

Friedrich-Alexander Ludwig

List of Publications by Year in descending order

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papers

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docs citations

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#	ARTICLE	IF	CITATIONS
1	Role of TRPC6 in kidney damage after acute ischemic kidney injury. <i>Scientific Reports</i> , 2022, 12, 3038.	1.6	7
2	Development and Biological Evaluation of the First Highly Potent and Specific Benzamide-Based Radiotracer [18F]BA3 for Imaging of Histone Deacetylases 1 and 2 in Brain. <i>Pharmaceuticals</i> , 2022, 15, 324.	1.7	0
3	A Pharmacokinetic and Metabolism Study of the TRPC6 Inhibitor SH045 in Mice by LC-MS/MS. <i>International Journal of Molecular Sciences</i> , 2022, 23, 3635.	1.8	0
4	(+)-[18F]Flubatine as a novel $\alpha 4\beta 2$ nicotinic acetylcholine receptor PET ligand—results of the first-in-human brain imaging application in patients with $\beta 2$ -amyloid PET-confirmed Alzheimer's disease and healthy controls. <i>European Journal of Nuclear Medicine and Molecular Imaging</i> , 2021, 48, 731-746.	3.3	10
5	Development of 18F-Labeled Radiotracers for PET Imaging of the Adenosine A2A Receptor: Synthesis, Radiolabeling and Preliminary Biological Evaluation. <i>International Journal of Molecular Sciences</i> , 2021, 22, 2285.	1.8	5
6	Preclinical Evaluation of [18F]FACH in Healthy Mice and Piglets: An 18F-Labeled Ligand for Imaging of Monocarboxylate Transporters with PET. <i>International Journal of Molecular Sciences</i> , 2021, 22, 1645.	1.8	0
7	Improved in vivo PET imaging of the adenosine A2A receptor in the brain using [18F]FLUDA, a deuterated radiotracer with high metabolic stability. <i>European Journal of Nuclear Medicine and Molecular Imaging</i> , 2021, 48, 2727-2736.	3.3	18
8	Validation of an LC-MS/MS Method to Quantify the New TRPC6 Inhibitor SH045 (Larixyl) <i>Tj ETQq0 0 0 rgBT /Overlock 10 Tf 50 467 Td (N</i> <i>Pharmaceuticals</i> , 2021, 14, 259.	1.7	3
9	Design, Radiosynthesis and Preliminary Biological Evaluation in Mice of a Brain-Penetrant 18F-Labeled $\beta 2$ Receptor Ligand. <i>International Journal of Molecular Sciences</i> , 2021, 22, 5447.	1.8	6
10	Development of fluorinated and methoxylated benzothiazole derivatives as highly potent and selective cannabinoid CB2 receptor ligands. <i>Bioorganic Chemistry</i> , 2021, 114, 105191.	2.0	4
11	Development of Novel Analogs of the Monocarboxylate Transporter Ligand FACH and Biological Validation of One Potential Radiotracer for Positron Emission Tomography (PET) Imaging. <i>Molecules</i> , 2020, 25, 2309.	1.7	4
12	Sigma-1 Receptor Positron Emission Tomography: A New Molecular Imaging Approach Using (S)-(α)-[18F]Fluspidine in Glioblastoma. <i>Molecules</i> , 2020, 25, 2170.	1.7	5
13	Preclinical Incorporation Dosimetry of [18F]FACH—A Novel 18F-Labeled MCT1/MCT4 Lactate Transporter Inhibitor for Imaging Cancer Metabolism with PET. <i>Molecules</i> , 2020, 25, 2024.	1.7	3
14	In vitro and in vivo Human Metabolism of (S)-[18F]Fluspidine — A Radioligand for Imaging $\beta 1$ Receptors With Positron Emission Tomography (PET). <i>Frontiers in Pharmacology</i> , 2019, 10, 534.	1.6	9
15	Synthesis and In Vitro Evaluation of 8-Pyridinyl-Substituted Benzo[e]imidazo[2,1-c][1,2,4]triazines as Phosphodiesterase 2A Inhibitors. <i>Molecules</i> , 2019, 24, 2791.	1.7	9
16	Targeting cyclic nucleotide phosphodiesterase 5 (PDE5) in brain: Toward the development of a PET radioligand labeled with fluorine-18. <i>Bioorganic Chemistry</i> , 2019, 86, 346-362.	2.0	14
17	Structure—Affinity Relationships of Fluorinated Spirocyclic $\beta 2$ Receptor Ligands with an Exocyclic Benzylamino Moiety. <i>ChemMedChem</i> , 2019, 14, 1392-1402.	1.6	1
18	Development and radiosynthesis of the first ¹⁸ F-labeled inhibitor of monocarboxylate transporters (MCTs). <i>Journal of Labelled Compounds and Radiopharmaceuticals</i> , 2019, 62, 411-424.	0.5	6

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19	Radiosynthesis and Biological Investigation of a Novel Fluorine-18 Labeled Benzoimidazotriazine-Based Radioligand for the Imaging of Phosphodiesterase 2A with Positron Emission Tomography. <i>Molecules</i> , 2019, 24, 4149.	1.7	5
20	μ -Acryloyllysine Piperazides as Irreversible Inhibitors of Transglutaminase 2: Synthesis, Structure-Activity Relationships, and Pharmacokinetic Profiling. <i>Journal of Medicinal Chemistry</i> , 2018, 61, 4528-4560.	2.9	27
21	Exploring the Metabolism of (+)-[18F]Flubatine In Vitro and In Vivo: LC-MS/MS Aided Identification of Radiometabolites in a Clinical PET Study. <i>Molecules</i> , 2018, 23, 464.	1.7	9
22	Radiosynthesis of (S)-[18F]T1 : The first PET radioligand for molecular imaging of $\alpha 4 \beta 2$ nicotinic acetylcholine receptors. <i>Applied Radiation and Isotopes</i> , 2017, 124, 106-113.	0.7	6
23	LC-MS Supported Studies on the in Vitro Metabolism of both Enantiomers of Flubatine and the in Vivo Metabolism of (+)-[18F]Flubatine: A Positron Emission Tomography Radioligand for Imaging $\alpha 4 \beta 2$ Nicotinic Acetylcholine Receptors. <i>Molecules</i> , 2016, 21, 1200.	1.7	12
24	Development of a New Radiofluorinated Quinoline Analog for PET Imaging of Phosphodiesterase 5 (PDE5) in Brain. <i>Pharmaceuticals</i> , 2016, 9, 22.	1.7	15
25	Comparison of in vitro Silico, Electrochemical, in vitro and in vivo Metabolism of a Homologous Series of (Radio)fluorinated $\alpha 1$ Receptor Ligands Designed for Positron Emission Tomography. <i>ChemMedChem</i> , 2016, 11, 2445-2458.	1.6	12
26	Development of a Novel Nonpeptidic ^{18}F -Labeled Radiotracer for in Vivo Imaging of Oxytocin Receptors with Positron Emission Tomography. <i>Journal of Medicinal Chemistry</i> , 2016, 59, 1800-1817.	2.9	17
27	Asymmetric Synthesis of Spirocyclic 2-Benzopyrans for Positron Emission Tomography of $\alpha 1$ Receptors in the Brain. <i>Pharmaceuticals</i> , 2014, 7, 78-112.	1.7	2
28	Synthesis of Natural Product Precursors by Baeyer-Villiger Oxidation with Cyclohexanone Monooxygenase from <i>Acinetobacter</i> . <i>Synthesis</i> , 2001, 2001, 0947-0951.	1.2	43