2016 , 21,	4.8	9
162	Enantioselective Palladium-Catalyzed Oxidative β -Fluoroarylation of β -Unsaturated Carbonyl Derivatives. <i>Angewandte Chemie</i> , 2016 , 128, 9191-9195	3.6	15

161	Addition of Nucleophiles to Fluorinated Michael Acceptors. <i>European Journal of Organic Chemistry</i> , 2016 , 2016, 1751-1759	3.2	6
160	Gold-Catalyzed Povarov-Type Reaction of Fluorinated Imino Esters and Furans. <i>Journal of Organic Chemistry</i> , 2016 , 81, 6515-24	4.2	12
159	Enantioselective Palladium-Catalyzed Oxidative α -Fluoroarylation of β -Unsaturated Carbonyl Derivatives. <i>Angewandte Chemie - International Edition</i> , 2016 , 55, 9045-9	16.4	47
158	Diastereodivergent Synthesis of Fluorinated Cyclic β -Amino Acid Derivatives. <i>Organic Letters</i> , 2015 , 17, 5412-5	6.2	16
157	Silylboronates in the chiral Brønsted acid-catalysed allylboration of aldehydes. <i>Chemical Communications</i> , 2015 , 51, 5246-9	5.8	33
156	Olefin metathesis reactions with fluorinated substrates, catalysts, and solvents. <i>Chemical Reviews</i> , 2015 , 115, 871-930	68.1	131
155	Sildenafil reduces neuroinflammation and restores spatial learning in rats with hepatic encephalopathy: underlying mechanisms. <i>Journal of Neuroinflammation</i> , 2015 , 12, 195	10.1	51
154	Asymmetric Synthesis of Fluorinated Isoindolinones through Palladium-Catalyzed Carbonylative Amination of Enantioenriched Benzylic Carbamates. <i>Chemistry - A European Journal</i> , 2015 , 21, 11579-84	4.8	27
153	A Versatile Approach to CF ₃ -Containing 2-Pyrrolidones by Tandem Michael Addition-Cyclization: Exemplification in the Synthesis of Amidine Class BACE1 Inhibitors. <i>Chemistry - A European Journal</i> , 2015 , 21, 11719-26	4.8	14
152	Tandem cross enyne metathesis (CEYM)-intramolecular Diels-Alder reaction (IMDAR). An easy entry to linear bicyclic scaffolds. <i>Beilstein Journal of Organic Chemistry</i> , 2015 , 11, 1486-93	2.5	6
151	Gold-catalyzed tandem hydroamination/formal aza-Diels-Alder reaction of homopropargyl amino esters: a combined computational and experimental mechanistic study. <i>Chemistry - A European Journal</i> , 2015 , 21, 5459-66	4.8	12
150	Differential reactivity of fluorinated homopropargylic amino esters vs gold(I) salts. The role of the nitrogen protecting group. <i>Journal of Fluorine Chemistry</i> , 2015 , 171, 60-66	2.1	8
149	A novel and selective fluoride opening of aziridines by XtalFluor-E. synthesis of fluorinated diamino acid derivatives. <i>Organic Letters</i> , 2015 , 17, 1074-7	6.2	38
148	Asymmetric intramolecular aza-Michael reaction in desymmetrization processes. Total synthesis of hippodamine and epi-hippodamine. <i>Organic Letters</i> , 2015 , 17, 960-3	6.2	24
147	Asymmetric allylation/Pauson-Khand reaction: a simple entry to polycyclic amines. Application to the synthesis of aminosteroid analogues. <i>Organic Letters</i> , 2014 , 16, 1224-7	6.2	30
146	Diastereoselective synthesis of 2-phenyl-3-(trifluoromethyl)piperazines as building blocks for drug discovery. <i>Journal of Organic Chemistry</i> , 2014 , 79, 5887-94	4.2	17
145	A Selective Synthesis of Fluorinated Cispentacin Derivatives. <i>European Journal of Organic Chemistry</i> , 2014 , 2014, 4070-4076	3.2	17
144	Fluorine in pharmaceutical industry: fluorine-containing drugs introduced to the market in the last decade (2001-2011). <i>Chemical Reviews</i> , 2014 , 114, 2432-506	68.1	2974

