

Friedrich Hammerschmidt

List of Publications by Year in descending order

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#	ARTICLE	IF	CITATIONS
1	Biosynthesis of the Fungal Organophosphonate Fosfonochlorin Involves an Iron(II) and 2-oxoglutarate Dependent Oxacyclase. <i>ChemBioChem</i> , 2022, 23, .	2.6	8
2	Essential Functional Interplay of the Catalytic Groups in Acid Phosphatase. <i>ACS Catalysis</i> , 2022, 12, 3357-3370.	11.2	5
3	Substrate Tunnel Engineering Aided by X-ray Crystallography and Functional Dynamics Swaps the Function of MIO-Enzymes. <i>ACS Catalysis</i> , 2021, 11, 4538-4549.	11.2	21
4	Overall Retention of Methyl Stereochemistry during B12-Dependent Radical SAM Methyl Transfer in Fosfomycin Biosynthesis. <i>Biochemistry</i> , 2021, 60, 1587-1596.	2.5	6
5	C-H Bond Cleavage Is Rate-Limiting for Oxidative C-P Bond Cleavage by the Mixed Valence Diiron-Dependent Oxygenase PhnZ. <i>Biochemistry</i> , 2019, 58, 5271-5280.	2.5	11
6	2-Nitroimidazole-Furanoside Derivatives for Hypoxia Imaging—Investigation of Nucleoside Transporter Interaction, 18F-Labeling and Preclinical PET Imaging. <i>Pharmaceuticals</i> , 2019, 12, 31.	3.8	5
7	An Oxidative Pathway for Microbial Utilization of Methylphosphonic Acid as a Phosphate Source. <i>ACS Chemical Biology</i> , 2019, 14, 735-741.	3.4	16
8	Preparation of Phosphonic Acid Analogues of Proline and Proline Analogues and Their Biological Evaluation as 1-Pyrroline-5-carboxylate Reductase Inhibitors. <i>ACS Omega</i> , 2018, 3, 4441-4452.	3.5	6
9	On the rearrangement of N-aryl-N-Boc-phosphoramidates to N-Boc-protected o-aminoarylphosphonates. <i>Monatshefte für Chemie</i> , 2018, 149, 87-98.	1.8	4
10	The 1-hydroxyphosphonate-phosphate rearrangement of a noncyclic substrate — some new observations. <i>Organic and Biomolecular Chemistry</i> , 2018, 16, 3672-3680.	2.8	8
11	Stereochemical Course of Methyl Transfer by Cobalamin-Dependent Radical SAM Methyltransferase in Fosfomycin Biosynthesis. <i>Biochemistry</i> , 2018, 57, 2069-2073.	2.5	13
12	Improved Synthesis of Racemate and Enantiomers of Taniguchi Lactone and Conversion of Their C=C Double Bonds into Triple Bonds. <i>Synthesis</i> , 2018, 50, 651-657.	2.3	6
13	C-Methylation Catalyzed by Fom3, a Cobalamin-Dependent Radical S-adenosyl-methionine Enzyme in Fosfomycin Biosynthesis, Proceeds with Inversion of Configuration. <i>Biochemistry</i> , 2018, 57, 4963-4966.	2.5	24
14	Formal synthesis of P-chiral [¹⁶ O, ¹⁷ O, ¹⁸ O]phosphoenol pyruvates by means of the 1-hydroxyphosphonate-phosphate rearrangement. <i>Phosphorus, Sulfur and Silicon and the Related Elements</i> , 2018, 193, 515-519.	1.6	3
15	Efficient preparation of 2-nitroimidazole nucleosides as precursors for hypoxia PET tracers. <i>Monatshefte für Chemie</i> , 2017, 148, 83-90.	1.8	1
16	Conversion of nitriles to 1-aminophosphonic acids and preparation of phosphahomocysteines of high enantiomeric excess. <i>Phosphorus, Sulfur and Silicon and the Related Elements</i> , 2017, 192, 737-744.	1.6	8
17	Preparation of Nonradioactive Standards and a Precursor for a Hypoxia 18F PET Tracer Derived from 1-(1 ² -d-Galactopyranosyl)-2-nitroimidazole. <i>Synthesis</i> , 2017, 49, 2933-2938.	2.3	1
18	A Methylidene Group in the Phosphonic Acid Analogue of Phenylalanine Reverses the Enantiopreference of Binding to Phenylalanine Ammonia-Lyases. <i>Advanced Synthesis and Catalysis</i> , 2017, 359, 2109-2120.	4.3	9

