

# Jordi Bujons

## List of Publications by Year in descending order

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80  
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citations

218592

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#	ARTICLE	IF	CITATIONS
1	Dynamic Combinatorial Optimization of <i>In Vitro</i> and <i>In Vivo</i> Heparin Antidotes. <i>Journal of Medicinal Chemistry</i> , 2022, 65, 4865-4877.	2.9	3
2	Chemoenzymatic Production of Enantiocomplementary $\alpha$ -Substituted $\beta$ -Hydroxycarboxylic Acids from $\alpha$ -Amino Acids. <i>Advanced Synthesis and Catalysis</i> , 2021, 363, 2866-2876.	2.1	7
3	Synthesis of $\beta$ -Hydroxy- $\alpha$ -amino Acid Derivatives by Enzymatic Tandem Aldol Addition-Transamination Reactions. <i>ACS Catalysis</i> , 2021, 11, 4660-4669.	5.5	25
4	Semaphorin 3A-Glycosaminoglycans Interaction as Therapeutic Target for Axonal Regeneration. <i>Pharmaceuticals</i> , 2021, 14, 906.	1.7	3
5	Inhibition of Sema-3A Promotes Cell Migration, Axonal Growth, and Retinal Ganglion Cell Survival. <i>Translational Vision Science and Technology</i> , 2021, 10, 16.	1.1	2
6	Titelbild: Live-Cell-Templated Dynamic Combinatorial Chemistry ( <i>Angew. Chem.</i> 39/2020). <i>Angewandte Chemie</i> , 2020, 132, 16949-16949.	1.6	0
7	Biocatalytic Construction of Quaternary Centers by Aldol Addition of 3,3-Disubstituted 2-Oxoacid Derivatives to Aldehydes. <i>Journal of the American Chemical Society</i> , 2020, 142, 19754-19762.	6.6	10
8	Live-Cell-Templated Dynamic Combinatorial Chemistry. <i>Angewandte Chemie</i> , 2020, 132, 17355-17359.	1.6	5
9	Live-Cell-Templated Dynamic Combinatorial Chemistry. <i>Angewandte Chemie - International Edition</i> , 2020, 59, 17202-17206.	7.2	20
10	Chemoenzymatic Hydroxymethylation of Carboxylic Acids by Tandem Stereodivergent Biocatalytic Aldol Reaction and Chemical Decarboxylation. <i>ACS Catalysis</i> , 2019, 9, 7568-7577.	5.5	15
11	Aldolase-Catalyzed Asymmetric Synthesis of $N$ -Heterocycles by Addition of Simple Aliphatic Nucleophiles to Aminoaldehydes. <i>Advanced Synthesis and Catalysis</i> , 2019, 361, 2673-2687.	2.1	19
12	Identification of benzo[ <i>cd</i> ]indol-2(1 <i>H</i> )-ones as novel Atg4B inhibitors via a structure-based virtual screening and a novel AlphaScreen assay. <i>European Journal of Medicinal Chemistry</i> , 2019, 178, 648-666.	2.6	15
13	Nucleophile Promiscuity of Engineered Class II Pyruvate Aldolase YfaU from <i>E. coli</i> . <i>Angewandte Chemie</i> , 2018, 130, 3645-3649.	1.6	11
14	Nucleophile Promiscuity of Engineered Class II Pyruvate Aldolase YfaU from <i>E. coli</i> . <i>Angewandte Chemie - International Edition</i> , 2018, 57, 3583-3587.	7.2	22
15	Titelbild: Nucleophile Promiscuity of Engineered Class II Pyruvate Aldolase YfaU from <i>E. coli</i> ( <i>Angew. Chem.</i> 14/2018). <i>Angewandte Chemie</i> , 2018, 130, 3581-3581.	1.6	0
16	Dynamic Covalent Identification of an Efficient Heparin Ligand. <i>Angewandte Chemie</i> , 2018, 130, 12149-12153.	1.6	8
17	Dynamic Covalent Identification of an Efficient Heparin Ligand. <i>Angewandte Chemie - International Edition</i> , 2018, 57, 11973-11977.	7.2	20
18	Biocatalytic Aldol Addition of Simple Aliphatic Nucleophiles to Hydroxyaldehydes. <i>ACS Catalysis</i> , 2018, 8, 8804-8809.	5.5	25

