

# Sulfur(VI) Fluoride Exchange (SuFEx): Another Good Re

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Citation Report

#	ARTICLE	IF	CITATIONS
3	Nâ€Heterocyclic Carbene Catalyzed Synthesis of Î€Sultones via Î±,Î²â€Unsaturated Sulfonyl Azolium Intermediates. <i>Angewandte Chemie - International Edition</i> , 2015, 54, 11780-11784.	7.2	60
4	SuFEx on the Surface: A Flexible Platform for Postpolymerization Modification of Polymer Brushes. <i>Angewandte Chemie - International Edition</i> , 2015, 54, 13370-13373. Gesellschaft Deutscher Chemiker Awards: Adolf von Baeyer Memorial Medal for Carsten Bolm / WÃ¼hler Prize for Sustainable Chemistry to Matthias Beller / Karl Ziegler Prize for Helmut Schwarz /	7.2	99
7	Honorary Membership for Henning Hopf / August Wilhelm von Hofmann Lectureship for K.â€...Barry Sharpless / Wilhelm Klemm Prize for Thomasâ€...F. FÃssler / Heinz Schmidkuntz Prize for Michael Tausch / Arfvedsonâ€Schlenk Prize for Clareâ€...P. Grey / Dr. Hermann Schnell Fellowship for Jiayin Yuan / Fresenius Prize for Renato Zenobi / V. <i>Angewandte Chemie - International Edition</i> , 2015, 54, 10068-10071.	7.2	0
8	Recent Advances in Click Chemistry Applied to Dendrimer Synthesis. <i>Molecules</i> , 2015, 20, 9263-9294.	1.7	112
9	Covalent Molecular Probes for Class A G Protein-Coupled Receptors: Advances and Applications. <i>ACS Chemical Biology</i> , 2015, 10, 1376-1386.	1.6	53
10	Transition-metal-free persulfuration to construct unsymmetrical disulfides and mechanistic study of the sulfur redox process. <i>Chemical Communications</i> , 2015, 51, 4208-4211.	2.2	90
11	Lateâ€Stage Fluorination: Fancy Novelty or Useful Tool?. <i>Angewandte Chemie - International Edition</i> , 2015, 54, 3216-3221.	7.2	219
12	Rational Targeting of Active-Site Tyrosine Residues Using Sulfonyl Fluoride Probes. <i>ACS Chemical Biology</i> , 2015, 10, 1094-1098.	1.6	153
14	Sulfonyl fluorides as privileged warheads in chemical biology. <i>Chemical Science</i> , 2015, 6, 2650-2659.	3.7	383
15	Nickel- and Palladium-Catalyzed Coupling of Aryl Fluorosulfonates with Aryl Boronic Acids Enabled by Sulfonyl Fluoride. <i>ACS Catalysis</i> , 2015, 5, 5041-5046.	5.5	78
16	PyFluor: A Low-Cost, Stable, and Selective Deoxyfluorination Reagent. <i>Journal of the American Chemical Society</i> , 2015, 137, 9571-9574.	6.6	222
17	Development and Application of O-(Trimethylsilyl)aryl Fluorosulfates for the Synthesis of Arynes. <i>Journal of Organic Chemistry</i> , 2015, 80, 6890-6896.	1.7	37
18	An efficient click-multicomponent strategy for the diversity oriented synthesis of 15â€18 membered macrocyclic peptidomimetic fluorophores. <i>Tetrahedron Letters</i> , 2015, 56, 2451-2455.	0.7	6
19	Cell permeable affinity- and activity-based probes. <i>Future Medicinal Chemistry</i> , 2015, 7, 2131-2141.	1.1	23
20	Combination of AGET ATRP and SuFEx for post-polymerization chain-end modifications. <i>Polymer</i> , 2015, 78, 37-41.	1.8	41
21	Selective Access to Heterocyclic Sulfonamides and Sulfonyl Fluorides via a Parallel Medicinal Chemistry Enabled Method. <i>ACS Combinatorial Science</i> , 2015, 17, 653-657.	3.8	20
22	Click chemistry patents and their impact on drug discovery and chemical biology. <i>Pharmaceutical Patent Analyst</i> , 2015, 4, 109-119.	0.4	6
23	Direct introduction of R-SO <sub>2</sub> F moieties into proteins and protein-polymer conjugation using SuFEx chemistry. <i>Polymer</i> , 2016, 99, 7-12.	1.8	35

