

Brad D Maxwell

List of Publications by Year in descending order

Source: <https://exaly.com/author-pdf/1680432/publications.pdf>

Version: 2024-02-01

20
papers

865
citations

1307594

7
h-index

888059

17
g-index

21
all docs

21
docs citations

21
times ranked

1209
citing authors

#	ARTICLE	IF	CITATIONS
1	A general alkyl-alkyl cross-coupling enabled by redox-active esters and alkylzinc reagents. <i>Science</i> , 2016, 352, 801-805.	12.6	579
2	Hydromethylation of Unactivated Olefins. <i>Journal of the American Chemical Society</i> , 2015, 137, 8046-8049.	13.7	137
3	Discovery of a Parenteral Small Molecule Coagulation Factor XIa Inhibitor Clinical Candidate (BMS-962212). <i>Journal of Medicinal Chemistry</i> , 2017, 60, 9703-9723.	6.4	45
4	Discovery of Pyrrolidine-Containing GPR40 Agonists: Stereochemistry Effects a Change in Binding Mode. <i>Journal of Medicinal Chemistry</i> , 2017, 60, 1417-1431.	6.4	25
5	Deuterated active pharmaceutical ingredients: A science-based proposal for synthesis, analysis, and control. Part 1: Framing the problem. <i>Journal of Labelled Compounds and Radiopharmaceuticals</i> , 2019, 62, 690-694.	1.0	16
6	Discovery of Clinical Candidate 2-((2 <i>S</i> ,6 <i>S</i>)-2-Phenyl-6-hydroxyadamantan-2-yl)-1-(3- ² -hydroxyazetid-1-yl)ethanone [BMS-816336], an Orally Active Novel Selective 11 ^β -Hydroxysteroid Dehydrogenase Type 1 Inhibitor. <i>Journal of Medicinal Chemistry</i> , 2017, 60, 4932-4948.	6.4	10
7	The syntheses and <i>in vitro</i> biotransformation studies of [¹⁴ C]apixaban, a highly potent, selective, efficacious and orally bioavailable inhibitor of blood coagulation Factor Xa. <i>Journal of Labelled Compounds and Radiopharmaceuticals</i> , 2011, 54, 418-425.	1.0	9
8	The synthesis of a carbon- ¹⁴ labeled pegylated Adnectin, for placental transfer studies in guinea pigs. <i>Journal of Labelled Compounds and Radiopharmaceuticals</i> , 2013, 56, 492-494.	1.0	5
9	The syntheses of isotopically labelled CB ¹ antagonists for the treatment of obesity. <i>Journal of Labelled Compounds and Radiopharmaceuticals</i> , 2016, 59, 665-672.	1.0	5
10	The syntheses of [¹⁴ C]BMS-823778 for use in a human ADME clinical study and of [¹³ CD ₃ ¹³ CD ₂]BMT-094817, a stable-isotope labeled standard of a newly detected human metabolite. <i>Journal of Labelled Compounds and Radiopharmaceuticals</i> , 2016, 59, 255-259.	1.0	4
11	New radical methods for the potential synthesis of carbon- ¹³ and carbon- ¹⁴ labeled complex products. <i>Journal of Labelled Compounds and Radiopharmaceuticals</i> , 2018, 61, 1024-1035.	1.0	4
12	A novel synthesis of [2- ¹⁴ C]2,5-dichloropyrimidine. <i>Journal of Labelled Compounds and Radiopharmaceuticals</i> , 2011, 54, 813-815.	1.0	3
13	The synthesis of [¹⁴ C]4-acetylphenylalanine, effect on cell viability, and assessment of protein incorporation in male rat hepatocytes. <i>Journal of Labelled Compounds and Radiopharmaceuticals</i> , 2017, 60, 352-356.	1.0	2
14	Discovery of Clinical Candidate BMS-823778 as an Inhibitor of Human 11 ^β -Hydroxysteroid Dehydrogenase Type 1 (11 ^β -HSD-1). <i>ACS Medicinal Chemistry Letters</i> , 2018, 9, 1170-1174.	2.8	2
15	An improved synthesis of [2- ¹⁴ C]2, 5-dichloropyrimidine. <i>Journal of Labelled Compounds and Radiopharmaceuticals</i> , 2012, 55, 300-302.	1.0	1
16	The synthesis of ¹⁴ C-labeled <i>N</i> -succinimidyl- ϵ -maleimidopropionate, a linker molecule for PEGylated biologics. <i>Journal of Labelled Compounds and Radiopharmaceuticals</i> , 2014, 57, 667-669.	1.0	1
17	The synthesis and analysis of [phenyl- ¹⁴ C(U)]BMS-770767 and [¹³ C ₆]BMS-770767 for use in discovery biotransformation, human ADME and bioanalytical studies. <i>Journal of Labelled Compounds and Radiopharmaceuticals</i> , 2016, 59, 657-664.	1.0	1
18	Editorial. <i>Journal of Labelled Compounds and Radiopharmaceuticals</i> , 2016, 59, 232-232.	1.0	0

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19	The syntheses of [¹³ C ₆] and [phenyl- ¹⁴ C(U)]BMS-16336, an inhibitor of 11 β -hydroxysteroid dehydrogenase type 1, for type 2 diabetes. Journal of Labelled Compounds and Radiopharmaceuticals, 2017, 60, 357-365.	1.0	0
20	The synthesis of [1- ¹⁴ C]2-(1 <i>H</i> -tetrazol-5-yl)acetic acid. Journal of Labelled Compounds and Radiopharmaceuticals, 2017, 60, 49-54.	1.0	0