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19	Combining Aldolases and Transaminases for the Synthesis of 2-Amino-4-hydroxybutanoic Acid. <i>ACS Catalysis</i> , 2017, 7, 1707-1711.	5.5	60
20	The first fluorogenic sensor for sphingosine-1-phosphate lyase activity in intact cells. <i>Chemical Communications</i> , 2017, 53, 5441-5444.	2.2	12
21	2-Keto-3-Deoxy- $\alpha$ -Rhamnonate Aldolase (YfaU) as Catalyst in Aldol Additions of Pyruvate to Amino Aldehyde Derivatives. <i>Advanced Synthesis and Catalysis</i> , 2017, 359, 2090-2100.	2.1	20
22	Entropy-driven homochiral self-sorting of a dynamic library. <i>Chemical Communications</i> , 2017, 53, 4274-4277.	2.2	17
23	Intramolecular Benzoin Reaction Catalyzed by Benzaldehyde Lyase from <i>Pseudomonas Fluorescens</i> Biovar I. <i>Angewandte Chemie - International Edition</i> , 2017, 56, 5304-5307.	7.2	13
24	Cationic Peptides and Peptidomimetics Bind Glycosaminoglycans as Potential Sema3A Pathway Inhibitors. <i>Biophysical Journal</i> , 2016, 110, 1291-1303.	0.2	17
25	Studies on the inhibition of sphingosine-1-phosphate lyase by stabilized reaction intermediates and stereodefined azido phosphates. <i>European Journal of Medicinal Chemistry</i> , 2016, 123, 905-915.	2.6	2
26	Bacterial versus human sphingosine-1-phosphate lyase (S1PL) in the design of potential S1PL inhibitors. <i>Bioorganic and Medicinal Chemistry</i> , 2016, 24, 4381-4389.	1.4	3
27	Positional Scanning Synthesis of a Peptoid Library Yields New Inducers of Apoptosis that Target Karyopherins and Tubulin. <i>ChemBioChem</i> , 2015, 16, 1580-1587.	1.3	10
28	Structure-Guided Engineering of <i>D</i> -Fructose-6-Phosphate Aldolase for Improved Acceptor Tolerance in Biocatalytic Aldol Additions. <i>Advanced Synthesis and Catalysis</i> , 2015, 357, 1787-1807.	2.1	20
29	Salt-Induced Adaptation of a Dynamic Combinatorial Library of Pseudopeptidic Macrocycles: Unraveling the Electrostatic Effects in Mixed Aqueous Media. <i>Chemistry - A European Journal</i> , 2015, 21, 6869-6878.	1.7	30
30	Efficient Synthesis of Conformationally Restricted Apoptosis Inhibitors Bearing a Triazole Moiety. <i>Chemistry - A European Journal</i> , 2015, 21, 14122-14128.	1.7	12
31	Engineered <i>L</i> -Serine Hydroxymethyltransferase from <i>Streptococcus thermophilus</i> for the Synthesis of $\pm$ -Dialkyl- $\alpha$ -Amino Acids. <i>Angewandte Chemie</i> , 2015, 127, 3056-3060.	1.6	12
32	Expedient Synthesis of C-Aryl Carbohydrates by Consecutive Biocatalytic Benzoin and Aldol Reactions. <i>Chemistry - A European Journal</i> , 2015, 21, 3335-3346.	1.7	13
33	Engineered <i>L</i> -Serine Hydroxymethyltransferase from <i>Streptococcus thermophilus</i> for the Synthesis of $\pm$ -Dialkyl- $\alpha$ -Amino Acids. <i>Angewandte Chemie - International Edition</i> , 2015, 54, 3013-3017.	7.2	35
34	Synthesis and Evaluation of Hydroxymethylaminocyclitols as Glycosidase Inhibitors. <i>Journal of Organic Chemistry</i> , 2015, 80, 3512-3529.	1.7	17
35	Asymmetric assembly of aldose carbohydrates from formaldehyde and glycolaldehyde by tandem biocatalytic aldol reactions. <i>Nature Chemistry</i> , 2015, 7, 724-729.	6.6	63
36	Apaf-1 Inhibitors Protect from Unwanted Cell Death in In Vivo Models of Kidney Ischemia and Chemotherapy Induced Ototoxicity. <i>PLoS ONE</i> , 2014, 9, e110979.	1.1	22