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19	Chemical Synthesis of (<i>R</i>)- and (<i>S</i>)-[¹⁶ O, ¹⁷ O, ¹⁸ O]Phosphoenol Pyruvate. <i>Journal of Organic Chemistry</i> , 2017, 82, 10310-10318.	3.2	7
20	Chemoenzymatic Synthesis of Racemic and Enantiomerically Pure Phosphoaspartic Acid and Phosphoarginine. <i>European Journal of Organic Chemistry</i> , 2017, 2017, 4836-4845.	2.4	6
21	Phosphonodifluoropyruvate is a mechanism-based inhibitor of phosphonopyruvate decarboxylase from <i>Bacteroides fragilis</i> . <i>Bioorganic and Medicinal Chemistry</i> , 2017, 25, 4368-4374.	3.0	9
22	Synthesis and preclinical characterization of 1-(6-deoxy-6-[¹⁸ F]fluoro- β -D)-Tj ETQq0 0 0 rgBT /Overlock 10 Tf 50 627 Td (-allofu assess tumor hypoxia. <i>Bioorganic and Medicinal Chemistry</i> , 2016, 24, 5326-5339.	3.0	13
23	[¹⁸ F]Fluoro-azomycin-2'-deoxy- β -D-ribofuranoside "A new imaging agent for tumor hypoxia in comparison with [¹⁸ F]FAZA. <i>Nuclear Medicine and Biology</i> , 2016, 43, 759-769.	0.6	4
24	The Stereochemical Course of the β -Hydroxyphosphonate "Phosphate Rearrangement. <i>Chemistry - A European Journal</i> , 2015, 21, 10200-10206.	3.3	21
25	Phosphonate "Phosphinate Rearrangement. <i>Journal of Organic Chemistry</i> , 2015, 80, 1082-1091.	3.2	9
26	Crystal structure of PhnZ in complex with substrate reveals a di-iron oxygenase mechanism for catabolism of organophosphonates. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2014, 111, 5171-5176.	7.1	43
27	Radiosynthesis of [¹²⁴ I]iodometomidate and Biological Evaluation Using Small-Animal PET. <i>Molecular Imaging and Biology</i> , 2014, 16, 317-321.	2.6	5
28	On the Configurational Stability of Chiral, Nonracemic Fluoro- and Iodo-[D ₁]Methylolithiums. <i>Chemistry - A European Journal</i> , 2014, 20, 4086-4091.	3.3	22
29	On the Configurational Stability of Chiral Heteroatom-Substituted [D ₁]Methylpalladium Complexes as Intermediates of Stille and Suzuki-Miyaura Cross-Coupling Reactions. <i>European Journal of Organic Chemistry</i> , 2013, 2013, 5143-5148.	2.4	19
30	Determination of absolute configuration of the phosphonic acid moiety of fosfazinomycins. <i>Organic and Biomolecular Chemistry</i> , 2013, 11, 7420.	2.8	9
31	[³ H]Metyrapol and 4-[¹³¹ I]iodometomidate Label Overlapping, but Not Identical, Binding Sites on Rat Adrenal Membranes. <i>Molecular Pharmaceutics</i> , 2013, 10, 1119-1130.	4.6	9
32	Improved Synthesis of No-Carrier-Added [¹ I]MIBG and Its Precursor. <i>Synthesis</i> , 2012, 44, 3387-3391.	2.3	5
33	On the Preparation and Determination of Configurational Stability of Chiral Thio- and Bromo-[D ₁]methylolithiums. <i>Journal of Organic Chemistry</i> , 2012, 77, 10021-10034.	3.2	14
34	Rearrangement of lithiated S-alkyl O,O-dialkyl thiophosphates: Scope and stereochemistry of the thiophosphate "mercaptophosphonate rearrangement. <i>Organic and Biomolecular Chemistry</i> , 2011, 9, 5220.	2.8	11
35	Chemoenzymatic Synthesis of Phosphonic Acid Analogues of L-Lysine, L-Proline, L-Ornithine, and L-Pipecolic Acid of 99% ee " Assignment of Absolute Configuration to "L-Proline. <i>European Journal of Organic Chemistry</i> , 2011, 2011, 1870-1879.	2.4	25
36	Studies on the Biodegradation of Fosfomycin: Synthesis of [¹³ C]-Labeled Intermediates, Feeding Experiments with <i>Rhizobium huakuii</i> PMY1, and Isolation of Labeled Amino Acids from Cell Mass by HPLC. <i>Chemistry - A European Journal</i> , 2011, 17, 13341-13348.	3.3	4

