

# Sabin Llona-Minguez

## List of Publications by Year in descending order

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16  
papers

332  
citations

840776

11  
h-index

888059

17  
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18  
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18  
docs citations

18  
times ranked

712  
citing authors

#	ARTICLE	IF	CITATIONS
1	Lysophosphatidic acid receptor (LPA) modulators: The current pharmacological toolbox. <i>Progress in Lipid Research</i> , 2015, 58, 51-75.	11.6	57
2	Targeted NUDT5 inhibitors block hormone signaling in breast cancer cells. <i>Nature Communications</i> , 2018, 9, 250.	12.8	56
3	Small-molecule inhibitors of I $\kappa$ B kinase (IKK) and IKK-related kinases. <i>Pharmaceutical Patent Analyst</i> , 2013, 2, 481-498.	1.1	43
4	Discovery of the First Potent and Selective Inhibitors of Human dCTP Pyrophosphatase 1. <i>Journal of Medicinal Chemistry</i> , 2016, 59, 1140-1148.	6.4	40
5	Vinylid MIDA Boronates: New Building Blocks for the Synthesis of Aza-Heterocycles. <i>Chemistry - A European Journal</i> , 2015, 21, 7394-7398.	3.3	23
6	Inhibitory Kappa B Kinase I $\kappa$ B (IKK) Inhibitors That Recapitulate Their Selectivity in Cells against Isoform-Related Biomarkers. <i>Journal of Medicinal Chemistry</i> , 2017, 60, 7043-7066.	6.4	23
7	Piperazin-1-ylpyridazine Derivatives Are a Novel Class of Human dCTP Pyrophosphatase 1 Inhibitors. <i>Journal of Medicinal Chemistry</i> , 2017, 60, 4279-4292.	6.4	19
8	A Convenient Microwave-Assisted Propylphosphonic Anhydride (T3P) Mediated One-Pot Pyrazolone Synthesis. <i>European Journal of Organic Chemistry</i> , 2013, 2013, 5879-5885.	2.4	16
9	Identification of Triazolothiadiazoles as Potent Inhibitors of the dCTP Pyrophosphatase 1. <i>Journal of Medicinal Chemistry</i> , 2017, 60, 2148-2154.	6.4	14
10	A facile and efficient synthesis of tetrahydro- $\beta$ -carbolines. <i>Tetrahedron Letters</i> , 2013, 54, 3554-3557.	1.4	13
11	Stereoselective synthesis of carbocyclic analogues of the nucleoside Q precursor (PreQ <sub>0</sub> ). <i>Beilstein Journal of Organic Chemistry</i> , 2014, 10, 1333-1338.	2.2	13
12	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.	2.2	4
13	Novel spirocyclic systems via multicomponent aza-Diels-Alder reaction. <i>Organic and Biomolecular Chemistry</i> , 2017, 15, 7758-7764.	2.8	4
14	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.	2.2	2
15	Tetrahydrobenzothiophene carboxamides: Beyond the kinase domain and into the fatty acid realm. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2017, 27, 4462-4466.	2.2	1
16	Structure-metabolism-relationships in the microsomal clearance of piperazin-1-ylpyridazines. <i>MedChemComm</i> , 2017, 8, 1553-1560.	3.4	1