

Kevin J Duffy

List of Publications by Year in descending order

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33
papers

3,157
citations

304602

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36
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docs citations

36
times ranked

4630
citing authors

#	ARTICLE	IF	CITATIONS
1	STING Activation and its Application in Immuno-Oncology. <i>Current Topics in Medicinal Chemistry</i> , 2019, 19, 2205-2227.	1.0	11
2	Design of amidobenzimidazole STING receptor agonists with systemic activity. <i>Nature</i> , 2018, 564, 439-443.	13.7	505
3	Discovery and Preclinical Characterization of GSK1278863 (Daprodustat), a Small Molecule Hypoxia Inducible Factor Prolyl Hydroxylase Inhibitor for Anemia. <i>Journal of Pharmacology and Experimental Therapeutics</i> , 2017, 363, 336-347.	1.3	65
4	Endothelial HIF-2 mediates protection and recovery from ischemic kidney injury. <i>Journal of Clinical Investigation</i> , 2014, 124, 2396-2409.	3.9	150
5	Targeting Lactate Dehydrogenase-A Inhibits Tumorigenesis and Tumor Progression in Mouse Models of Lung Cancer and Impacts Tumor-Initiating Cells. <i>Cell Metabolism</i> , 2014, 19, 795-809.	7.2	411
6	Quinoline 3-sulfonamides inhibit lactate dehydrogenase A and reverse aerobic glycolysis in cancer cells. <i>Cancer & Metabolism</i> , 2013, 1, 19.	2.4	163
7	Preischemic targeting of HIF prolyl hydroxylation inhibits fibrosis associated with acute kidney injury. <i>American Journal of Physiology - Renal Physiology</i> , 2012, 302, F1172-F1179.	1.3	104
8	Chronic Inhibition of Hypoxia-inducible Factor Prolyl 4-hydroxylase Improves Ventricular Performance, Remodeling, and Vascularity After Myocardial Infarction in the Rat. <i>Journal of Cardiovascular Pharmacology</i> , 2010, 56, 147-155.	0.8	84
9	Discovery of the First Potent and Selective Inhibitor of Centromere-Associated Protein E: GSK923295. <i>ACS Medicinal Chemistry Letters</i> , 2010, 1, 30-34.	1.3	54
10	Substituted benzothiadiazine inhibitors of Hepatitis C virus polymerase. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2009, 19, 4350-4353.	1.0	43
11	Synthesis and biological activity of heteroaryl 3-(1,1-dioxo-2H-(1,2,4)-benzothiadiazin-3-yl)-4-hydroxy-2(1H)-quinolinone derivatives as hepatitis C virus NS5B polymerase inhibitors. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2009, 19, 4354-4358.	1.0	24
12	Preclinical Activity of Eltrombopag (SB-497115), an Oral, Nonpeptide Thrombopoietin Receptor Agonist. <i>Stem Cells</i> , 2009, 27, 424-430.	1.4	243
13	Biochemical Characterization of Human Prolyl Hydroxylase Domain Protein 2 Variants Associated with Erythrocytosis. <i>Biochemistry</i> , 2008, 47, 11165-11167.	1.2	23
14	Novel ATP-Competitive Kinesin Spindle Protein Inhibitors. <i>Journal of Medicinal Chemistry</i> , 2007, 50, 4939-4952.	2.9	73
15	THE DISCOVERY OF ELTROMBOPAG, AN ORALLY BIOAVAILABLE TpoR AGONIST. , 2007, , 241-254.		8
16	GSK626616: A DYRK3 Inhibitor as a Potential New Therapy for the Treatment of Anemia.. <i>Blood</i> , 2007, 110, 510-510.	0.6	5
17	3-(1,1-Dioxo-2H-(1,2,4)-benzothiadiazin-3-yl)-4-hydroxy-2(1H)-quinolinones, Potent Inhibitors of Hepatitis C Virus RNA-Dependent RNA Polymerase. <i>Journal of Medicinal Chemistry</i> , 2006, 49, 971-983.	2.9	157
18	Discovery and characterization of a selective, nonpeptidyl thrombopoietin receptor agonist. <i>Experimental Hematology</i> , 2005, 33, 85-93.	0.2	176