- 143 Microwave-assisted tandem organocatalytic peptide-coupling intramolecular aza-Michael reaction: β -unsaturated N-acyl pyrazoles as Michael acceptors. *Chemistry - A European Journal*, **2014**, 20, 15697-7014. 4.8 15
- 142 Tandem gold self-relay catalysis for the synthesis of 2,3-dihydropyridin-4(1H)-ones: combination of β -keto ester and Lewis acid properties of gold salts. *Chemistry - A European Journal*, **2014**, 20, 14126-31. 4.8 18
- 141 Fluorinated Pyrazoles and Indazoles **2014**, 279-321. 7
- 140 Synthesis of Fluorinated and Nonfluorinated Tebufenpyrad Analogues for the Study of Anti-angiogenesis MOA. *Organic Process Research and Development*, **2014**, 18, 1027-1036. 3.9 14
- 139 Biochemical quantitation of the eIF5A hypusination in Arabidopsis thaliana uncovers ABA-dependent regulation. *Frontiers in Plant Science*, **2014**, 5, 202. 6.2 8
- 138 Unique reactivity of fluorinated molecules with transition metals. *Chimia*, **2014**, 68, 382-409. 1.3 13
- 137 An Approach to 2,4-Substituted Pyrazolo[1,5-a]pyridines and Pyrazolo[1,5-a]azepines by Ring-Closing Metathesis. *European Journal of Organic Chemistry*, **2013**, 2013, 7164-7174. 3.2 10
- 136 Base-dependent stereodivergent intramolecular aza-Michael reaction: asymmetric synthesis of 1,3-disubstituted isoindolines. *Chemistry - A European Journal*, **2013**, 19, 11776-85. 4.8 44
- 135 Gold catalyzed stereoselective tandem hydroamination-formal aza-Diels-Alder reaction of propargylic amino esters. *Chemical Communications*, **2013**, 49, 1336-8. 5.8 30
- 134 Asymmetric Tandem Reactions: New Strategies and Applications. *Phosphorus, Sulfur and Silicon and the Related Elements*, **2013**, 188, 331-339. 1
- 133 Relay Catalysis: Enantioselective Synthesis of Cyclic Benzo-Fused Homoallylic Alcohols by Chiral Brønsted Acid-Catalyzed Allylboration/Ring Closing Metathesis. *Advanced Synthesis and Catalysis*, **2013**, 355, 1058-1064. 5.6 38
- 132 Synthesis and application of β -substituted Pauson-Khand adducts: trifluoromethyl as a removable steering group. *Angewandte Chemie - International Edition*, **2013**, 52, 5355-9. 16.4 21
- 131 Gold-catalyzed intramolecular hydroamination of o-alkynylbenzyl carbamates: a route to chiral fluorinated isoindoline and isoquinoline derivatives. *Organic Letters*, **2013**, 15, 832-5. 6.2 69
- 130 Asymmetric allylation/ring closing metathesis: one-pot synthesis of benzo-fused cyclic homoallylic amines. Application to the formal synthesis of Sertraline derivatives. *Organic Letters*, **2013**, 15, 3770-3. 6.2 21
- 129 Structure-based design of an RNA-binding p-terphenylene scaffold that inhibits HIV-1 Rev protein function. *Angewandte Chemie - International Edition*, **2013**, 52, 13405-9. 16.4 21
- 128 Synthesis and Application of β -Substituted Pauson-Khand Adducts: Trifluoromethyl as a Removable Steering Group. *Angewandte Chemie*, **2013**, 125, 5463-5467. 3.6 8
- 127 One-pot cross-ene-yne metathesis (CEYM)-Diels-Alder reaction of gem-difluoropropargylic alkynes. *Beilstein Journal of Organic Chemistry*, **2013**, 9, 2688-95. 2.5 7
- 126 Efficient regio- and stereoselective access to novel fluorinated β -aminocyclohexanecarboxylates. *Beilstein Journal of Organic Chemistry*, **2013**, 9, 1164-9. 2.5 12

125	Efficacy and activity prediction by molecular topology of new drugs against the <i>Tetranychus urticae</i> plague. <i>Combinatorial Chemistry and High Throughput Screening</i> , 2013 , 16, 473-83	1.3	1
124	A practical entry to α -alkyl amino alcohols: application to the synthesis of a potent BACE1 inhibitor. <i>Organic and Biomolecular Chemistry</i> , 2012 , 10, 6758-66	3.9	10
