Allyn T Londregan

List of Publications by Year
in descending order

21	807	623734 14	713466 21
papers	citations	h-index	g-index

24	24	24	1026
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	General and Mild Preparation of 2-Aminopyridines. Organic Letters, 2010, 12, 5254-5257.	4.6	126
2	Mild Addition of Nucleophiles to Pyridine-N-Oxides. Organic Letters, 2011, 13, 1840-1843.	4.6	110
3	Selective stalling of human translation through small-molecule engagement of the ribosome nascent chain. PLoS Biology, 2017, 15, e2001882.	5.6	104
4	Discovery of PF-5190457, a Potent, Selective, and Orally Bioavailable Ghrelin Receptor Inverse Agonist Clinical Candidate. ACS Medicinal Chemistry Letters, 2014, 5, 474-479.	2.8	51
5	A New and Useful Method for the Macrocyclization of Linear Peptides. Organic Letters, 2012, 14, 2890-2893.	4.6	47
6	Identification of Tetrahydropyrido[4,3-d]pyrimidine Amides as a New Class of Orally Bioavailable TGR5 Agonists. ACS Medicinal Chemistry Letters, 2013, 4, 63-68.	2.8	45
7	Expedient Synthesis of Î±-(2-Azaheteroaryl) Acetates via the Addition of Silyl Ketene Acetals to Azine-N-oxides. Organic Letters, 2014, 16, 3336-3339.	4.6	44
8	Preparation of Heteroaryl Ethers from Azine N-Oxides and Alcohols. Organic Letters, 2016, 18, 1362-1365.	4.6	44
9	Small Molecule Proprotein Convertase Subtilisin/Kexin Type 9 (PCSK9) Inhibitors: Hit to Lead Optimization of Systemic Agents. Journal of Medicinal Chemistry, 2018, 61, 5704-5718.	6.4	37
10	Liverâ€Targeted Smallâ€Molecule Inhibitors of Proprotein Convertase Subtilisin/Kexin Type 9 Synthesis. Angewandte Chemie - International Edition, 2017, 56, 16218-16222.	13.8	35
11	Discovery of an in Vivo Tool to Establish Proof-of-Concept for MAP4K4-Based Antidiabetic Treatment. ACS Medicinal Chemistry Letters, 2015, 6, 1128-1133.	2.8	33
12	Rapid and Selective in situ Reduction of Pyridine-N-oxides with TetraÂhydroxydiboron. Synlett, 2013, 24, 2695-2700.	1.8	30
13	Discovery of 5-phenoxy-1,3-dimethyl-1H-pyrazole-4-carboxamides as potent agonists of TGR5 via sequential combinatorial libraries. Bioorganic and Medicinal Chemistry Letters, 2013, 23, 1407-1411.	2.2	22
14	Expedient Synthesis of Î±-Heteroaryl Piperidines Using a Pd-Catalyzed Suzuki Cross-Couplingâ€“Reduction Sequence. Organic Letters, 2014, 16, 413-415.	4.6	21
15	An improved amide coupling procedure for the synthesis of N-(pyridin-2-yl)amides. Tetrahedron Letters, 2009, 50, 1986-1988.	1.4	14
16	Discovery of Selective Small Molecule Inhibitors of Monoacylglycerol Acyltransferase 3. Journal of Medicinal Chemistry, 2015, 58, 7164-7172.	6.4	14
17	Sequential Xanthalation and O-Trifluoromethylation of Phenols: AÂProcedure for the Synthesis of Aryl Trifluoromethyl Ethers. Journal of Organic Chemistry, 2019, 84, 15767-15776.	3.2	12
18	Discovery of N-(piperidin-3-yl)-N-(pyridin-2-yl)piperidine/piperazine-1-carboxamides as small molecule inhibitors of PCSK9. Bioorganic and Medicinal Chemistry Letters, 2018, 28, 3685-3688.	2.2	11

#	Article	IF	Citations
19	The enantioselective synthesis of (R)- and (S)-3-amino-3,4-dihydro-1H-[1,8]naphthyridin-2-one. Tetrahedron: Asymmetry, 2010, 21, 2072-2075.	1.8	4
20	Synthesis and Analysis of Macrocyclic Peptides with 310-Helical Structure. Synlett, 2015, 26, 1164-1168.	1.8	2
21	Liverâ€Targeted Smallâ€Molecule Inhibitors of Proprotein Convertase Subtilisin/Kexin Type 9 Synthesis. Angewandte Chemie, 2017, 129, 16436-16440.	2.0	1