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37	Synthesis of bipyridine analogues of metomidate for conjugate formation with the $^{99m}\text{Tc}(\text{I})$ -tricarbonyl complex. <i>Monatshefte für Chemie</i> , 2010, 141, 437-443.	1.8	1
38	Novel formal synthesis of stereospecifically C-6 deuterated d-glucose employing configurationally stable alkoxyethylolithiums. <i>Tetrahedron</i> , 2010, 66, 591-598.	1.9	6
39	Preparation of Enantiopure Chiral Amino- $[\text{D}^1]$ methylolithium Compounds and Determination of Their Micro- and Macroscopic Configurational Stabilities. <i>Chemistry - A European Journal</i> , 2009, 15, 5729-5739.	3.3	21
40	Enantiopure Chiral (2,4,6-Triisopropylbenzoyl)oxy- $[\text{D}^1]$ methylolithium: Configurational Stability, Reactions, and Mechanistic Studies. <i>Journal of Organic Chemistry</i> , 2009, 74, 2380-2388.	3.2	32
41	Studies on the biodegradation of fosfomicin: Growth of <i>Rhizobium huakuii</i> PMY1 on possible intermediates synthesised chemically. <i>Organic and Biomolecular Chemistry</i> , 2009, 7, 1944.	2.8	18
42	Preparation of L^* -Aminobenzylphosphonic Acids with a Stereogenic Quaternary Carbon Atom via Microscopically Configurationally Stable L^* -Aminobenzylolithiums. <i>Chemistry - A European Journal</i> , 2008, 14, 8603-8614.	3.3	33
43	On the conversion of structural analogues of (S)-2-hydroxypropylphosphonic acid to epoxides by the final enzyme of fosfomicin biosynthesis in <i>S. fradiae</i> . <i>Bioorganic and Medicinal Chemistry Letters</i> , 2008, 18, 3056-3059.	2.2	9
44	New Selective Inhibitors of Steroid 11β -Hydroxylation in the Adrenal Cortex. Synthesis and Structure-Activity Relationship of Potent Etomidate Analogues. <i>Journal of Medicinal Chemistry</i> , 2008, 51, 2244-2253.	6.4	45
45	Preparation and Configurational Stability of Chiral Chloro- $[\text{D}^1]$ methylolithiums of 98% Enantiomeric Excess. <i>Journal of the American Chemical Society</i> , 2008, 130, 2329-2335.	13.7	38
46	Preparation of Chiral L^* -Oxy- $[\text{2H}^1]$ methylolithiums of 99% ee and Determination of Their Configurational Stability. <i>Journal of the American Chemical Society</i> , 2007, 129, 914-923.	13.7	40
47	Configurational Stability of Oxymethylolithiums as Intermediates in Intramolecular Rearrangements. <i>Chemistry - A European Journal</i> , 2007, 13, 9582-9588.	3.3	31
48	Chemoenzymatic Synthesis of Stannylated Metomidate as a Precursor for Electrophilic Radiohalogenations - Regioselective Alkylation of Methyl 1H-Imidazole-5-carboxylate.. <i>ChemInform</i> , 2005, 36, no.	0.0	0
49	Chemoenzymatic Synthesis of Stannylated Metomidate as a Precursor for Electrophilic Radiohalogenations ? Regioselective Alkylation of Methyl 1H-Imidazole-5-carboxylate [1]. <i>Monatshefte für Chemie</i> , 2005, 136, 229-239.	1.8	8
50	Structure and reactivity of hydroxypropylphosphonic acid epoxidase in fosfomicin biosynthesis by a cation- and flavin-dependent mechanism. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2005, 102, 14221-14226.	7.1	31
51	Direct Chemical Synthesis of Chiral Methanol of 98% ee and Its Conversion to $[\text{2H}^1, \text{3H}]$ Methyl Tosylate and $[\text{2H}^1, \text{3H-Methyl}]$ Methionine. <i>Journal of the American Chemical Society</i> , 2005, 127, 13934-13940.	13.7	30
52	Synthesis of Chiral, Nonracemic L^* -Sulfanylphosphonates and Derivatives.. <i>ChemInform</i> , 2003, 34, no.	0.0	0
