Alaric J Dyckman

List of Publications by Year
in descending order

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24 papers	527 citations	686830 13 h-index	642321 23 g-index
26 all docs	26 docs citations	26 times ranked	717 citing authors

#	Article	IF	Citations
1	Design, Synthesis, and Anti-inflammatory Properties of Orally Active 4-(Phenylamino)-pyrrolo[2,1-f][1,2,4]triazine p38Î± Mitogen-Activated Protein Kinase Inhibitors. Journal of Medicinal Chemistry, 2008, 51, 4-16.	2.9	81
2	Discovery of 4-(5-(Cyclopropylcarbamoyl)-2-methylphenylamino)-5-methyl-N-propylpyrrolo[1,2-f][1,2,4]triazine-6-carboxamide (BMS-582949), a Clinical p38Î± MAP Kinase Inhibitor for the Treatment of Inflammatory Diseases. Journal of Medicinal Chemistry, 2010, 53, 6629-6639.	2.9	57
3	The Discovery of Orally Active Triaminotriazine Aniline Amides as Inhibitors of p38 MAP Kinase. Journal of Medicinal Chemistry, 2004, 47, 6283-6291.	2.9	56
4	Modulators of Sphingosine-1-phosphate Pathway Biology: Recent Advances of Sphingosine-1-phosphate Receptor 1 (S1P₁) Agonists and Future Perspectives. Journal of Medicinal Chemistry, 2017, 60, 5267-5289.	2.9	48
5	Novel Tricyclic Inhibitors of IÎºB Kinase. Journal of Medicinal Chemistry, 2009, 52, 1994-2005.	2.9	25
6	Identification and Preclinical Pharmacology of BMS-986104: A Differentiated S1P₁ Receptor Modulator in Clinical Trials. ACS Medicinal Chemistry Letters, 2016, 7, 283-288.	1.3	25
7	Regioselective Epoxide Ring Opening for the Stereospecific Scale-Up Synthesis of BMS-960, A Potent and Selective Isoxazole-Containing S1P₁ Receptor Agonist. Organic Process Research and Development, 2017, 21, 200-207.	1.3	25
8	Discovery of Potent and Orally Bioavailable Small Molecule Antagonists of Toll-like Receptors 7/8/9 (TLR7/8/9). ACS Medicinal Chemistry Letters, 2020, 11, 1751-1758.	1.3	24
9	Discovery and Structureâ€“Activity Relationship (SAR) of a Series of Ethanolamine-Based Direct-Acting Agonists of Sphingosine-1-phosphate (S1P₁). Journal of Medicinal Chemistry, 2016, 59, 6248-6264.	2.9	22
10	Potent and Selective Agonists of Sphingosine 1-Phosphate 1 (S1P₁): Discovery and SAR of a Novel Isoxazole Based Series. Journal of Medicinal Chemistry, 2016, 59, 2820-2840.	2.9	20
11	Synthesis, initial SAR and biological evaluation of 1,6-dihydroimidazo[4,5-d]pyrrolo[2,3-b]pyridin-4-amine derived inhibitors of IÎºB kinase. Bioorganic and Medicinal Chemistry Letters, 2009, 19, 2646-2649.	1.0	19
12	Development of a Practical Synthesis of a p38 Kinase Inhibitor via a Safe and Robust Amination. Organic Process Research and Development, 2012, 16, 1618-1625.	1.3	19
13	An Efficient Scale-Up Synthesis of BMS-520, a Potent and Selective Isoxazole-Containing S1P₁ Receptor Agonist. Organic Process Research and Development, 2016, 20, 989-995.	1.3	15
14	Discovery of pyrrolo[2,1-f][1,2,4]triazine C6-ketones as potent, orally active p38Î± MAP kinase inhibitors. Bioorganic and Medicinal Chemistry Letters, 2011, 21, 4633-4637.	1.0	13
15	Identification and Preclinical Pharmacology of ((1R,3S)-1-Amino-3-((S)-6-(2-methoxyphenethyl)-5,6,7,8-tetrahydronaphthalen-2-yl)cyclopentyl)methanol (BMS-986166): A Differentiated Sphingosine-1-phosphate Receptor 1 (S1P₁) Modulator Advanced into Clinical Trials. Journal of Medicinal Chemistry, 2019, 62, 2265-2285.	2.9	13
16	Imidazo[4,5-d]thiazolo[5,4-b]pyridine based inhibitors of IKK2: Synthesis, SAR, PK/PD and activity in a preclinical model of rheumatoid arthritis. Bioorganic and Medicinal Chemistry Letters, 2011, 21, 383-386.	1.0	11
17	Asymmetric Hydroboration Approach to the Scalable Synthesis of ((1R,3S)-1-Amino-3-((R)-6-hexyl-5,6,7,8-tetrahydronaphthalen-2-yl)cyclopentyl)methanol (BMS-986104) as a Potent S1P₁ Receptor Modulator. Journal of Medicinal Chemistry, 2016, 59, 11138-11147.	2.9	10
18	Novel tricyclic inhibitors of IKK2: Discovery and SAR leading to the identification of 2-methoxy-N-((6-(1-methyl-4-(methylamino)-1,6-dihydroimidazo[4,5-d]pyrrolo[2,3-b]pyridin-7-yl)pyridin-2-yl)methyl)acetamide (BMS-066). Bioorganic and Medicinal Chemistry Letters, 2011, 21, 7006-7012.	1.0	9

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19	Identification of Tricyclic Agonists of Sphingosine-1-phosphate Receptor 1 (S1P₁) Employing Ligand-Based Drug Design. Journal of Medicinal Chemistry, 2016, 59, 9837-9854.	2.9	8
20	Integrating High-Content Analysis into a Multiplexed Screening Approach to Identify and Characterize GPCR Agonists. Journal of Biomolecular Screening, 2014, 19, 1079-1089.	2.6	7
21	Identification of potent tricyclic prodrug S1P1 receptor modulators. MedChemComm, 2017, 8, 725-729.	3.5	6
22	Aryl Ether-Derived Sphingosine-1-Phosphate Receptor (S1P1) Modulators: Optimization of the PK, PD, and Safety Profiles. ACS Medicinal Chemistry Letters, 2020, 11, 1766-1772.	1.3	5
23	Identification of 2-Pyridinylindole-Based Dual Antagonists of Toll-like Receptors 7 and 8 (TLR7/8). ACS Medicinal Chemistry Letters, 2022, 13, 812-818.	1.3	5
24	Bicyclic Ligand-Biased Agonists of S1P₁: Exploring Side Chain Modifications to Modulate the PK, PD, and Safety Profiles. Journal of Medicinal Chemistry, 2021, 64, 1454-1480.	2.9	4