

Antoine Maruani

List of Publications by Year in descending order

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Version: 2024-02-01

25
papers

1,699
citations

331670

21
h-index

552781

26
g-index

30
all docs

30
docs citations

30
times ranked

2089
citing authors

#	ARTICLE	IF	CITATIONS
1	Aerobically-initiated C(sp ³)â€“H bond amination through the use of activated azodicarboxylates. <i>Organic and Biomolecular Chemistry</i> , 2020, 18, 6258-6264.	2.8	11
2	A Plug-and-Play Approach for the <i>De Novo</i> Generation of Dually Functionalized Bispecifics. <i>Bioconjugate Chemistry</i> , 2020, 31, 520-529.	3.6	31
3	Disulfide Modified IgG1: An Investigation of Biophysical Profile and Clinically Relevant Fc Interactions. <i>Bioconjugate Chemistry</i> , 2019, 30, 1048-1054.	3.6	28
4	Use of pyridazinediones as extracellular cleavable linkers through reversible cysteine conjugation. <i>Chemical Communications</i> , 2019, 55, 14829-14832.	4.1	9
5	Highly homogeneous antibody modification through optimisation of the synthesis and conjugation of functionalised dibromopyridazinediones. <i>Organic and Biomolecular Chemistry</i> , 2018, 16, 1359-1366.	2.8	60
6	Synthesis of a novel HER2 targeted aza-BODIPYâ€“antibody conjugate: synthesis, photophysical characterisation and <i>in vitro</i> evaluation. <i>Organic and Biomolecular Chemistry</i> , 2018, 16, 1144-1149.	2.8	17
7	Assembly of High-Potency Photosensitizerâ€“Antibody Conjugates through Application of Dendron Multiplier Technology. <i>Bioconjugate Chemistry</i> , 2018, 29, 176-181.	3.6	27
8	Bispecifics and antibodyâ€“drug conjugates: A positive synergy. <i>Drug Discovery Today: Technologies</i> , 2018, 30, 55-61.	4.0	29
9	Enabling the controlled assembly of antibody conjugates with a loading of two modules without antibody engineering. <i>Chemical Science</i> , 2017, 8, 2056-2060.	7.4	52
10	Antibody fragments as nanoparticle targeting ligands: a step in the right direction. <i>Chemical Science</i> , 2017, 8, 63-77.	7.4	209
11	Pyridazinediones deliver potent, stable, targeted and efficacious antibodyâ€“drug conjugates (ADCs) with a controlled loading of 4 drugs per antibody. <i>RSC Advances</i> , 2017, 7, 9073-9077.	3.6	62
12	Dual modification of biomolecules. <i>Organic and Biomolecular Chemistry</i> , 2016, 14, 6165-6178.	2.8	50
13	A facile, one-pot procedure for the conversion of aromatic aldehydes to esters, as well as thioesters and amides, via acyl hydrazide intermediates. <i>RSC Advances</i> , 2016, 6, 3372-3376.	3.6	19
14	The Use of 3,6-Pyridazinediones in Organic Synthesis and Chemical Biology. <i>Journal of Chemical Research</i> , 2016, 40, 1-9.	1.3	21
15	Recent advances in the construction of antibodyâ€“drug conjugates. <i>Nature Chemistry</i> , 2016, 8, 114-119.	13.6	289
16	Next-generation disulfide stapling: reduction and functional re-bridging all in one. <i>Chemical Science</i> , 2016, 7, 799-802.	7.4	72
17	A platform for efficient, thiol-stable conjugation to albumin's native single accessible cysteine. <i>Organic and Biomolecular Chemistry</i> , 2015, 13, 7946-7949.	2.8	47
18	A plug-and-play approach to antibody-based therapeutics via a chemoselective dual click strategy. <i>Nature Communications</i> , 2015, 6, 6645.	12.8	203

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19	Site-selective multi-porphyrin attachment enables the formation of a next-generation antibody-based photodynamic therapeutic. <i>Chemical Communications</i> , 2015, 51, 15304-15307.	4.1	50
20	A mild TCEP-based para-azidobenzyl cleavage strategy to transform reversible cysteine thiol labelling reagents into irreversible conjugates. <i>Chemical Communications</i> , 2015, 51, 5279-5282.	4.1	42
21	A rapid, site-selective and efficient route to the dual modification of DARPs. <i>Chemical Communications</i> , 2014, 50, 4898-4900.	4.1	16
22	Regioselective and Stoichiometrically Controlled Conjugation of Photodynamic Sensitizers to a HER2 Targeting Antibody Fragment. <i>Bioconjugate Chemistry</i> , 2014, 25, 611-617.	3.6	65
23	Next generation maleimides enable the controlled assembly of antibody-drug conjugates via native disulfide bond bridging. <i>Organic and Biomolecular Chemistry</i> , 2014, 12, 7261-7269.	2.8	135
24	Acid-cleavable thiomaleamic acid linker for homogeneous antibody-drug conjugation. <i>Chemical Communications</i> , 2013, 49, 8187.	4.1	67
25	A mild synthesis of N-functionalised bromomaleimides, thiomaleimides and bromopyridazinediones. <i>Tetrahedron Letters</i> , 2013, 54, 3493-3495.	1.4	46