## Jose Manuel Otero Casas

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Highly functionalized cyclic and bicyclic βâ^'amino acids from sugar βâ^'nitroesters. Tetrahedron, 2020, 76, 130837.	1.9	0
2	Hydroxylammonium derivatives for selective active-site lysine modification in the anti-virulence bacterial target DHQ1 enzyme. Organic Chemistry Frontiers, 2019, 6, 3127-3135.	4.5	4
3	Adsorption of Pharmaceutical Pollutants from Water Using Covalent Organic Frameworks. Chemistry - A European Journal, 2018, 24, 10601-10605.	3.3	106
4	6â€Deoxyhexoses from <scp>l</scp> â€Rhamnose in the Search for Inducers of the Rhamnose Operon: Synergy of Chemistry and Biotechnology. Chemistry - A European Journal, 2016, 22, 12557-12565.	3.3	8
5	Synthetic Chemical Inducers and Genetic Decoupling Enable Orthogonal Control of the <i>rhaBAD</i> Promoter. ACS Synthetic Biology, 2016, 5, 1136-1145.	3.8	47
6	Structure of the Receptor-Binding Carboxy-Terminal Domain of the Bacteriophage T5 L-Shaped Tail Fibre with and without Its Intra-Molecular Chaperone. Viruses, 2015, 7, 6424-6440.	3.3	46
7	Chemical Modification of a Dehydratase Enzyme Involved in Bacterial Virulence by an Ammonium Derivative: Evidence of its Active Site Covalent Adduct. Journal of the American Chemical Society, 2015, 137, 9333-9343.	13.7	12
8	Irreversible covalent modification of type I dehydroquinase with a stable Schiff base. Organic and Biomolecular Chemistry, 2015, 13, 706-716.	2.8	8
9	Exploring the Water-Binding Pocket of the Type II Dehydroquinase Enzyme in the Structure-Based Design of Inhibitors. Journal of Medicinal Chemistry, 2014, 57, 3494-3510.	6.4	8
10	Insights into substrate binding and catalysis in bacterial typeÂl dehydroquinase. Biochemical Journal, 2014, 462, 415-424.	3.7	8
11	Mycobacterium tuberculosis Shikimate Kinase Inhibitors: Design and Simulation Studies of the Catalytic Turnover. Journal of the American Chemical Society, 2013, 135, 12366-12376.	13.7	51
12	Inhibiting and Reversing Amyloidâ€Î² Peptide (1–40) Fibril Formation with Gramicidinâ€S and Engineered Analogues. Chemistry - A European Journal, 2013, 19, 17338-17348.	3.3	39
13	Mechanistic Basis of the Inhibition of Type II Dehydroquinase by (2 <i>S</i> )- and (2 <i>R</i> )-2-Benzyl-3-dehydroquinic Acids. ACS Chemical Biology, 2013, 8, 568-577.	3.4	11
14	Design, Synthesis, and Structural Analysis of Turn Modified <i>cyclo</i> -(αβ <sup>3</sup> αβ <sup>2</sup> α) <sub>2</sub> Peptide Derivatives toward Crystalline Hexagon-Shaped Cationic Nanochannel Assemblies. Crystal Growth and Design, 2013, 13, 4355-4367.	3.0	6
15	An overview of key routes for the transformation of sugars into carbasugars and related compounds. Carbohydrate Chemistry, 2012, , 263-302.	0.3	13
16	High-resolution structures of <i>Thermus thermophilus</i> enoyl-acyl carrier protein reductase in the apo form, in complex with NAD <sup>+</sup> and in complex with NAD <sup>+</sup> and triclosan. Acta Crystallographica Section F: Structural Biology Communications, 2012, 68, 1139-1148.	0.7	7
17	â€~Inverted' analogs of the antibiotic gramicidin S with an improved biological profile. Bioorganic and Medicinal Chemistry, 2012, 20, 6059-6062.	3.0	8
18	A Nitro Sugarâ€Mediated Stereocontrolled Synthesis of β <sup>2</sup> â€Amino Acids: Synthesis of a Polyhydroxylated <i>trans</i> â€2â€AminoÂcyclohexanecarboxylic Acid. European Journal of Organic Chemistry, 2012, 2012, 2969-2979.	2.4	3