123	Stereoselective access to fluorinated and non-fluorinated quaternary piperidines: synthesis of pipercolic acid and iminosugar derivatives. <i>Chemistry - A European Journal</i> , 2012 , 18, 3753-64	4.8	25
122	1,7-octadiene-assisted tandem multicomponent cross-enyne metathesis (CEYM)-Diels-Alder reactions: a useful alternative to Mori's conditions. <i>Chemistry - A European Journal</i> , 2012 , 18, 10991-7	4.8	24
121	Synthetic and biological applications of fluorous reagents as phase tags. <i>Topics in Current Chemistry</i> , 2012 , 308, 45-67		6
120	A New Tandem Cross Metathesis-Intramolecular Aza-Michael Reaction for the Synthesis of α,α -Difluorinated Lactams. <i>Synthesis</i> , 2012 , 44, 1863-1873	2.9	24
119	Intramolecular Michael reaction of tert-butylsulfinyl ketimines: asymmetric synthesis of 3-substituted indanones. <i>Organic Letters</i> , 2011 , 13, 6564-7	6.2	32
118	Organocatalytic enantioselective synthesis of quinolizidine alkaloids (+)-myrtine, (+)-lupinine, and (+)-epiepipiquinamide. <i>Tetrahedron</i> , 2011 , 67, 7412-7417	2.4	31
117	From 2000 to mid-2010: a fruitful decade for the synthesis of pyrazoles. <i>Chemical Reviews</i> , 2011 , 111, 6984-7034	68.1	758
116	Asymmetric synthesis of quaternary α -amino acid derivatives and their fluorinated analogues. <i>Amino Acids</i> , 2011 , 41, 559-73	3.5	14
115	Selective Synthesis of New Fluorinated Alicyclic α -Amino Ester Stereoisomers. <i>European Journal of Organic Chemistry</i> , 2011 , 2011, 4993-5001	3.2	30
114	Synthesis of Fluorinated and Non-Fluorinated Bicyclic Amidines through Ring-Closing Metathesis. <i>European Journal of Organic Chemistry</i> , 2011 , 2011, 7317-7323	3.2	5
113	New cathepsin inhibitors to explore the fluorophilic properties of the S2 pocket of cathepsin B: design, synthesis, and biological evaluation. <i>Chemistry - A European Journal</i> , 2011 , 17, 5256-60	4.8	13
112	Chiral monofluorobenzyl carbanions: synthesis of enantiopure α -fluorinated β -phenylethylamines. <i>Chemistry - A European Journal</i> , 2011 , 17, 6142-7	4.8	22
111	Microwave-assisted organocatalytic enantioselective intramolecular aza-Michael reaction with α,β -unsaturated ketones. <i>Chemistry - A European Journal</i> , 2011 , 17, 14267-72	4.8	46
110	Design, synthesis, and biological evaluation of novel fluorinated ethanolamines. <i>Chemistry - A European Journal</i> , 2011 , 17, 14772-84	4.8	13
109	Regio- and diastereoselective fluorination of alicyclic α -amino acids. <i>Organic and Biomolecular Chemistry</i> , 2011 , 9, 6528-34	3.9	30
108	Synthesis of Fluorinated α -Amino Acids. <i>Synthesis</i> , 2011 , 2011, 3045-3079	2.9	35

107	Asymmetric tandem reactions: New synthetic strategies. <i>Pure and Applied Chemistry</i> , 2010 , 82, 669-677	2.1	31
106	A Mild, Efficient Synthesis of gem-Difluorodihydrouracils. <i>Synthesis</i> , 2010 , 2010, 651-660	2.9	1
105	Diastereoselective intramolecular additions of allyl- and propargylsilanes to iminium ions: synthesis of cyclic and bicyclic quaternary amino acids. <i>Organic Letters</i> , 2010 , 12, 3014-7	6.2	22
104	Regioselectivity in intermolecular Pauson-Khand reactions of dissymmetric fluorinated alkynes. <i>Organic Letters</i> , 2010 , 12, 5620-3	6.2	19
103	Tandem asymmetric Michael reaction-intramolecular Michael addition. An easy entry to chiral fluorinated 1,4-dihydropyridines. <i>Organic Letters</i> , 2010 , 12, 3484-7	6.2	44