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24	General Cycloaddition Between a Trimethylsilyl-Capped Alkyne and an Azide Catalyzed by an N-Heterocyclic Carbene-Copper Complex and Pyridine-Bis-carboxamide. <i>Advanced Synthesis and Catalysis</i> , 2016, 358, 1736-1740.	2.1	9
25	Ethenesulfonyl Fluoride: The Most Perfect Michael Acceptor Ever Found?. <i>Angewandte Chemie - International Edition</i> , 2016, 55, 12664-12667.	7.2	81
26	Ethensulfonylfluorid: der beste je entdeckte Michael-Akzeptor?. <i>Angewandte Chemie</i> , 2016, 128, 12854-12858.	1.6	19
27	Proteasome inhibition by new dual warhead containing peptido vinyl sulfonyl fluorides. <i>Bioorganic and Medicinal Chemistry</i> , 2016, 24, 3429-3435.	1.4	39
28	Cellular thermal shift and clickable chemical probe assays for the determination of drug-target engagement in live cells. <i>Organic and Biomolecular Chemistry</i> , 2016, 14, 6179-6183.	1.5	22
29	Arylfluorosulfates Inactivate Intracellular Lipid Binding Protein(s) through Chemoselective SuFEx Reaction with a Binding Site Tyr Residue. <i>Journal of the American Chemical Society</i> , 2016, 138, 7353-7364.	6.6	212
30	Palladium- and Nickel-Catalyzed Amination of Aryl Fluorosulfonates. <i>ACS Catalysis</i> , 2016, 6, 3515-3519.	5.5	77
31	Patent highlights: December 2015-January 2016. <i>Pharmaceutical Patent Analyst</i> , 2016, 5, 147-153.	0.4	0
32	A Heck-Matsuda Process for the Synthesis of Arylethenesulfonyl Fluorides: Selectively Addressable Bis-electrophiles for SuFEx Click Chemistry. <i>Angewandte Chemie</i> , 2016, 128, 14361-14364.	1.6	46
33	A Heck-Matsuda Process for the Synthesis of Arylethenesulfonyl Fluorides: Selectively Addressable Bis-electrophiles for SuFEx Click Chemistry. <i>Angewandte Chemie - International Edition</i> , 2016, 55, 14155-14158.	7.2	151
34	Chemoselective Preparation of Clickable Aryl Sulfonyl Fluoride Monomers: A Toolbox of Highly Functionalized Intermediates for Chemical Biology Probe Synthesis. <i>ChemBioChem</i> , 2016, 17, 1925-1930.	1.3	47
35	SuFEx Click: New Materials from SO <sub>x</sub> F and Silyl Ethers. <i>Chemistry - A European Journal</i> , 2016, 22, 16348-16354.	1.7	50
36	A Click Ligation Based on SuFEx for the Metal-Free Synthesis of Sugar and Iminosugar Clusters. <i>European Journal of Organic Chemistry</i> , 2016, 2016, 5102-5116.	1.2	35
37	Discovery of novel anti-HIV agents via Cu(I)-catalyzed azide-alkyne cycloaddition (CuAAC) click chemistry-based approach. <i>Expert Opinion on Drug Discovery</i> , 2016, 11, 857-871.	2.5	39
38	Reaction of Alkyl Halides with Rongalite: One-Pot and Telescoped Syntheses of Aliphatic Sulfonamides, Sulfonyl Fluorides, and Unsymmetrical Sulfones. <i>Organic Letters</i> , 2016, 18, 5848-5851.	2.4	84
39	Ethenesulfonyl Fluoride (ESF): An On-Water Procedure for the Kilogram-Scale Preparation. <i>Journal of Organic Chemistry</i> , 2016, 81, 11360-11362.	1.7	82
40	Synthesis of Sulfotyrosine-Containing Peptides by Incorporating Fluorosulfated Tyrosine Using an Fmoc-Based Solid-Phase Strategy. <i>Angewandte Chemie</i> , 2016, 128, 1867-1870.	1.6	17
41	Synthesis of Sulfotyrosine-Containing Peptides by Incorporating Fluorosulfated Tyrosine Using an Fmoc-Based Solid-Phase Strategy. <i>Angewandte Chemie - International Edition</i> , 2016, 55, 1835-1838.	7.2	43

