Scott D Taylor

List of Publications by Year
in descending order

65 papers	1,496 citations	331670 21 h-index	361022 35 g-index
66 all docs	66 docs citations	66 times ranked	1587 citing authors

#	Article	IF	Citations
1	The Chiral Target of Daptomycin Is the 2R,2â€²S Stereoisomer of Phosphatidylglycerol. Angewandte Chemie - International Edition, 2022, 61, e202114858.	13.8	11
2	Discovery of Highly Active Derivatives of Daptomycin by Assessing the Effect of Amino Acid Substitutions at Positions 8 and 11 on a Daptomycin Analogue. ACS Infectious Diseases, 2022, 8, 778-789.	3.8	4
3	Synthesis of Azido Acids and Their Application in the Preparation of Complex Peptides. Synthesis, 2021, 53, 391-417.	2.3	1
4	A high-yielding solid-phase total synthesis of daptomycin using a Fmoc SPPS stable kynurenine synthon. Organic and Biomolecular Chemistry, 2021, 19, 3144-3153.	2.8	13
5	Nonthermoresponsive and Thermoresponsive Cationic Starch for the Flocculation of Oil Sands Mature Fine Tailings. Energy & Fuels, 2021, 35, 5163-5171.	5.1	1
6	Solid-Phase Total Synthesis of Dehydrotryptophan-Bearing Cyclic Peptides Tunicyclin B, Sclerotide A, CDA3a, and CDA4a using a Protected Î²-Hydroxytryptophan Building Block. Organic Letters, 2021, 23, 3048-3052.	4.6	6
7	Synthesis of Î²-Hydroxy-Î±,Î±-difluorosulfonamides from Carbanions of Difluoromethanesulfonamides. Journal of Organic Chemistry, 2021, 86, 6577-6591.	3.2	0
8	Enantioselective Synthesis and Application of Small and Environmentally Sensitive Fluorescent Amino Acids for Probing Biological Interactions. Journal of Organic Chemistry, 2021, 86, 11407-11418.	3.2	1
9	Total Synthesis of Analogs of A54145D and A54145A1 for Structureâ€“Activity Relationship Studies. Journal of Organic Chemistry, 2020, 85, 2213-2219.	3.2	7
10	Mild, Rapid, and Chemoselective Procedure for the Introduction of the 9-Phenyl-9-fluorenyl Protecting Group into Amines, Acids, Alcohols, Sulfonamides, Amides, and Thiols. Journal of Organic Chemistry, 2020, 85, 2068-2081.	3.2	3
11	Discovery of 5-aryl-3-thiophen-2-yl-1H-pyrazoles as a new class of Hsp90 inhibitors in hepatocellular carcinoma. Bioorganic Chemistry, 2020, 94, 103433.	4.1	8
12	Thermoresponsive Starch for the Flocculation of Oil Sands Mature Fine Tailings. Environmental Science & Technology, 2020, 54, 13981-13991.	10.0	7
13	Highly efficient and enantioselective syntheses of (2S,3R)-3-alkyl- and alkenylglutamates from Fmoc-protected Garnerâ€™s aldehyde. Amino Acids, 2020, 52, 987-998.	2.7	8
14	Synthesis of Fmoc-Protected Amino Alcohols via the Sharpless Asymmetric Aminohydroxylation Reaction Using FmocNHCl as the Nitrogen Source. Journal of Organic Chemistry, 2019, 84, 15476-15485.	3.2	4
15	Total Synthesis of A54145 Factor D. Journal of Organic Chemistry, 2019, 84, 12021-12030.	3.2	13
16	Total Synthesis of Paenibacterin and Its Analogues. Journal of Organic Chemistry, 2019, 84, 5339-5347.	3.2	9
17	The effect of replacing the ester bond with an amide bond and of overall stereochemistry on the activity of daptomycin. Bioorganic and Medicinal Chemistry, 2019, 27, 240-246.	3.0	15
18	Thermoresponsive hydroxybutylated starch nanoparticles. Carbohydrate Polymers, 2019, 209, 145-151.	10.2	11