102	Recent Developments in the Synthesis of Fluorinated α -Amino Acids. <i>Current Organic Chemistry</i> , 2010 , 14, 928-949	1.7	69
101	Tandem nucleophilic addition-intramolecular aza-Michael reaction: facile synthesis of chiral fluorinated isoindolines. <i>Organic Letters</i> , 2010 , 12, 5494-7	6.2	70
100	N-sulfinyl amines as a nitrogen source in the asymmetric intramolecular aza-Michael reaction: total synthesis of (-)-pinidinol. <i>Chemistry - A European Journal</i> , 2010 , 16, 9835-45	4.8	68
99	Nitrogen-Containing Organofluorine Derivatives: An Overview. <i>Synlett</i> , 2009 , 2009, 525-549	2.2	61
98	Organocatalytic anti-Selective Mannich Reactions with Fluorinated Aldimines: Synthesis of anti-Fluoroalkyl- β -amino Alcohols. <i>European Journal of Organic Chemistry</i> , 2009 , 2009, 5208-5214	3.2	24
97	A new strategy for the synthesis of fluorinated 3,4-dihydropyrimidinones. <i>Journal of Fluorine Chemistry</i> , 2009 , 130, 1145-1150	2.1	5
96	Synthesis and enzymatic evaluation of novel partially fluorinated thiol dual ACE/NEP inhibitors. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2009 , 19, 4715-9	2.9	6
95	Recent Advances in the Synthesis of Pyrazoles. A Review. <i>Organic Preparations and Procedures International</i> , 2009 , 41, 253-290	1.1	167
94	Fluorous TBAF: a convenient and selective reagent for fluoride-mediated deprotections. <i>Journal of Organic Chemistry</i> , 2009 , 74, 6398-401	4.2	8
93	AuX ₃ -mediated selective head-to-head dimerization of difluoropropargyl amides. <i>Journal of Organic Chemistry</i> , 2009 , 74, 7690-6	4.2	17
92	Cross-metathesis reactions as an efficient tool in the synthesis of fluorinated cyclic β -amino acids. <i>Journal of Organic Chemistry</i> , 2009 , 74, 3414-23	4.2	34
91	Straightforward stereoselective access to cyclic peptidomimetics. <i>Journal of Organic Chemistry</i> , 2009 , 74, 4429-32	4.2	35
90	New Fluorinated Pyrazol and Uracil Derivatives: Synthesis and Biological Activity. <i>ACS Symposium Series</i> , 2009 , 182-209	0.4	7

89	Solution versus fluorous versus solid-phase synthesis of 2,5-disubstituted 1,3-azoles. Preliminary antibacterial activity studies. <i>Journal of Organic Chemistry</i> , 2009 , 74, 8988-96	4.2	57
88	New fluorinated peptidomimetics through tandem aza-michael addition to alpha-trifluoromethyl acrylamide acceptors: synthesis and conformational study in solid state and solution. <i>Journal of Organic Chemistry</i> , 2009 , 74, 3122-32	4.2	18
87	Asymmetric synthesis of indolines through intramolecular shifting of aromatic sulfinyl groups. Role of the pi,pi-stacking interactions in these unusual S(N)Ar processes. <i>Journal of the American Chemical Society</i> , 2009 , 131, 9432-41	16.4	35
86	A new strategy for the synthesis of optically pure beta-fluoroalkyl beta-amino acid derivatives. <i>Organic Letters</i> , 2009 , 11, 641-4	6.2	33
85	Synthesis of new fluorinated Tebufenpyrad analogs with acaricidal activity through regioselective pyrazole formation. <i>Journal of Organic Chemistry</i> , 2008 , 73, 8545-52	4.2	105
84	An efficient entry to optically active anti- and syn-beta-amino-alpha-trifluoromethyl alcohols. <i>Organic Letters</i> , 2008 , 10, 605-8	6.2	26
83	Concise preparation of 2,2-difluorohomopropargyl carbonyl derivatives. Application to the synthesis of 4,4-difluoroisoquinolinone congeners. <i>Journal of Organic Chemistry</i> , 2008 , 73, 2656-61	4.2	38
82	Solution and fluorous phase synthesis of beta,beta-difluorinated 1-amino-1-cyclopentane carboxylic acid derivatives. <i>Journal of Fluorine Chemistry</i> , 2008 , 129, 943-950	2.1	12