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42	Selective and Orthogonal Post-Polymerization Modification using Sulfur(VI) Fluoride Exchange (SuFEx) and Copper-Catalyzed Azide-Alkyne Cycloaddition (CuAAC) Reactions. <i>Macromolecules</i> , 2016, 49, 4473-4479.	2.2	92
43	Multifunctional Surface Manipulation Using Orthogonal Click Chemistry. <i>Langmuir</i> , 2016, 32, 6600-6605.	1.6	45
44	Electrophilic, Activation-Free Fluorogenic Reagent for Labeling Bioactive Amines. <i>Bioconjugate Chemistry</i> , 2016, 27, 1430-1434.	1.8	22
45	Chemoselective Synthesis of Polysubstituted Pyridines from Heteroaryl Fluorosulfates. <i>Chemistry - A European Journal</i> , 2016, 22, 5692-5697.	1.7	72
46	Proximity-enabled bioreactivity to generate covalent peptide inhibitors of p53-Mdm4. <i>Chemical Communications</i> , 2016, 52, 5140-5143.	2.2	62
47	On-Water Synthesis of Biaryl Sulfonyl Fluorides. <i>Journal of Organic Chemistry</i> , 2016, 81, 2618-2623.	1.7	49
48	Rational design of a water-soluble, lipid-compatible fluorescent probe for Cu with sub-part-per-trillion sensitivity. <i>Chemical Science</i> , 2016, 7, 1468-1473.	3.7	27
49	Discovery of bioactive molecules from CuAAC click-chemistry-based combinatorial libraries. <i>Drug Discovery Today</i> , 2016, 21, 118-132.	3.2	138
50	Nucleophilic Deoxyfluorination of Phenols via Aryl Fluorosulfonate Intermediates. <i>Journal of the American Chemical Society</i> , 2017, 139, 1452-1455.	6.6	174
51	SuFEx: a metal-free click ligation for multivalent biomolecules. <i>Organic and Biomolecular Chemistry</i> , 2017, 15, 1549-1553.	1.5	40
52	A Synthesis of $\alpha$ -Dual Warhead- $\beta$ -Aryl Ethenesulfonyl Fluorides and One-Pot Reaction to $\beta$ -Sultams. <i>Organic Letters</i> , 2017, 19, 480-483.	2.4	91
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54	Click Reaction Based on the Biosynthesis of Firefly Luciferin. <i>Chemistry Letters</i> , 2017, 46, 753-755.	0.7	7
55	Copper-catalyzed sulfonylative Suzuki-Miyaura cross-coupling. <i>Chemical Science</i> , 2017, 8, 3249-3253.	3.7	127
56	Multidimensional SuFEx Click Chemistry: Sequential Sulfur(VI) Fluoride Exchange Connections of Diverse Modules Launched From An SOF <sub>4</sub> Hub. <i>Angewandte Chemie - International Edition</i> , 2017, 56, 2903-2908.	7.2	136
57	A facile and mild Pd-catalyzed one-pot process for direct hydrodeoxygenation (HDO) phenols to arenes through a ArOSO <sub>2</sub> F intermediates transformation. <i>Tetrahedron Letters</i> , 2017, 58, 2340-2343.	0.7	42
58	Bifluoride-catalysed sulfur(VI) fluoride exchange reaction for the synthesis of polysulfates and polysulfonates. <i>Nature Chemistry</i> , 2017, 9, 1083-1088.	6.6	222
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61	Sulfurâ€‘Fluoride Exchange (SuFEx)â€‘Mediated Synthesis of Sterically Hindered and Electronâ€‘Deficient Secondary and Tertiary Amides via Acyl Fluoride Intermediates. <i>Chemistry - A European Journal</i> , 2017, 23, 9990-9995.	1.7	37
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63	Aryl Fluorosulfate Trapped Staudinger Reduction. <i>Organic Letters</i> , 2017, 19, 1582-1585.	2.4	21
64	Palladiumâ€‘Catalyzed Fluorosulfonylvinylation of Organic Iodides. <i>Angewandte Chemie - International Edition</i> , 2017, 56, 4849-4852.	7.2	95
65	Palladiumâ€‘Catalyzed Fluorosulfonylvinylation of Organic Iodides. <i>Angewandte Chemie</i> , 2017, 129, 4927-4930.	1.6	31
