Marc A Labroli

List of Publications by Year
in descending order

14	494	1170033 9	1181555 14
papers	citations	h-index	g-index

14	14	14	931
all docs	docs citations	times ranked	citing authors

#	Article	IF	Citations
1	Informing the Selection of Screening Hit Series with in Silico Absorption, Distribution, Metabolism, Excretion, and Toxicity Profiles. Journal of Medicinal Chemistry, 2017, 60, 6771-6780.	2.9	17
2	Can We Make Small Molecules Lean? Optimization of a Highly Lipophilic TarO Inhibitor. Journal of Medicinal Chemistry, 2017, 60, 3851-3865.	2.9	8
3	Chemical Genetic Analysis and Functional Characterization of Staphylococcal Wall Teichoic Acid 2-Epimerases Reveals Unconventional Antibiotic Drug Targets. PLoS Pathogens, 2016, 12, e1005585.	2.1	35
4	A convergent preparation of the CHK1 inhibitor MK-8776 (SCH 900776). Tetrahedron Letters, 2016, 57, 2601-2603.	0.7	4
5	Benzimidazole analogs as WTA biosynthesis inhibitors targeting methicillin resistant Staphylococcus aureus. Bioorganic and Medicinal Chemistry Letters, 2016, 26, 4743-4747.	1.0	10
6	TarO-specific inhibitors of wall teichoic acid biosynthesis restore Î²-lactam efficacy against methicillin-resistant staphylococci. Science Translational Medicine, 2016, 8, 329ra32.	5.8	95
7	NMR Binding and Functional Assays for Detecting Inhibitors of S. aureus MnaA. Journal of Biomolecular Screening, 2016, 21, 579-589.	2.6	5
8	Discovery of potent wall teichoic acid early stage inhibitors. Bioorganic and Medicinal Chemistry Letters, 2016, 26, 3999-4002.	1.0	12
9	Syntheses of 5â€²-amino-2â€²,5â€²-dideoxy-2â€²,2â€²-difluorocytidine derivatives as novel anticancer nucleoside analogs. Tetrahedron Letters, 2014, 55, 598-602.	0.7	6
10	The identification of novel 5â€²-amino gemcitabine analogs as potent RRM1 inhibitors. Bioorganic and Medicinal Chemistry, 2014, 22, 2303-2310.	1.4	8
11	Discovery of pyrazolo[1,5-a]pyrimidine-based Pim inhibitors: A template-based approach. Bioorganic and Medicinal Chemistry Letters, 2013, 23, 6178-6182.	1.0	27
12	Discovery of pyrazolo[1,5-a]pyrimidine-based CHK1 inhibitors: A template-based approachâ€”Part 1. Bioorganic and Medicinal Chemistry Letters, 2011, 21, 467-470.	1.0	58
13	Discovery of pyrazolo[1,5-a]pyrimidine-based CHK1 inhibitors: A template-based approachâ€”Part 2. Bioorganic and Medicinal Chemistry Letters, 2011, 21, 471-474.	1.0	54
14	Targeting the Replication Checkpoint Using SCH 900776, a Potent and Functionally Selective CHK1 Inhibitor Identified via High Content Screening. Molecular Cancer Therapeutics, 2011, 10, 591-602.	1.9	155