

# Francisco J Sayago

## List of Publications by Year in descending order

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times ranked

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#	ARTICLE	IF	CITATIONS
1	New approaches towards the synthesis of 1,2,3,4-tetrahydro isoquinoline-3-phosphonic acid (TicP). <i>Amino Acids</i> , 2021, 53, 451-459.	2.7	6
2	Synthesis and biological activity of dehydrophos derivatives. <i>Organic and Biomolecular Chemistry</i> , 2019, 17, 1097-1112.	2.8	3
3	1-aminovinylphosphonate Esters as Substrates for the Diels-Alder Reaction: First Synthetic and Theoretical Study. <i>European Journal of Organic Chemistry</i> , 2019, 2019, 1268-1272.	2.4	3
4	An Improved Synthesis of the Antibiotic Dehydrophos. <i>European Journal of Organic Chemistry</i> , 2018, 2018, 3965-3973.	2.4	5
5	Synthesis of trans-fused Octahydroisoindole-1-carboxylic Acids. <i>Letters in Organic Chemistry</i> , 2018, 15, 404-411.	0.5	3
6	Stereoselective Synthesis of $\hat{\pm}$ -Amino-H-phosphinic Acids and Derivatives. <i>Synthesis</i> , 2017, 49, 987-997.	2.3	7
7	Ru-catalyzed C-H functionalization of phenylglycine derivatives: Synthesis of isoquinoline-1-carboxylates and isoindoline-1-carboxylates. <i>Journal of Molecular Catalysis A</i> , 2017, 426, 407-418.	4.8	16
8	Stereoselective Synthesis of $\hat{\pm}$ -Amino-C-phosphinic Acids and Derivatives. <i>Molecules</i> , 2016, 21, 1141.	3.8	24
9	First Synthesis of ( <i>R</i> )- and ( <i>S</i> )-1,2,3,4-tetrahydroisoquinoline- $\beta$ -phosphonic Acid (Tic <sup>P</sup> ) Using a Pictet-Spengler Reaction. <i>European Journal of Organic Chemistry</i> , 2016, 2016, 2711-2719.	2.4	13
10	Lipase-catalyzed dynamic kinetic resolution of dimethyl (1,3-dihydro-2H-isoindol-1-yl)phosphonate. <i>Tetrahedron</i> , 2016, 72, 7311-7316.	1.9	7
11	Enzymatic and chromatographic resolution procedures applied to the synthesis of the phosphoprolin enantiomers. <i>Tetrahedron: Asymmetry</i> , 2015, 26, 1469-1477.	1.8	10
12	Synthesis of [ <i>c</i> ]-Fused Bicyclic Proline Analogues. <i>European Journal of Organic Chemistry</i> , 2015, 2015, 1633-1658.	2.4	8
13	An update on the stereoselective synthesis of $\hat{\pm}$ -aminophosphonic acids and derivatives. <i>Tetrahedron</i> , 2015, 71, 1745-1784.	1.9	82
14	First Practical and Efficient Synthesis of 3-Phosphorylated $\hat{\pm}$ -Carboline Derivatives Using the Pictet-Spengler Reaction. <i>European Journal of Organic Chemistry</i> , 2015, 2015, 1084-1091.	2.4	11
15	Remote Substituent Effects on the Stereoselectivity and Organocatalytic Activity of Densely Substituted Unnatural Proline Esters in Aldol Reactions. <i>European Journal of Organic Chemistry</i> , 2015, 2015, 2503-2516.	2.4	23
16	Amide-triazole isosteric substitution for tuning self-assembly and incorporating new functions into soft supramolecular materials. <i>Chemical Communications</i> , 2015, 51, 5294-5297.	4.1	45
17	Aldolase-Catalyzed Synthesis of Conformationally Constrained Iminocyclitols: Preparation of Polyhydroxylated Benzopyrrolizidines and Cyclohexapyrrolizidines. <i>Organic Letters</i> , 2014, 16, 1422-1425.	4.6	17
18	Cyclopalladation and Reactivity of Amino Esters through C-H Bond Activation: Experimental, Kinetic, and Density Functional Theory Mechanistic Studies. <i>Chemistry - A European Journal</i> , 2013, 19, 17398-17412.	3.3	30