53	Synthesis of chiral, nonracemic L^* -sulfanylphosphonates and derivatives. <i>Tetrahedron: Asymmetry</i> , 2003, 14, 1829-1836.	1.8	32
54	Indirect Evidence for the Biosynthesis of (1S,2S)-1,2-Epoxypropylphosphonic Acid as a Co-Metabolite of Fosfomicin [(1R,2S)-1,2-Epoxypropylphosphonic Acid] by <i>Streptomyces fradiae</i> . <i>European Journal of Organic Chemistry</i> , 2002, 2002, 1139-1142.	2.4	16

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55	On the Transformation of (S)-2-Hydroxypropylphosphonic Acid into Fosfomycin in <i>Streptomyces fradiae</i> —A Unique Method of Epoxide Ring Formation. <i>ChemBioChem</i> , 2002, 3, 829-835.	2.6	18
56	Strong versus weak chiral cation exchangers: Comparative evaluation for enantiomer separation of chiral bases by non-aqueous CEC. <i>Journal of Separation Science</i> , 2002, 25, 1269-1283.	2.5	35
57	THE PHOSPHATE-PHOSPHONATE AND PHOSPHONATE-PHOSPHATE REARRANGEMENTS AND THEIR APPLICATIONS - 7[1]: USE OF <i>t</i> -BUTYL AS PROTECTING GROUP AND SYNTHESIS OF CHIRAL, NONRACEMIC $\hat{\pm}$ -HYDROXYPHOSPHONATES. <i>Phosphorus, Sulfur and Silicon and the Related Elements</i> , 2001, 174, 101-118.	1.6	17
58	Simultaneous separation of the stereoisomers of 1-amino-2-hydroxy and 2-amino-1-hydroxypropane phosphonic acids by stereoselective capillary electrophoresis employing a quinine carbamate type chiral selector. <i>Electrophoresis</i> , 2001, 22, 1182-1187.	2.4	24
59	Metallation of Phosphorylated Aliphatic Alcohols to Configurationally Stable $\hat{\pm}$ -Oxyalkyllithium Compounds $\hat{\sim}$ Use of the Phosphoryl Group as an Activating Group and Electrophile. <i>European Journal of Organic Chemistry</i> , 2000, 2000, 2239-2245.	2.4	24
60	Direct liquid chromatographic enantioseparation of chiral $\hat{\pm}$ - and $\hat{2}$ -aminophosphonic acids employing quinine-derived chiral anion exchangers: determination of enantiomeric excess and verification of absolute configuration. <i>Analytica Chimica Acta</i> , 2000, 404, 169-177.	5.4	57
61	Enzymes in organic chemistry. Part 10: Chemo-enzymatic synthesis of l-phosphoserine and l-phosphisoserine and enantioseparation of amino-hydroxyethylphosphonic acids by non-aqueous capillary electrophoresis with quinine carbamate as chiral ion pair agent. <i>Tetrahedron: Asymmetry</i> , 2000, 11, 2955-2964.	1.8	46
62	Transformation of Arylmethylamines into $\hat{\pm}$ -Aminophosphonic Acids via Metalated Phosphoramidates: $\hat{\sim}$ Rearrangement of Partly Configurationally Stable N-Phosphorylated $\hat{\pm}$ -Aminocarbanions. <i>Journal of Organic Chemistry</i> , 2000, 65, 6121-6131.	3.2	86
63	Chemoenzymatic Synthesis of $\hat{\pm}$ -Aminophosphonic Acids. <i>Phosphorus, Sulfur and Silicon and the Related Elements</i> , 1999, 147, 439-439.	1.6	2
64	Enzymes in organic chemistry. Part 9: Chemo-enzymatic synthesis of phosphonic acid analogues of l-valine, l-leucine, l-isoleucine, l-methionine and l- $\hat{\pm}$ -aminobutyric acid of high enantiomeric excess. <i>Tetrahedron: Asymmetry</i> , 1999, 10, 1709-1721.	1.8	48
65	Phosphate-Phosphonate Rearrangement of Aliphatic Phosphates. <i>Phosphorus, Sulfur and Silicon and the Related Elements</i> , 1999, 147, 377-377.	1.6	2
66	ENZYMES IN ORGANIC CHEMISTRY, 8. ^[1] PROTEASE-CATALYZED KINETIC RESOLUTION OF $\hat{\pm}$ -CHLOROACETOXYPHOSPHONATES IN A BIPHASIC SYSTEM. <i>Phosphorus, Sulfur and Silicon and the Related Elements</i> , 1998, 141, 231-238.	1.6	9