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37	Inhibition of dihydroceramide desaturase activity by the sphingosine kinase inhibitor SKI II. <i>Journal of Lipid Research</i> , 2014, 55, 1711-1720.	2.0	66
38	Engineering the Donor Selectivity of <i>D</i> -Fructose-6-Phosphate Aldolase for Biocatalytic Asymmetric Cross-Aldol Additions of Glycolaldehyde. <i>Chemistry - A European Journal</i> , 2014, 20, 12572-12583.	1.7	35
39	Selective chaperone effect of aminocyclitol derivatives on G202R and other mutant glucocerebrosidases causing Gaucher disease. <i>International Journal of Biochemistry and Cell Biology</i> , 2014, 54, 245-254.	1.2	8
40	Sequential Biocatalytic Aldol Reactions in Multistep Asymmetric Synthesis: Pipecolic Acid, Piperidine and Pyrrolidine (Homo)Iminocyclitol Derivatives from Achiral Building Blocks. <i>Advanced Synthesis and Catalysis</i> , 2014, 356, 3007-3024.	2.1	31
41	Optimizing the control of apoptosis by amide/triazole isosteric substitution in a constrained peptoid. <i>European Journal of Medicinal Chemistry</i> , 2013, 63, 892-896.	2.6	18
42	<sup>15</sup> N NMR spectroscopic and theoretical GIAO-DFT studies for the unambiguous characterization of disubstituted 1,2,3-triazoles. <i>Organic and Biomolecular Chemistry</i> , 2013, 11, 7318.	1.5	15
43	Chemo-enzymatic synthesis and glycosidase inhibitory properties of DAB and LAB derivatives. <i>Organic and Biomolecular Chemistry</i> , 2013, 11, 2005.	1.5	25
44	Allopregnanolone and Pregnanolone Analogues Modified in the C Ring: Synthesis and Activity. <i>Journal of Medicinal Chemistry</i> , 2013, 56, 2323-2336.	2.9	11
45	Chemoenzymatic synthesis, structural study and biological activity of novel indolizidine and quinolizidine iminocyclitols. <i>Organic and Biomolecular Chemistry</i> , 2012, 10, 6309.	1.5	30
46	Highly efficient aldol additions of DHA and DHAP to N-Cbz-amino aldehydes catalyzed by l-rhamnulose-1-phosphate and l-fuculose-1-phosphate aldolases in aqueous borate buffer. <i>Organic and Biomolecular Chemistry</i> , 2011, 9, 8430.	1.5	26
47	Structure-guided redesign of d-fructose-6-phosphate aldolase from <i>E. coli</i> : remarkable activity and selectivity towards acceptor substrates by two-point mutation. <i>Chemical Communications</i> , 2011, 47, 5762.	2.2	41
48	Computational Prediction of Structure-Activity Relationships for the Binding of Aminocyclitols to $\beta$ -Glucocerebrosidase. <i>Journal of Chemical Information and Modeling</i> , 2011, 51, 601-611.	2.5	14
49	New Glucocerebrosidase Inhibitors by Exploration of Chemical Diversity of <i>N</i> -Substituted Aminocyclitols Using Click Chemistry and in Situ Screening. <i>Journal of Medicinal Chemistry</i> , 2011, 54, 2069-2079.	2.9	36
50	Triazine-Based Vanilloid 1 Receptor Open Channel Blockers: Design, Synthesis, Evaluation, and SAR Analysis. <i>Journal of Medicinal Chemistry</i> , 2011, 54, 7441-7452.	2.9	21
51	Redesign of the Phosphate Binding Site of <i>L</i> -Rhamnulose-1-Phosphate Aldolase towards a Dihydroxyacetone Dependent Aldolase. <i>Advanced Synthesis and Catalysis</i> , 2011, 353, 89-99.	2.1	38
52	Chemical Modulation of Peptoids: Synthesis and Conformational Studies on Partially Constrained Derivatives. <i>Chemistry - A European Journal</i> , 2011, 17, 7927-7939.	1.7	33
53	Structure-Guided Minimalist Redesign of the <i>L</i> -Fuculose-1-Phosphate Aldolase Active Site: Expedient Synthesis of Novel Polyhydroxylated Pyrrolizidines and their Inhibitory Properties Against Glycosidases and Intestinal Disaccharidases. <i>Chemistry - A European Journal</i> , 2010, 16, 10691-10706.	1.7	39
54	Click Chemistry Approach to New <i>N</i> -Substituted Aminocyclitols as Potential Pharmacological Chaperones for Gaucher Disease. <i>Journal of Medicinal Chemistry</i> , 2010, 53, 5248-5255.	2.9	55