81	Solution-, solid-phase, and fluorous synthesis of beta,beta-difluorinated cyclic quaternary alpha-amino acid derivatives: a comparative study. <i>Chemistry - A European Journal</i> , 2008 , 14, 7019-29	4.8	28
80	Organocatalytic approach to benzofused nitrogen-containing heterocycles: enantioselective total synthesis of (+)-angustureine. <i>Chemistry - A European Journal</i> , 2008 , 14, 9868-72	4.8	111
79	Anionic-anionic asymmetric tandem reactions: one-pot synthesis of optically pure fluorinated indolines from 2-p-tolylsulfinyl alkylbenzenes. <i>Angewandte Chemie - International Edition</i> , 2008 , 47, 7941-4	16.4	47
78	Anionic-Anionic Asymmetric Tandem Reactions: One-Pot Synthesis of Optically Pure Fluorinated Indolines from 2-p-Tolylsulfinyl Alkylbenzenes. <i>Angewandte Chemie</i> , 2008 , 120, 8059-8062	3.6	7
77	An efficient synthesis of uracil derivatives from 2-alkyl-2-oxazolines and nitriles. <i>Journal of Fluorine Chemistry</i> , 2008 , 129, 836-847	2.1	2
76	Improved regioselectivity in pyrazole formation through the use of fluorinated alcohols as solvents: synthesis and biological activity of fluorinated tebufenpyrad analogs. <i>Journal of Organic Chemistry</i> , 2008 , 73, 3523-9	4.2	154
75	Selective formal transesterification of fluorinated 2-(trimethylsilyl)ethyl alpha-imino esters mediated by TBAF. <i>Journal of Organic Chemistry</i> , 2008 , 73, 5617-20	4.2	5
74	Tiratricol neutralizes bacterial endotoxins and reduces lipopolysaccharide-induced TNF-alpha production in the cell. <i>Chemical Biology and Drug Design</i> , 2008 , 72, 320-8	2.9	3
73	Asymmetric synthesis of fluorinated amino macrolactones through ring-closing metathesis. <i>Journal of Organic Chemistry</i> , 2007 , 72, 8716-23	4.2	12
72	The role of fluorine in the stereoselective tandem aza-Michael addition to acrylamide acceptors: an experimental and theoretical mechanistic study. <i>Chemistry - A European Journal</i> , 2007 , 13, 8530-42	4.8	15

71	Enantioselective organocatalytic intramolecular aza-Michael reaction: a concise synthesis of (+)-sedamine, (+)-allosedamine, and (+)-coniine. <i>Organic Letters</i> , 2007 , 9, 5283-6	6.2	157
70	Synthesis of a new fluorinated oxazolidinone and its reactivity as a chiral auxiliary in Aldol reactions. <i>Journal of Fluorine Chemistry</i> , 2007 , 128, 647-653	2.1	9
69	Synthesis of fluorinated allylic amines: Reaction of 2-(trimethylsilyl)ethyl sulfones and sulfoxides with fluorinated imines. <i>Journal of Fluorine Chemistry</i> , 2007 , 128, 1248-1254	2.1	6
68	Nitrogen-Containing Organofluorine Compounds through Metathesis Reactions. <i>ACS Symposium Series</i> , 2007 , 54-68	0.4	1
67	Microwave-assisted tandem cross metathesis intramolecular Aza-Michael reaction: an easy entry to cyclic beta-amino carbonyl derivatives. <i>Journal of the American Chemical Society</i> , 2007 , 129, 6700-1	16.4	119
66	Intramolecular hydroamination of difluoropropargyl amides: regioselective synthesis of fluorinated beta- and gamma-lactams. <i>Organic Letters</i> , 2007 , 9, 4251-3	6.2	64
65	An Enantio- and Diastereoselective Synthesis of Fluorinated β -Aminoalkylpiperine Derivatives through Mannich and Ring-Closing Metathesis Reactions. <i>Synthesis</i> , 2006 , 2006, 4087-4091	2.9	13
64	Asymmetric synthesis of fluorinated cyclic beta-amino acid derivatives through cross metathesis. <i>Organic Letters</i> , 2006 , 8, 4633-6	6.2	36