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19	A Highly Efficient, Asymmetric Synthesis of Benzothiadiazine-Substituted Tetramic Acids:â€‰ Potent Inhibitors of Hepatitis C Virus RNA-Dependent RNA Polymerase. <i>Organic Letters</i> , 2005, 7, 5521-5524.	2.4	36
20	Species Specificity and Receptor Domain Interaction of a Small Molecule TPO Receptor Agonist.. <i>Blood</i> , 2004, 104, 2909-2909.	0.6	20
21	Discovery of SB-497115, a Small-Molecule Thrombopoietin (TPO) Receptor Agonist for the Treatment of Thrombocytopenia.. <i>Blood</i> , 2004, 104, 2910-2910.	0.6	15
22	Biological Activity and Selectivity for Tpo Receptor of the Orally Bioavailable, Small Molecule Tpo Receptor Agonist, SB-497115.. <i>Blood</i> , 2004, 104, 2912-2912.	0.6	11
23	Resistance Profile of a Hepatitis C Virus RNA-Dependent RNA Polymerase Benzothiadiazine Inhibitor. <i>Antimicrobial Agents and Chemotherapy</i> , 2003, 47, 3525-3530.	1.4	95
24	Selective Binding and Oligomerization of the Murine Granulocyte Colony-stimulating Factor Receptor by a Low Molecular Weight, Nonpeptidyl Ligand. <i>Journal of Biological Chemistry</i> , 2003, 278, 9426-9434.	1.6	15
25	Arresting Initiation of Hepatitis C Virus RNA Synthesis Using Heterocyclic Derivatives. <i>Journal of Biological Chemistry</i> , 2003, 278, 16602-16607.	1.6	65
26	Identification and Biological Characterization of Heterocyclic Inhibitors of the Hepatitis C Virus RNA-dependent RNA Polymerase. <i>Journal of Biological Chemistry</i> , 2002, 277, 38322-38327.	1.6	190
27	Identification of a Pharmacophore for Thrombopoietic Activity of Small, Non-Peptidyl Molecules. 2. Rational Design of Naphtho[1,2-d]imidazole Thrombopoietin Mimics. <i>Journal of Medicinal Chemistry</i> , 2002, 45, 3576-3578.	2.9	53
28	Identification of a Pharmacophore for Thrombopoietic Activity of Small, Non-Peptidyl Molecules. 1. Discovery and Optimization of Salicylaldehyde Thiosemicarbazone Thrombopoietin Mimics. <i>Journal of Medicinal Chemistry</i> , 2002, 45, 3573-3575.	2.9	83
29	Hydrazinonaphthalene and Azonaphthalene Thrombopoietin Mimics Are Nonpeptidyl Promoters of Megakaryocytopoiesis. <i>Journal of Medicinal Chemistry</i> , 2001, 44, 3730-3745.	2.9	106
30	Design and synthesis of diaminopyrrolidinone inhibitors of human osteoclast cathepsin K. <i>Bioorganic and Medicinal Chemistry Letters</i> , 1999, 9, 1907-1910.	1.0	17
31	Total Synthesis of Altohyrin A (Spongistatin 1): Part 1. <i>Angewandte Chemie - International Edition</i> , 1998, 37, 187-190.	7.2	130
32	A novel synthesis of 2-(2-quinoxalino)-3,5-diarylfurans. <i>Tetrahedron Letters</i> , 1998, 39, 8017-8020.	0.7	7
33	The scope and mechanism of a novel synthesis of 3,4-fused isoxazoles. <i>Journal of the Chemical Society Chemical Communications</i> , 1995, , 2457.	2.0	8