Markus Laube

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	CRISPR/Cas9 Mediated Knockout of Cyclooxygenase-2 Gene Inhibits Invasiveness in A2058 Melanoma Cells. Cells, 2022, 11, 749.	1.8	7
2	"Clickable―Albumin Binders for Modulating the Tumor Uptake of Targeted Radiopharmaceuticals. Journal of Medicinal Chemistry, 2022, 65, 710-733.	2.9	13
3	Carboranyl Analogues of Mefenamic Acid and Their Biological Evaluation. ACS Omega, 2022, 7, 24282-24291.	1.6	13
4	Development of an ¹⁸ F-Labeled Irreversible Inhibitor of Transglutaminase 2 as Radiometric Tool for Quantitative Expression Profiling in Cells and Tissues. Journal of Medicinal Chemistry, 2021, 64, 3462-3478.	2.9	16
5	Modulation of Î ³ -Secretase Activity by a Carborane-Based Flurbiprofen Analogue. Molecules, 2021, 26, 2843.	1.7	10
6	T Cell Mediated Conversion of a Non-Anti-La Reactive B Cell to an Autoreactive Anti-La B Cell by Somatic Hypermutation. International Journal of Molecular Sciences, 2021, 22, 1198.	1.8	9
7	Radiolabeled Silicon-Rhodamines as Bimodal PET/SPECT-NIR Imaging Agents. Pharmaceuticals, 2021, 14, 1155.	1.7	4
8	Deuteration <i>versus</i> ethylation – strategies to improve the metabolic fate of an ¹⁸ F-labeled celecoxib derivative. RSC Advances, 2020, 10, 38601-38611.	1.7	6
9	Adjuvant Drug-Assisted Bone Healing: Advances and Challenges in Drug Delivery Approaches. Pharmaceutics, 2020, 12, 428.	2.0	26
10	Carboranyl Derivatives of Rofecoxib with Cytostatic Activity against Human Melanoma and Colon Cancer Cells. Scientific Reports, 2020, 10, 4827.	1.6	15
11	Fluorine-18 Labeling of S100 Proteins for Small Animal Positron Emission Tomography. Methods in Molecular Biology, 2019, 1929, 461-485.	0.4	0
12	Regulation of A375 melanoma cell adhesion and migration by EphB4 and EphrinB2 – insights from co-culture experiments. Journal of Cellular Biotechnology, 2019, 5, 27-42.	0.1	0
13	Carboranyl Analogues of Ketoprofen with Cytostatic Activity against Human Melanoma and Colon Cancer Cell Lines. ACS Omega, 2019, 4, 8824-8833.	1.6	11
14	Synthesis and preliminary radiopharmacological characterisation of an ¹¹ C″abelled azadipeptide nitrile as potential PET tracer for imaging of cysteine cathepsins. Journal of Labelled Compounds and Radiopharmaceuticals, 2019, 62, 448-459.	0.5	9
15	Synthesis and Cyclooxygenase Inhibition of Sulfonamide-Substituted (Dihydro)Pyrrolo[3,2,1-hi]indoles and Their Potential Prodrugs. Molecules, 2019, 24, 3807.	1.7	11
16	Carboranyl Analogues of Celecoxib with Potent Cytostatic Activity against Human Melanoma and Colon Cancer Cell Lines. ChemMedChem, 2019, 14, 315-321.	1.6	20
17	Technetium-99m based small molecule radiopharmaceuticals and radiotracers targeting inflammation and infection. Dalton Transactions, 2017, 46, 14435-14451.	1.6	23
18	"Hydrous 18 F-fluoroethylation―– Leaving off the azeotropic drying. Applied Radiation and Isotopes, 2017, 127, 260-268.	0.7	9

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19	Insights into binding of S100 proteins to scavenger receptors: class B scavenger receptor CD36 binds S100A12 with high affinity. Amino Acids, 2017, 49, 183-191.	1.2	22
20	Development of Antioxidant COX-2 Inhibitors as Radioprotective Agents for Radiation Therapy—A Hypothesis-Driven Review. Antioxidants, 2016, 5, 14.	2.2	56
21	Controlled immobilization of His-tagged proteins for protein-ligand interaction experiments using Ni2+-NTA layer on glass surfaces. Clinical Hemorheology and Microcirculation, 2016, 61, 523-539.	0.9	4
22	Development of a ¹⁸ F″abeled Diarylâ€6ubstituted Dihydropyrrolo[3,2,1â€ <i>hi</i> jindole as Potential Probe for Functional Imaging of Cyclooxygenaseâ€2 with PET. ChemistrySelect, 2016, 1, 5812-5820.	0.7	8
23	Protective effects of 2,3-diaryl-substituted indole-based cyclooxygenase-2 inhibitors on oxidative modification of human low density lipoproteins in vitro. Clinical Hemorheology and Microcirculation, 2016, 61, 615-632.	0.9	4
24	Optical imaging of COX-2: Studies on an autofluorescent 2,3-diaryl-substituted indole-based cyclooxygenase-2 inhibitor. Biochemical and Biophysical Research Communications, 2015, 458, 40-45.	1.0	12
25	Diaryl-Substituted (Dihydro)pyrrolo[3,2,1- <i>hi</i>]indoles, a Class of Potent COX-2 Inhibitors with Tricyclic Core Structure. Journal of Organic Chemistry, 2015, 80, 5611-5624.	1.7	27
26	2-[¹⁸ F]Fluoroethyl tosylate – a versatile tool for building ¹⁸ F-based radiotracers for positron emission tomography. MedChemComm, 2015, 6, 1714-1754.	3.5	37
27	Organotypical vascular model for characterization of radioprotective compounds: Studies on antioxidant 2,3-diaryl-substituted indole-based cyclooxygenase-2 inhibitors. Clinical Hemorheology and Microcirculation, 2014, 58, 281-295.	0.9	7
28	2,3-Diaryl-substituted indole based COX-2 inhibitors as leads for imaging tracer development. RSC Advances, 2014, 4, 38726-38742.	1.7	24
29	Visualization of cyclooxygenase-2 using a 2,3-diarylsubstituted indole-based inhibitor and confocal laser induced cryofluorescence microscopy at 20K in melanoma cells in vitro. Biochemical and Biophysical Research Communications, 2013, 430, 301-306.	1.0	6
30	2â€Carbaboraneâ€3â€phenylâ€1 <i>H</i> â€indoles—Synthesis via McMurry Reaction and Cyclooxygenase (CC Inhibition Activity. ChemMedChem, 2013, 8, 329-335.	DX) _{1.6}	26
31	Radiolabeled COX-2 Inhibitors for Non-Invasive Visualization of COX-2 Expression and Activity $\hat{a} \in \mathbb{C}^{n}$ A Critical Update. Molecules, 2013, 18, 6311-6355.	1.7	65
32	Radiosynthesis of a 18F-labeled 2,3-diarylsubstituted indole via McMurry coupling for functional characterization of cyclooxygenase-2 (COX-2) in vitro and in vivo. Bioorganic and Medicinal Chemistry, 2012, 20, 3410-3421.	1.4	47