63	Role of the gem-difluoro moiety in the tandem ring-closing metathesis-olefin isomerization: regioselective preparation of unsaturated lactams. <i>Journal of Organic Chemistry</i> , 2006 , 71, 2706-14	4.2	75
62	Fluorous (trimethylsilyl)ethanol: a new reagent for carboxylic acid tagging and protection in peptide synthesis. <i>Journal of Organic Chemistry</i> , 2006 , 71, 3299-302	4.2	39
61	Asymmetric synthesis of new beta,beta-difluorinated cyclic quaternary alpha-amino acid derivatives. <i>Organic Letters</i> , 2006 , 8, 4129-32	6.2	40
60	First Fluorous Synthesis of Fluorinated Uracils. <i>QSAR and Combinatorial Science</i> , 2006 , 25, 753-760		16
59	New fluorinated 1,3-vinylogous amidines as versatile intermediates: synthesis of fluorinated pyrimidin-2(1H)-ones. <i>Tetrahedron</i> , 2006 , 62, 1444-1451	2.4	10
58	Synthesis and biological evaluation of new bicyclic fluorinated uracils through ring-closing metathesis. <i>Journal of Organic Chemistry</i> , 2006 , 71, 4010-3	4.2	17
57	1,4-Benzodiazepine N-nitrosoamidines: useful intermediates in the synthesis of tricyclic benzodiazepines. <i>Molecules</i> , 2006 , 11, 583-8	4.8	17
56	Highly enantioselective synthesis of fluorinated gamma-amino alcohols through proline-catalyzed cross-Mannich reaction. <i>Organic Letters</i> , 2005 , 7, 3433-6	6.2	82
55	Stereoselective Synthesis of Fluorine-Containing β -Amino Acids 2005 , 319-350		0
54	Fluorinated β -Enamino Esters as Versatile Synthetic Intermediates: Synthesis of Fluorinated β -Amino Acids and Uracils. <i>ACS Symposium Series</i> , 2005 , 593-610	0.4	0

53	New approaches to the synthesis of organofluorine nitrogenated derivatives. <i>Journal of Fluorine Chemistry</i> , 2004 , 125, 621-627	2.1	22
52	A versatile synthesis of fluorinated uracils in solution and on solid-phase. <i>Organic Letters</i> , 2004 , 6, 1417-202		34
51	Highly Stereoselective Tandem Aza-Michael Addition-Enolate Protonation to Form Partially Modified Retropeptide Mimetics Incorporating a Trifluoroalanine Surrogate. <i>Angewandte Chemie</i> , 2003 , 115, 2106-2109	3.6	8
50	Highly stereoselective tandem aza-Michael addition-enolate protonation to form partially modified retropeptide mimetics incorporating a trifluoroalanine surrogate. <i>Angewandte Chemie - International Edition</i> , 2003 , 42, 2060-3	16.4	56
49	Stereocontrolled solid-phase synthesis of fluorinated partially-modified retropeptides via tandem aza-Michael/enolate-protonation. <i>Tetrahedron Letters</i> , 2003 , 44, 7019-7022	2	14
48	Novel approach for asymmetric synthesis of fluorinated beta-amino sulfones and allylic amines. <i>Organic Letters</i> , 2003 , 5, 2707-10	6.2	35
47	Diastereoselective synthesis of fluorinated, seven-membered beta-amino acid derivatives via ring-closing metathesis. <i>Organic Letters</i> , 2003 , 5, 2523-6	6.2	42
46	An efficient synthesis of new fluorinated uracil derivatives. <i>Chemical Communications</i> , 2003 , 844-845	5.8	14
45	New strategy for the stereoselective synthesis of fluorinated beta-amino acids. <i>Journal of Organic Chemistry</i> , 2002 , 67, 4667-79	4.2	62
44	A concise, asymmetric synthesis of tetramic acid derivatives. <i>Organic Letters</i> , 2002 , 4, 3651-4	6.2	14
43	A new and expeditious strategy for the synthesis of β -amino acids from α -oxazolines. <i>Tetrahedron</i> , 2001 , 57, 703-712	2.4	10
42	Novel strategy for the synthesis of fluorinated β -amino acid derivatives from α -oxazolines. <i>Tetrahedron</i> , 2001 , 57, 6475-6486	2.4	21