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19	An entirely fmoc solid phase approach to the synthesis of daptomycin analogs. Peptide Science, 2019, 111, e23094.	1.8	17
20	Asymmetric Synthesis of Fmoc-Protected Î²-Hydroxy and Î²-Methoxy Amino Acids via a Sharpless Aminohydroxylation Reaction Using FmocNHCl. Organic Letters, 2018, 20, 7717-7720.	4.6	11
21	Efficient One-Pot, Two-Component Modular Synthesis of 3,5-Disubstituted Pyrazoles. ACS Omega, 2018, 3, 15566-15574.	3.5	4
22	Mechanistic studies on the effect of membrane lipid acyl chain composition on daptomycin pore formation. Chemistry and Physics of Lipids, 2018, 216, 73-79.	3.2	27
23	Synthesis of Nucleosideâ€5â€²â€Oâ€Tetraphosphates from Activated Trimetaphosphate and Nucleosideâ€5â€²â€Oâ€Monophosphates. Current Protocols in Nucleic Acid Chemistry, 2018, 75, e62.	0.5	2
24	An Acyl-Linked Dimer of Daptomycin Is Strongly Inhibited by the Bacterial Cell Wall. ACS Infectious Diseases, 2017, 3, 462-466.	3.8	5
25	Synthesis of Î²-Ketosulfonamides Derived from Amino Acids and Their Conversion to Î²-Keto-Î±,Î±-difluorosulfonamides via Electrophilic Fluorination. Journal of Organic Chemistry, 2017, 82, 11157-11165.	3.2	5
26	Daptomycin Pore Formation Is Restricted by Lipid Acyl Chain Composition. ACS Infectious Diseases, 2017, 3, 797-801.	3.8	19
27	Increased Electromer Formation and Charge Trapping in Solution-Processed versus Vacuum-Deposited Small Molecule Host Materials of Organic Light-Emitting Devices. ACS Applied Materials & Interfaces, 2017, 9, 40564-40572.	8.0	34
28	Two successive calcium-dependent transitions mediate membrane binding and oligomerization of daptomycin and the related antibiotic A54145. Biochimica Et Biophysica Acta - Biomembranes, 2016, 1858, 1999-2005.	2.6	27
29	A Fresh Look at the Staudinger Reaction on Azido Esters: Formation of 2H-1,2,3-Triazol-4-ols from Î±-Azido Esters Using Trialkyl Phosphines. Organic Letters, 2016, 18, 4412-4415.	4.6	16
30	Membrane Binding and Oligomerization of the Lipopeptide A54145 Studied by Pyrene Fluorescence. Biophysical Journal, 2016, 111, 1267-1277.	0.5	20
31	Exploring the Potent Inhibition of CTP Synthase by Gemcitabineâ€5â€²â€Triphosphate. ChemBioChem, 2016, 17, 2240-2249.	2.6	19
32	Î±-Azido Esters in Depsipeptide Synthesis: Câ€“O Bond Cleavage during Azido Group Reduction. Journal of Organic Chemistry, 2016, 81, 11831-11840.	3.2	9
33	One flask synthesis of 2â€²,3â€²-cyclic nucleoside monophosphates from unprotected nucleosides using activated cyclic trimetaphosphate. Tetrahedron Letters, 2016, 57, 5457-5459.	1.4	6
34	The action mechanism of daptomycin. Bioorganic and Medicinal Chemistry, 2016, 24, 6253-6268.	3.0	203
35	Synthesis of Nucleoside Triphosphates from 2â€²-3â€²-Protected Nucleosides Using Trimetaphosphate. Organic Letters, 2016, 18, 580-583.	4.6	22
36	Î±-Azido Acids in Solid-Phase Peptide Synthesis: Compatibility with Fmoc Chemistry and an Alternative Approach to the Solid Phase Synthesis of Daptomycin Analogs. Journal of Organic Chemistry, 2016, 81, 2624-2628.	3.2	14