67	ENZYMES IN ORGANIC CHEMISTRY 7. ^[1] EVALUATION OF HOMOCHIRAL <i>t</i> -BUTYL(PHENYL)PHOSPHINOTHIOIC ACID FOR THE DETERMINATION OF ENANTIOMERIC EXCESSES AND ABSOLUTE CONFIGURATIONS OF $\hat{\pm}$ -SUBSTITUTED PHOSPHONATES. <i>Phosphorus, Sulfur and Silicon and the Related Elements</i> , 1998, 140, 79-93.	1.6	25
68	Biodegradation of Phosphonomycin by <i>Rhizobium huakuii</i> PMY1. <i>Applied and Environmental Microbiology</i> , 1998, 64, 356-358.	3.1	58
69	Enzymes in organic chemistry, part 2: Lipase-catalysed hydrolysis of 1-acyloxy-2-arylethylphosphonates and synthesis of phosphonic acid analogues of L-phenylalanine and L-tyrosine. <i>Tetrahedron</i> , 1995, 51, 4933-4946.	1.9	66
70	Determination of absolute configuration of $\hat{\pm}$ -hydroxyphosphonates by ^{31}P NMR spectroscopy of corresponding Mosher esters. <i>Tetrahedron</i> , 1994, 50, 10253-10264.	1.9	57
71	Incorporation of L-[Methyl- $^2\text{H}_3$]methionine and 2-[Hydroxy- ^{18}O]hydroxyethylphosphonic Acid into Fosfomycin in <i>Streptomyces fradiae</i> —An Unusual Methyl Transfer. <i>Angewandte Chemie International Edition in English</i> , 1994, 33, 341-342.	4.4	30
72	Enzymes in organic chemistry, part 1: Enantioselective hydrolysis of $\hat{\pm}$ -(acyloxy)phosphonates by esterolytic enzymes. <i>Tetrahedron: Asymmetry</i> , 1993, 4, 109-120.	1.8	94

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73	Biosynthese von Naturstoffen mit einer P=C-Bindung, IX. Synthese und Einbau von (S)- und (R)-2-Hydroxy-[2-2H1]ethylphosphonsäure in Fosfomycin durch Streptomyces fradiae. Liebigs Annalen Der Chemie, 1992, 1992, 553-557.	0.8	12
74	Biosynthesis of natural products with a P=C bond. Part 8: on the origin of the oxirane oxygen atom of fosfomycin in Streptomyces fradiae. Journal of the Chemical Society Perkin Transactions 1, 1991, , 1993-1996.	0.9	40
75	Biosynthesis of natural products with a phosphorus-carbon bond. 7. Synthesis of [1,1-2H2]-, [2,2-2H2]-, (R)- and (S)-[1-2H1](2-hydroxyethyl)phosphonic acid and (R,S)-[1-2H1](1,2-dihydroxyethyl)phosphonic acid and incorporation studies into fosfomycin in Streptomyces fradiae. Journal of Organic Chemistry, 1991, 56, 2364-2370.	3.2	63
76	Addition von Dialkylphosphiten und Dialkyl(trimethylsilyl)phosphiten an 2-(Benzyloxy)propanal Darstellung aller vier stereoisomeren (1,2-Dihydroxyäthylpropyl)phosphonsäuren aus chiralen Lactaten. Liebigs Annalen Der Chemie, 1991, 1991, 469-475.	0.8	23
77	Biosynthese von Naturstoffen mit einer P=C-Bindung, V. Das Oxiran-Sauerstoff-Atom des Fosfomycins entstammt nicht dem Luft-Sauerstoff. Liebigs Annalen Der Chemie, 1990, 1990, 1055-1061.	0.8	37
78	Absolute Konfiguration der (2-Aminoäthyl)phosphonsäure aus Acanthamoeba castellanii (Neff) - Darstellung der Phosphonsäure-Analoga von (+)- und (-)-Serin. Liebigs Annalen Der Chemie, 1989, 1989, 577-583.	0.8	74
79	Biosynthese von Naturstoffen mit einer P=C-Bindung, I. Einbau von D-Glucose in (2-Aminoethyl)phosphonsäure in Tetrahymena thermophila. Liebigs Annalen Der Chemie, 1988, 1988, 531-535.	0.8	15
80	Biosynthese von Naturstoffen mit einer P=C-Bindung, IV. Synthese der (R)- und (S)- (2-Amino[2-2H1]ethyl)phosphonsäure und Hydroxylierung zu (2-Aminoäthyl)phosphonsäure in Acanthamoeba castellanii (Neff). Liebigs Annalen Der Chemie, 1988, 1988, 961-964.	0.8	8
81	Zur Stereochemie der Phosphat-Phosphonat-Umlagerung. Liebigs Annalen Der Chemie, 1986, 1986, 2053-2064.	0.8	24