41	C α versus C β Annulation Reactions of 2-Alkyl-2-oxazolines and 2-Alkyl-2-thiazolines: A Simple Synthesis of Novel 3-Aminoindene, Phthalimidine, Pyrrolidine, and Piperidine Derivatives. <i>European Journal of Organic Chemistry</i> , 2001 , 2001, 1195-1201	3.2	6
40	Synthesis of Nonracemic β -Trifluoromethyl β -Amino Acids from Sulfinimines of Trifluoropyruvate. <i>European Journal of Organic Chemistry</i> , 2001 , 2001, 1449-1458	3.2	52
39	Enantioselective synthesis of fluorinated alpha-amino acids and derivatives in combination with ring-closing metathesis: intramolecular pi-stacking interactions as a source of stereocontrol. <i>Organic Letters</i> , 2001 , 3, 2621-4	6.2	50
38	Synthesis of Nonracemic β -Trifluoromethyl β -Amino Acids from Sulfinimines of Trifluoropyruvate 2001 , 2001, 1449		1
37	1-Chloro-1,1-difluoro-N-(4-methoxyphenyl)-3-(pyrrolidin-2-ylidene)propan-2-imine. <i>Acta Crystallographica Section C: Crystal Structure Communications</i> , 2000 , 56, E213-4		1
36	Stereoselective Syntheses of Allylic Amines Through Reduction of 1-Azadiene Intermediates. <i>Tetrahedron</i> , 2000 , 56, 8179-8187	2.4	7

- 35 A new and mild synthesis of β -amino acids from masked β -amino acid derivatives. *Tetrahedron Letters*, **1999**, 40, 1005-1008 2 5
- 34 Synthesis and C-alkylation of C-protected β -amino acid derivatives from oxa- thia- and imidazolines. *Tetrahedron*, **1999**, 55, 2695-2712 2.4 11
- 33 Stereoselective Mannich-Type Reaction of an Acyclic Ketimine with a Substituted Chlorotitanium Enolate: Efficient Approach to d-erythro- β -trifluoromethylhydroxyaspartic Units. *Journal of Organic Chemistry*, **1999**, 64, 8731-8735 4.2 52
- 32 New Strategies for the Synthesis of Fluorinated Vinylogous Amidines and beta-Enamino Ketones. *Journal of Organic Chemistry*, **1999**, 64, 5551-5556 4.2 35
- 31 First Highly Diastereoselective Synthesis of syn β -Methyl β -Fluoroalkyl β -Amino Esters. *Organic Letters*, **1999**, 1, 977-980 6.2 40
- 30 Two Practical and Efficient Approaches to Fluorinated and Nonfluorinated Chiral beta-Imino Sulfoxides. *Journal of Organic Chemistry*, **1998**, 63, 6210-6219 4.2 26
- 29 Synthesis and Reactivity of New β -Enamino Acid Derivatives: A Simple and General Approach to β -Enamino Esters and Thioesters. *Journal of Organic Chemistry*, **1998**, 63, 8825-8836 4.2 36
- 28 A new and general synthesis of N-substituted fluorinated β -aminosulfoxides. *Tetrahedron Letters*, **1997**, 38, 4891-4894 2 8
- 27 N-Substituted β -Enamino Acid Derivatives: A New Approach to Fluorinated β -Enamino Esters. *Tetrahedron Letters*, **1997**, 38, 6771-6774 2 18
- 26 An Efficient and Simple Entry to N-Substituted beta-Enamino Acid Derivatives from 2-Alkyl-2-oxazolines and 2-Alkyl-2-thiazolines. *Journal of Organic Chemistry*, **1996**, 61, 8849-8859 4.2 30
- 25 Synthesis and NMR configurational study of imidazo[2,1-b]thiazoles from 1H-1,4-diazepine-7(6H)-thiones. *Tetrahedron*, **1993**, 49, 6619-6626 2.4 3
- 24 A new method for the regioselective synthesis of β -amino acid derivatives. *Tetrahedron Letters*, **1993**, 34, 725-728 2 10
- 23 Stereoselective synthesis of 1,3-amino alcohols and 1,3-amino ketones. *Journal of Organic Chemistry*, **1992**, 57, 1219-1223 4.2 36
- 22 Synthesis and ¹H-NMR configurational study of β -thiazolines from 2-aza-1,3-dienes. *Tetrahedron*, **1992**, 48, 9745-9752 2.4 1
- 21 Synthesis and reactivity of β -amino- β -unsaturated oxa- and thiazolines. *Tetrahedron Letters*, **1992**, 33, 3801-3804 2 16