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55	Dihydroxyacetone Phosphate Aldolase Catalyzed Synthesis of Structurally Diverse Polyhydroxylated Pyrrolidine Derivatives and Evaluation of their Glycosidase Inhibitory Properties. <i>Chemistry - A European Journal</i> , 2009, 15, 7310-7328.	1.7	49
56	Novel separation of bioactive catechin derivatives from complex plant mixtures by anion-exchange chromatography. <i>Separation and Purification Technology</i> , 2008, 62, 317-322.	3.9	7
57	Activity of B-Nor Analogues of Neurosteroids on the GABAAR Receptor in Primary Neuronal Cultures. <i>Journal of Medicinal Chemistry</i> , 2006, 49, 3225-3234.	2.9	27
58	Biosynthesis of 10,12-dienoic fatty acids by a bifunctional $\Delta^11$ desaturase in <i>Spodoptera littoralis</i> . <i>Insect Biochemistry and Molecular Biology</i> , 2006, 36, 634-641.	1.2	29
59	Allosteric positive interaction of thymol with the GABAA receptor in primary cultures of mouse cortical neurons. <i>Neuropharmacology</i> , 2006, 50, 25-35.	2.0	113
60	Influence of N-amino protecting group on aldolase-catalyzed aldol additions of dihydroxyacetone phosphate to amino aldehydes. <i>Tetrahedron</i> , 2006, 62, 2648-2656.	1.0	25
61	Aldol Additions of Dihydroxyacetone Phosphate to N-Cbz-Amino Aldehydes Catalyzed by L-Fuculose-1-Phosphate Aldolase in Emulsion Systems: Inversion of Stereoselectivity as a Function of the Acceptor Aldehyde. <i>Chemistry - A European Journal</i> , 2005, 11, 1392-1401.	1.7	50
62	Studies on Toxic Oil Syndrome: $\Delta$ Stereoselective Hydrolysis of 3-(Phenylamino)propane-1,2-diol Esters by Human Pancreatic Lipase. <i>Chemical Research in Toxicology</i> , 2004, 17, 889-895.	1.7	7
63	Stereoselective Aldol Additions Catalyzed by Dihydroxyacetone Phosphate-Dependent Aldolases in Emulsion Systems: Preparation and Structural Characterization of Linear and Cyclic Iminopolys from Aminoaldehydes. <i>Chemistry - A European Journal</i> , 2003, 9, 4887-4899.	1.7	88
64	The organochlorine pesticides $\Delta^3$ -hexachlorocyclohexane (lindane), $\Delta^{\pm}$ -endosulfan and dieldrin differentially interact with GABAA and glycine-gated chloride channels in primary cultures of cerebellar granule cells. <i>Neuroscience</i> , 2003, 117, 397-403.	1.1	83
65	Metabolism of (R)- and (S)-3-(Phenylamino)propane-1,2-diol in C57BL/6- and A/J-Strain Mice. Identification of New Metabolites with Potential Toxicological Significance to the Toxic Oil Syndrome. <i>Chemical Research in Toxicology</i> , 2001, 14, 1097-1106.	1.7	15
66	Metabolism of R,S Enantiomers of 3-Phenylamino-1,2-Propanediol, a Compound Associated with the Toxic Oil Syndrome, in C57BL/6- and A/J-Strain Mice. <i>Advances in Experimental Medicine and Biology</i> , 2001, 500, 525-529.	0.8	1
67	Cytochrome c Peroxidase $\Delta$ Cytochrome c Complex: $\Delta$ Locating the Second Binding Domain on Cytochrome c Peroxidase with Site-Directed Mutagenesis. <i>Biochemistry</i> , 2000, 39, 10132-10139.	1.2	52
68	Mutants That Alter the Covalent Structure of Catalase Hydroperoxidase II from <i>Escherichia coli</i> xs. <i>Journal of Biological Chemistry</i> , 1999, 274, 27717-27725.	1.6	30
69	Biotransformation and Clearance of 3-(Phenylamino)propane-1,2-diol, a Compound Present in Samples Related to Toxic Oil Syndrome, in C57BL/6 and A/J Mice. <i>Chemical Research in Toxicology</i> , 1999, 12, 1127-1137.	1.7	18
70	$\Delta^{\mu}$ , $\Delta^{\mu}$ -Dimethyl-lysine cytochrome c as an NMR probe for lysine involvement in protein $\Delta$ protein complex formation. <i>Biochemical Journal</i> , 1998, 332, 439-449.	1.7	24
71	Charge Reversal of a Critical Active-Site Residue of Cytochrome-c Peroxidase. Characterization of the Arg48Glu Variant. <i>FEBS Journal</i> , 1997, 243, 72-84.	0.2	21
72	Mutagenic evidence concerning the location of two binding sites for cytochrome C on the surface of cytochrome C peroxidase. <i>Journal of Inorganic Biochemistry</i> , 1995, 59, 432.	1.5	0