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37	Solid-Phase Total Synthesis of Daptomycin and Analogs. Organic Letters, 2015, 17, 748-751.	4.6	55
38	A-ring substituted 17Î²-arylsulfonamides of 17Î²-aminoestra-1,3,5(10)-trien-3-ol as highly potent reversible inhibitors of steroid sulfatase. Bioorganic and Medicinal Chemistry, 2015, 23, 5681-5692.	3.0	8
39	Solid-phase synthesis and in vitro biological activity of a Thr4â†'Ser4 analog of daptomycin. Bioorganic and Medicinal Chemistry Letters, 2015, 25, 5490-5494.	2.2	17
40	Cardiolipin Prevents Membrane Translocation and Permeabilization by Daptomycin. Journal of Biological Chemistry, 2014, 289, 11584-11591.	3.4	136
41	Steroid derivatives as inhibitors of steroid sulfatase. Journal of Steroid Biochemistry and Molecular Biology, 2013, 137, 183-198.	2.5	28
42	Synthesis of Nucleoside Tetraphosphates and Dinucleoside Pentaphosphates via Activation of Cyclic Trimetaphosphate. Organic Letters, 2013, 15, 2612-2615.	4.6	33
43	Mutual inhibition through hybrid oligomer formation of daptomycin and the semisynthetic lipopeptide antibiotic CB-182,462. Biochimica Et Biophysica Acta - Biomembranes, 2013, 1828, 302-308.	2.6	24
44	Characterization of daptomycin oligomerization with perylene excimer fluorescence: Stoichiometric binding of phosphatidylglycerol triggers oligomer formation. Biochimica Et Biophysica Acta - Biomembranes, 2012, 1818, 673-678.	2.6	66
45	17Î²-Arylsulfonamides of 17Î²-aminoestra-1,3,5(10)-trien-3-ol as highly potent inhibitors of steroid sulfatase. Bioorganic and Medicinal Chemistry, 2012, 20, 1535-1544.	3.0	14
46	Efficient syntheses of 17-Î²-amino steroids. Steroids, 2011, 76, 1098-1102.	1.8	13
47	Inhibition of steroid sulfatase with 4-substituted estrone and estradiol derivatives. Bioorganic and Medicinal Chemistry, 2011, 19, 5999-6005.	3.0	25
48	Multiple Pathways for the Irreversible Inhibition of Steroid Sulfatase with Quinone Methideâ€Generating Suicide Inhibitors. ChemBioChem, 2009, 10, 1457-1461.	2.6	37
49	Ground State, Intermediate, and Multivalent Nucleotide Analogue Inhibitors of Cytidine 5â€²â€Triphosphate Synthase. ChemMedChem, 2008, 3, 1853-1857.	3.2	7
50	Bismethylene Triphosphate Nucleotides of Uridine 4-Phosphate Analogues:â€‰ A New Class of Anionic Pyrimidine Nucleotide Analogues. Journal of Organic Chemistry, 2008, 73, 1403-1412.	3.2	25
51	Synthesis of 4-Formyl Estrone Using a Positional Protecting Group and Its Conversion to Other C-4-Substituted Estrogens. Journal of Organic Chemistry, 2007, 72, 8824-8830.	3.2	18
52	Synthesis of Î±-Fluorosulfonate and Î±-Fluorosulfonamide Analogues of a Sulfated Carbohydrate. Organic Letters, 2006, 8, 5617-5620.	4.6	16
53	An Unsymmetrical Approach to the Synthesis of Bismethylene Triphosphate Analogues. Organic Letters, 2006, 8, 4243-4246.	4.6	21
54	Boronic acids as inhibitors of steroid sulfatase. Bioorganic and Medicinal Chemistry, 2006, 14, 8564-8573.	3.0	42

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55	Synthesis of Methylene- and Difluoromethylenephosphonate Analogues of Uridine-4-phosphate and 3-Deazauridine-4-phosphate. Journal of Organic Chemistry, 2006, 71, 9420-9430.	3.2	18
56	A fluorogenic substrate for the continuous assaying of aryl sulfatases. Analytical Biochemistry, 2005, 340, 80-88.	2.4	22
57	Synthesis of a non-hydrolyzable estrone sulfate analogue bearing the difluoromethanesulfonamide group and its evaluation as a steroid sulfatase inhibitor. Organic and Biomolecular Chemistry, 2005, 3, 3329.	2.8	22
58	Recent advances in protein tyrosine phosphatase 1B inhibitors. Expert Opinion on Investigational Drugs, 2004, 13, 199-214.	4.1	80
59	Synthesis of Î±-Fluorosulfonamides by Electrophilic Fluorination. Organic Letters, 2004, 6, 4285-4288.	4.6	50
60	The difluoromethylene group as a replacement for the labile oxygen in steroid sulfates: a new approach to steroid sulfatase inhibitors. Bioorganic and Medicinal Chemistry Letters, 2004, 14, 151-155.	2.2	29
61	Antibody-catalyzed activation of a model tripartate prodrug by a tandem hydrolysisâ€“1,6-elimination reaction. Chemical Communications, 2001, , 1386-1387.	4.1	7
62	Synthesis of Protectedl-4-[Sulfono(difluoromethyl)]phenylalanine and Its Incorporation into a Peptide. Organic Letters, 2001, 3, 1571-1574.	4.6	28
63	Preparation of chiral Î±-monofluoroalkylphosphonic acids and their evaluation as inhibitors of protein tyrosine phosphatase 1B. Journal of the Chemical Society, Perkin Transactions 1, 2000, , 1271-1281.	1.3	22
64	On the origins of enhanced reactivity of five-membered cyclic phosphate esters. The relative contributions of enthalpic and entropic factors. Journal of the American Chemical Society, 1990, 112, 6669-6671.	13.7	51
65	The Chiral Target of Daptomycin is the 2R,2'S Stereoisomer of Phosphatidylglycerol. Angewandte Chemie, 0, , .	2.0	0