- 20 Synthesis of 1,3-Amino Alcohols from 2-Aza-1,3-dienes by Reduction of 5,6-Dihydro-2H-1,3-oxazines. *Synthesis*, **1991**, 1991, 387-393 2.9 7
- 19 Tandem cycloaddition/rearrangement of 2-aza-1,3-dienes. A simple and efficient synthesis of 1H-1,4-diazepine-7(6H)-thiones. *Journal of the Chemical Society Chemical Communications*, **1991**, 1704-1705 5
- 18 2-Aza 1,3-dienes: a new and simple method for the synthesis of functionalized pyridine derivatives. *Journal of Organic Chemistry*, **1991**, 56, 6751-6754 4.2 17

17	New perspectives of carbo- and hetero-1,3-dienes in organic synthesis. <i>Pure and Applied Chemistry</i> , 1990 , 62, 1957-1966	2.1	18
16	Unactivated 2-Aza-1,3-dienes: Halogenation and face selectivity in diels-alder reactions. <i>Tetrahedron Letters</i> , 1990 , 31, 397-398	2	11
15	Synthesis of 4(1H)-pyridones by carbonylation of 2-Aza-1,3-Dienes.. <i>Tetrahedron Letters</i> , 1990 , 31, 3793-3796		9
14	An exceptionally simple and unexpected synthesis of new 4,6-diazasemibullvalenes. <i>Journal of the Chemical Society Chemical Communications</i> , 1990 , 1057-1059		1
13	Diels-alder reaction of unactivated 2-aza-1,3-dienes with diethyl ketomalonate: A carbon dioxide equivalent. <i>Tetrahedron Letters</i> , 1989 , 30, 2685-2688	2	8
12	Reduction of 5,6-dihydro-2H-1,3-oxazines. A simple approach to 1,3-aminoalcohols from 2-aza-1,3-dienes. <i>Tetrahedron Letters</i> , 1989 , 30, 2001-2004	2	5
11	A simple stereoselective synthesis of primary allylic amines from 4-amino-1-azadienes. <i>Journal of the Chemical Society Chemical Communications</i> , 1989 , 1132		11
10	A simple regiospecific synthesis of substituted pyridines from 2-aza-1,3-dienes. <i>Journal of Organic Chemistry</i> , 1988 , 53, 5960-5963	4.2	16
9	An efficient and general strategy for the synthesis of 1,4-dihydro- β - and β -azaphosphinines from 2-aza-1,3-dienes. <i>Journal of the Chemical Society Chemical Communications</i> , 1988 , 1596-1597		11
8	Diastereo- and enantio-selective synthesis of dihydro- and tetrahydro-pyrimidines. A new strategy for the asymmetric synthesis of β -amino ketones and β -amino alcohols. <i>Journal of the Chemical Society Chemical Communications</i> , 1988 , 410b-412		7
7	Cycloaddition of unactivated 2-aza-1,3-dienes with heterocumulenes: a convenient route to the synthesis of 1,3-difunctionalized compounds. <i>Journal of the Chemical Society Perkin Transactions 1</i> , 1988 , 1739-1744		10
6	Synthesis of N-Aryl-substituted 2-Aminoalkyl Ketones and 1,3-Alkanediamines. <i>Synthesis</i> , 1986 , 1986, 469-473	2.9	7
5	Silylation of 2-Aza-1,3-dienes. The first example of a thermally stable N-trimethyl-silyldivinyllamine. <i>Journal of the Chemical Society Chemical Communications</i> , 1986 , 361		6
4	Diels-Alder cycloaddition reaction of unactivated 2-aza-1,3-dienes with dialkyl azodicarboxylates and heterocumulenes. <i>Journal of the Chemical Society Chemical Communications</i> , 1986 , 1179-1180		12
3	Preparation and reactivity of 2-Aza-1,3-butadienes: A Diels-Alder route to 5,6-dihydro-2H-1,3-oxazine derivatives. <i>Chemische Berichte</i> , 1985 , 118, 3652-3663		25
2	Diastereoselective synthesis of γ -amino alcohols with three chiral centers by reduction of β -amino ketones and derivatives. <i>Journal of Organic Chemistry</i> , 1985 , 50, 4052-4056	4.2	22
1	Reduction of 1,3-diimines. A new and general method of synthesis of γ -diamines, β -amino ketones, and derivatives with two and three chiral centers. <i>Journal of Organic Chemistry</i> , 1983 , 48, 2255-2259	4.2	15