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73	Aflatoxin M1 8,9-Epoxyde: Preparation and Mutagenic Activity. <i>Chemical Research in Toxicology</i> , 1995, 8, 328-332.	1.7	12
74	A Convenient Synthesis of the Toxicophoric Furo[2,3-b]benzofuran Fragment Present in Aflatoxin M1. <i>Synlett</i> , 1994, 1994, 437-438.	1.0	4
75	A study of the interconversion between 3,4-dihydro-4-formyl-2-hydroxy-2H-benzopyran and 2,3,3a,8a-tetrahydro-2-hydroxyfuro[2,3-b]benzofuran moieties, and its application to a formal synthesis of (±)-aflatoxin B1. <i>Tetrahedron</i> , 1994, 50, 7597-7610.	1.0	10
76	A convenient entry to the toxicophoric furo[2,3-b] benzofuran fragment present in aflatoxins. <i>Tetrahedron Letters</i> , 1992, 33, 6387-6388.	0.7	4
77	Synthesis and mutagenicity of the aflatoxin B1 model 3a,8a-dihydro-4,6-dimethoxyfuro[2,3-b]benzofuran and its 2,3-epoxy derivative. <i>Journal of Agricultural and Food Chemistry</i> , 1991, 39, 1723-1728.	2.4	5
78	Use of dimethyldioxirane in the preparation of highly reactive compounds: First direct epoxidation of precocenes. <i>Tetrahedron Letters</i> , 1990, 31, 5235-5236.	0.7	35
79	NEW CARBON-PHOSPHORUS BOND CONTAINING COMPOUND IN THE REACTION OF 2-ACETYLTHIOPHENE WITH PHOSPHORUS PENTACHLORIDE. <i>Phosphorus, Sulfur and Silicon and the Related Elements</i> , 1990, 54, 205-207.	0.8	1
80	Enantioselective enzymic sterol synthesis by ultrasonically stimulated bakers' yeast. <i>Journal of the American Chemical Society</i> , 1988, 110, 604-606.	6.6	47