

F Javier Piedrafita

List of Publications by Year in descending order

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16
papers

382
citations

949033

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docs citations

17
times ranked

926
citing authors

#	ARTICLE	IF	CITATIONS
1	Targeted NUDT5 inhibitors block hormone signaling in breast cancer cells. <i>Nature Communications</i> , 2018, 9, 250.	5.8	56
2	Identification of Triazolothiadiazoles as Potent Inhibitors of the dCTP Pyrophosphatase 1. <i>Journal of Medicinal Chemistry</i> , 2017, 60, 2148-2154.	2.9	14
3	Piperazin-1-ylpyridazine Derivatives Are a Novel Class of Human dCTP Pyrophosphatase 1 Inhibitors. <i>Journal of Medicinal Chemistry</i> , 2017, 60, 4279-4292.	2.9	19
4	Crystal Structure of the Emerging Cancer Target MTHFD2 in Complex with a Substrate-Based Inhibitor. <i>Cancer Research</i> , 2017, 77, 937-948.	0.4	67
5	Diverse heterocyclic scaffolds as dCTP pyrophosphatase 1 inhibitors. Part 2: Pyridone- and pyrimidinone-derived systems. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2017, 27, 3219-3225.	1.0	4
6	Diverse heterocyclic scaffolds as dCTP pyrophosphatase 1 inhibitors. Part 1: Triazoles, triazolopyrimidines, triazinoindoles, quinoline hydrazones and arylpiperazines. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2017, 27, 3897-3904.	1.0	2
7	Novel spirocyclic systems via multicomponent aza-Diels-Alder reaction. <i>Organic and Biomolecular Chemistry</i> , 2017, 15, 7758-7764.	1.5	4
8	Tetrahydrobenzothiophene carboxamides: Beyond the kinase domain and into the fatty acid realm. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2017, 27, 4462-4466.	1.0	1
9	Structure-metabolism-relationships in the microsomal clearance of piperazin-1-ylpyridazines. <i>MedChemComm</i> , 2017, 8, 1553-1560.	3.5	1
10	Discovery of the First Potent and Selective Inhibitors of Human dCTP Pyrophosphatase 1. <i>Journal of Medicinal Chemistry</i> , 2016, 59, 1140-1148.	2.9	40
11	Lysophosphatidic acid receptor (LPA) modulators: The current pharmacological toolbox. <i>Progress in Lipid Research</i> , 2015, 58, 51-75.	5.3	57
12	Vinylidene Boronates: New Building Blocks for the Synthesis of Aza-Heterocycles. <i>Chemistry - A European Journal</i> , 2015, 21, 7394-7398.	1.7	23
13	Stereoselective synthesis of carbocyclic analogues of the nucleoside Q precursor (PreQ ₀). <i>Beilstein Journal of Organic Chemistry</i> , 2014, 10, 1333-1338.	1.3	13
14	Chemical strategies for development of ATR inhibitors. <i>Expert Reviews in Molecular Medicine</i> , 2014, 16, e10.	1.6	19
15	A Convenient Microwave-Assisted Propylphosphonic Anhydride (T3P [®]) Mediated One-Pot Pyrazolone Synthesis. <i>European Journal of Organic Chemistry</i> , 2013, 2013, 5879-5885.	1.2	16
16	Small-molecule inhibitors of Î² kinase (IKK) and IKK-related kinases. <i>Pharmaceutical Patent Analyst</i> , 2013, 2, 481-498.	0.4	43