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19	Engineering strategy to improve peptide analogs: from structure-based computational design to tumor homing. <i>Journal of Computer-Aided Molecular Design</i> , 2013, 27, 31-43.	2.9	14
20	Synthesis of quaternary $\hat{\pm}$ -aminophosphonic acids. <i>Tetrahedron</i> , 2012, 68, 6369-6412.	1.9	82
21	Dynamic Kinetic Resolution of 1,3-Dihydro-2H-isoindole-1-carboxylic Acid Methyl Ester: Asymmetric Transformations toward Isoindoline Carbamates. <i>Organic Letters</i> , 2012, 14, 1696-1699.	4.6	28
22	Quantum-chemical predictions of redox potentials of carbamates in methanol. <i>Physical Chemistry Chemical Physics</i> , 2011, 13, 17696.	2.8	16
23	Access to the <i>cis</i> -Fused Stereoisomers of Proline Analogues Containing an Octahydroindole Core. <i>European Journal of Organic Chemistry</i> , 2011, 2011, 2011-2028.	2.4	32
24	Stereodivergent Synthesis of Two Novel $\hat{\pm}$ -Aminophosphonic Acids Characterised by a <i>cis</i> -Fused Octahydroindole System. <i>European Journal of Organic Chemistry</i> , 2011, 2011, 3074-3081.	2.4	27
25	Synthesis of Phosphopline Derivatives with an Octahydroisoindole Structure. <i>European Journal of Organic Chemistry</i> , 2011, 2011, 6732-6738.	2.4	28
26	Practical access to the proline analogs ( <i>S</i> , <i>S</i> , <i>S</i> ) and ( <i>R</i> , <i>R</i> , <i>R</i> )-methyl octahydroindole-2-carboxylic acids by HPLC enantioseparation. <i>Chirality</i> , 2011, 23, 507-513.	2.6	15
27	Efficient access to (1H)-isoindolin-1-one-3-carboxylic acid derivatives by orthopalladation and carbonylation of methyl arylglycinate substrates. <i>Tetrahedron</i> , 2011, 67, 4185-4191.	1.9	34
28	Nanoparticle-induced vascular blockade in human prostate cancer. <i>Blood</i> , 2010, 116, 2847-2856.	1.4	149
29	A straightforward route to enantiopure $\hat{\pm}$ -substituted derivatives of (2S,3aS,7aS)-octahydroindole-2-carboxylic acid. <i>Tetrahedron</i> , 2009, 65, 5174-5180.	1.9	10
30	Towards the stereoselective synthesis of $\hat{\pm}$ -methylated (2S,3aS,7aS)-octahydroindole-2-carboxylic acid. <i>Tetrahedron: Asymmetry</i> , 2008, 19, 2763-2766.	1.8	4
31	Versatile methodology for the synthesis and $\hat{\pm}$ -functionalization of (2R,3aS,7aS)-octahydroindole-2-carboxylic acid. <i>Tetrahedron</i> , 2008, 64, 84-91.	1.9	18
32	Efficient access to enantiomerically pure cyclic $\hat{\pm}$ -amino esters through a lipase-catalyzed kinetic resolution. <i>Tetrahedron: Asymmetry</i> , 2008, 19, 1714-1719.	1.8	22
33	Stereocontrolled synthesis of iminocyclitols with an ether bridge. <i>Tetrahedron</i> , 2007, 63, 4695-4702.	1.9	6
34	Efficient access to N-protected derivatives of (R,R,R)- and (S,S,S)-octahydroindole-2-carboxylic acid by HPLC resolution. <i>Tetrahedron: Asymmetry</i> , 2007, 18, 2358-2364.	1.8	18
35	Expedient synthesis of sulfoazetidino spiro-C-glycosides from ketose acetals. <i>Tetrahedron</i> , 2006, 62, 915-921.	1.9	18
36	Anhydroazasugars as key intermediates in the stereocontrolled preparation of azasugars and their ethyl thioglycosides. <i>Tetrahedron: Asymmetry</i> , 2004, 15, 603-615.	1.8	12

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37	Ring contraction of glycopyranosyl enamines: an easy route to furanoid thioglycosides of 5-aminosugars. <i>Tetrahedron: Asymmetry</i> , 2004, 15, 2003-2010.	1.8	3
38	d-Ribofuranosylenamine: a versatile starting material for preparing azasugar thioglycosides and building blocks for thioureylene-di-nucleosides. <i>Tetrahedron: Asymmetry</i> , 2004, 15, 3783-3789.	1.8	11
39	Stereoselective synthesis of azasugar thioglycosides. <i>Tetrahedron Letters</i> , 2003, 44, 6605-6608.	1.4	10
40	An easy route to seven-membered iminocyclitols from aldohexopyranosyl enamines. <i>Tetrahedron: Asymmetry</i> , 2002, 13, 1743-1753.	1.8	37