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19	Studies on the Michael addition of naphthoquinones to sugar nitro olefins: first synthesis of polyhydroxylated hexahydro-11H-benzo[a]carbazole-5,6-diones and hexahydro-11bH-benzo[b]carbazole-6,11-diones. Tetrahedron, 2012, 68, 1612-1621.	1.9	15
20	Design, synthesis and structural analysis of mixed $\hat{I}\pm/\hat{I}^2$ -peptides that adopt stable cyclic hairpin-like conformations. Tetrahedron, 2012, 68, 2391-2400.	1.9	12
21	On a Possible Neutral Charge State for the Catalytic Dyad in Î <sup>2</sup> -Secretase When Bound to Hydroxyethylene Transition State Analogue Inhibitors. Journal of Medicinal Chemistry, 2011, 54, 3081-3085.	6.4	13
22	A Prodrug Approach for Improving Antituberculosis Activity of Potent Mycobacterium tuberculosis Type II Dehydroquinase Inhibitors. Journal of Medicinal Chemistry, 2011, 54, 6063-6084.	6.4	32
23	Target highlights in CASP9: Experimental target structures for the critical assessment of techniques for protein structure prediction. Proteins: Structure, Function and Bioinformatics, 2011, 79, 6-20.	2.6	19
24	Tetrahydrobenzothiophene Derivatives: Conformationally Restricted Inhibitors of Typeâ€II Dehydroquinase. ChemMedChem, 2011, 6, 266-272.	3.2	15
25	Evaluation of Readily Accessible Azoles as Mimics of the Aromatic Ring of <scp>D</scp> â€Phenylalanine in the Turn Region of Gramicidinâ€S. ChemMedChem, 2011, 6, 840-847.	3.2	17
26	Exploring the Conformational and Biological Versatility of βâ€Turnâ€Modified Gramicidin S by Using Sugar Amino Acid Homologues that Vary in Ring Size. Chemistry - A European Journal, 2011, 17, 3995-4004.	3.3	33
27	Synthesis and evaluation of strand and turn modified ring-extended gramicidin S derivatives. Bioorganic and Medicinal Chemistry, 2011, 19, 3402-3409.	3.0	9
28	Understanding the Key Factors that Control the Inhibition of Typeâ€II Dehydroquinase by (2 <i>R</i> )â€2â€Benzylâ€3â€dehydroquinic Acids. ChemMedChem, 2010, 5, 1726-1733.	3.2	22
29	Gramicidin S Derivatives Containing <i>cis</i> ―and <i>trans</i> â€Morpholine Amino Acids (MAAs) as Turn Mimetics. Chemistry - A European Journal, 2010, 16, 4259-4265.	3.3	15
30	An Adamantyl Amino Acid Containing Gramicidinâ€S Analogue with Broad Spectrum Antibacterial Activity and Reduced Hemolytic Activity. Chemistry - A European Journal, 2010, 16, 12174-12181.	3.3	33
31	Studies on the transformation of nitrosugars into iminosugars III: synthesis of (2R,3R,4R,5R,6R)-2-(hydroxymethyl)azepane-3,4,5,6-tetraol and (2R,3R,4R,5R,6S)-2-(hydroxymethyl)azepane-3,4,5,6-tetraol. Tetrahedron: Asymmetry, 2010, 21, 21-26.	1.8	17
32	Effect of the Protonation State of the Titratable Residues on the Inhibitor Affinity to BACE-1. Biochemistry, 2010, 49, 7255-7263.	2.5	51
33	Structure of the bacteriophage T4 long tail fiber receptor-binding tip. Proceedings of the National Academy of Sciences of the United States of America, 2010, 107, 20287-20292.	7.1	159
34	Synthesis and Biological Evaluation of New Nanomolar Competitive Inhibitors of Helicobacter pylori Type II Dehydroquinase. Structural Details of the Role of the Aromatic Moieties with Essential Residues. Journal of Medicinal Chemistry, 2010, 53, 191-200.	6.4	21
35	Bisphosphineâ€Functionalized Cyclic Decapeptides Based on the Natural Product Gramicidinâ€S: A Potential Scaffold for Transitionâ€Metal Coordination. Chemistry - A European Journal, 2009, 15, 8134-8145.	3.3	17
36	Synthesis and biological evaluation of asymmetric gramicidin S analogues containing modified d-phenylalanine residues. Bioorganic and Medicinal Chemistry, 2009, 17, 6318-6328.	3.0	14

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37	Studies on the transformation of nitrosugars into branched chain iminosugars. Part II: Synthesis of (3R,4R,5R,6S)-2,2-bis(hydroxymethyl)azepane-3,4,5,6-tetraol. Tetrahedron: Asymmetry, 2008, 19, 2443-2446.	1.8	22
38	Preliminary Studies on the Michael Addition of Quinones to Nitroolefins: (6bR,10aS)-7,8,9,10,10a,11-Hexahydro-6bH-benzo[a]carbazole-5,6-diones, (4aR,11bS)-1,2,3,4,4a,5-Hexahydro-11bH-benzo[b]carbazole-6,11-diones, and 1,2,3,4-Tetrahydro-5H-benzo[b]carbazole-6,11-diones, Synlett, 2007, 2007, 1399-1402.	1.8	1
39	Preliminary Studies on the Transformation of Nitrosugars into Branched Chain Iminosugars: Synthesis of 1,4-Dideoxy-4-C-hydroxymethyl- 1,4-imino-pentanols. Organic Letters, 2007, 9, 623-626.	4.6	25
40	Total synthesis of (5S,6S)-6-amino-2,8-dimethylnonan-5-ol and (5S,6S)-6-amino-7-cyclohexyl-2-methylheptan-5-ol. Arkivoc, 2007, 2007, 380-388.	0.5	0
41	Preliminary studies on a novel synthesis of β-amino acids: stereocontrolled transformation of d- and l-glyceraldehyde into 3-amino-2-(2′,2′-dimethyl-1′,3′-dioxolan-4′-yl)propanoic acids. Tetrahedron: Asymmetry, 2006, 17, 3063-3066.	1.8	9
42	Preliminary studies on the incorporation of sugars into naphthoquinones: synthesis of (1R,2S,3S,4R,4aS,11bS)-2-(benzyloxy)-1,2,3,4,4a,5-hexahydro-1,3,4-trihydroxy-11bH-benzo[b]carbazole-6,11-dior Tetrahedron: Asymmetry, 2005, 16, 11-14.	ne1.8	11
43	Preliminary studies on the synthesis of rancinamycins from nitrosugars: first total synthesis of (3S,4S,5S,6R)-5-benzyloxy-6-hydroxy-3,4-(isopropylidendioxy)-cyclohex-1-enecarbaldehyde. Tetrahedron: Asymmetry, 2005, 16, 4045-4049.	1.8	11