66	Pd(PPh <sub>3</sub> ) <sub>4</sub> â€‘Catalyzed Buchwaldâ€‘Hartwig Amination of Aryl Fluorosulfonates with Aryl Amines. <i>Asian Journal of Organic Chemistry</i> , 2017, 6, 1222-1225.	1.3	22
67	Broad-Spectrum Kinase Profiling in Live Cells with Lysine-Targeted Sulfonyl Fluoride Probes. <i>Journal of the American Chemical Society</i> , 2017, 139, 680-685.	6.6	256
68	Trifluoromethylfluorosulfonylation of Unactivated Alkenes Using Readily Available Ag(O <sub>2</sub> CCF <sub>2</sub> SO <sub>2</sub> F) and <i>N</i> -Fluorobenzenesulfonimide. <i>Angewandte Chemie - International Edition</i> , 2017, 56, 15432-15435.	7.2	63
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70	Trifluoromethylfluorosulfonylation of Unactivated Alkenes Using Readily Available Ag(O <sub>2</sub> CCF <sub>2</sub> SO <sub>2</sub> F) and <i>N</i> -Fluorobenzenesulfonimide. <i>Angewandte Chemie</i> , 2017, 129, 15634-15637.	1.6	19
71	Click-on fluorescent triazolyl coumarin peptidomimetics as inhibitors of human breast cancer cell line MCF-7. <i>New Journal of Chemistry</i> , 2017, 41, 13483-13489.	1.4	17
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75	Auf Lysin zielende, kovalente Inhibitoren. <i>Angewandte Chemie</i> , 2017, 129, 15398-15408.	1.6	22
76	<i>Ex Situ</i> Generation of Sulfonyl Fluoride for the Synthesis of Aryl Fluorosulfates. <i>Organic Letters</i> , 2017, 19, 5244-5247.	2.4	83
77	Fabrication of Photocontrolled Surfaces for Oil/Water Separation through Sulfur(VI) Fluoride Exchange. <i>Chemistry - A European Journal</i> , 2017, 23, 14712-14717.	1.7	25

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79	Synthesis of a Class of Fused Sulfone Heterocycles <i>via</i> DBU-Catalyzed Direct Annulative SuFEx Click of Ethenesulfonyl Fluorides and Pyrazolones or 1,3-Dicarbonyl Compounds. Advanced Synthesis and Catalysis, 2017, 359, 3254-3260.	2.1	36
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99	Direct Access to Aryl Bis(trifluoromethyl)carbinols from Aryl Bromides or Fluorosulfates: Palladium-Catalyzed Carbonylation. <i>Angewandte Chemie</i> , 2018, 130, 6974-6978.	1.6	9
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102	Direct Access to Aryl Bis(trifluoromethyl)carbinols from Aryl Bromides or Fluorosulfates: Palladium-Catalyzed Carbonylation. <i>Angewandte Chemie - International Edition</i> , 2018, 57, 6858-6862.	7.2	38
103	Nucleophilic Fluorination with Aqueous Bifluoride Solution: Effect of the Phase-Transfer Catalyst. <i>ACS Sustainable Chemistry and Engineering</i> , 2018, 6, 6693-6701.	3.2	29
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112	SuFEx Chemistry of Thionyl Tetrafluoride ( $SO_4$ ) with Organolithium Nucleophiles: Synthesis of Sulfonimidoyl Fluorides, Sulfoximines, Sulfonimidamides, and Sulfonimidates. <i>Angewandte Chemie</i> , 2018, 130, 1957-1961.	1.6	43
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114	A New Portal to SuFEx Click Chemistry: A Stable Fluorosulfuryl Imidazolium Salt Emerging as an $\text{SO}_2$ -Donor of Unprecedented Reactivity, Selectivity, and Scope. <i>Angewandte Chemie</i> , 2018, 130, 2635-2640.	1.6	100
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127	A Vinyl Sulfone-Based Fluorogenic Probe Capable of Selective Labeling of PHGDH in Live Mammalian Cells. <i>Angewandte Chemie</i> , 2018, 130, 588-592.	1.6	11
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