Hariprasad Gali

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Nanoparticles for Targeting of Prostate Cancer. Current Pharmaceutical Design, 2020, 26, 5393-5413.	0.9	4
2	Discovery and Development of a Novel mPGES-1/5-LOX Dual Inhibitor LFA-9 for Prevention and Treatment of Chronic Inflammatory Diseases. Journal of Inflammation Research, 2020, Volume 13, 1261-1278.	1.6	7
3	Production of [13N]ammonia from [13C]methanol on a 7.5†MeV cyclotron using 13C(p, n)13N reaction: Detection of myocardial infarction in a mouse model. Applied Radiation and Isotopes, 2019, 150, 19-24.	0.7	1
4	Preliminary Human Radiation Dose Estimates of PET Renal Agents, Para-18F-Fluorohippuric Acid and Ortho-124I-Iodohippuric Acid from Rat Biodistribution Data. Current Radiopharmaceuticals, 2018, 11, 58-63.	0.3	2
5	Chemodrug delivery using integrin-targeted PLGA-Chitosan nanoparticle for lung cancer therapy. Scientific Reports, 2017, 7, 14674.	1.6	88
6	Lack of chemopreventive effects of P2X7R inhibitors against pancreatic cancer. Oncotarget, 2017, 8, 97822-97834.	0.8	16
7	Primary radiation dosimetry of a novel PET radiopharmaceutical Ga-NODAGA-glycine in comparison with Tc-DTPA in renal studies. Hellenic Journal of Nuclear Medicine, 2017, 20, 241-246.	0.2	2
8	Synthesis and in vivo evaluation of ortho-[124I]iodohippurate for PET renography in healthy rats. Applied Radiation and Isotopes, 2016, 115, 251-255.	0.7	3
9	Small-Molecule Inhibition of GCNT3 Disrupts Mucin Biosynthesis and Malignant Cellular Behaviors in Pancreatic Cancer. Cancer Research, 2016, 76, 1965-1974.	0.4	34
10	Facile synthesis of para -[18 F]fluorohippurate via iodonium ylide-mediated radiofluorination for PET renography. Bioorganic and Medicinal Chemistry Letters, 2016, 26, 479-483.	1.0	10
11	Evaluation of [18 F]PFH PET renography to predict future disease progression in a rat model of autosomal dominant polycystic kidney disease. Nuclear Medicine and Biology, 2016, 43, 1-5.	0.3	7
12	Synthesis and <i>in vivo</i> evaluation of galliumâ€68â€labeled glycine and hippurate conjugates for positron emission tomography renography. Journal of Labelled Compounds and Radiopharmaceuticals, 2015, 58, 14-19.	0.5	7
13	Novel Retro-Inverso Peptide Inhibitor Reverses Angiotensin Receptor Autoantibody–Induced Hypertension in the Rabbit. Hypertension, 2015, 65, 793-799.	1.3	26
14	Raloxifene and Antiestrogenic Gonadorelin Inhibits Intestinal Tumorigenesis by Modulating Immune Cells and Decreasing Stem-like Cells. Cancer Prevention Research, 2014, 7, 300-309.	0.7	9
15	Role of (Drug) Transporters in Imaging in Health and Disease. Drug Metabolism and Disposition, 2014, 42, 2007-2015.	1.7	11
16	Evaluation of ^{99m} Tc-Probestin SPECT As a Novel Technique for Noninvasive Imaging of Kidney Aminopeptidase N Expression. Molecular Pharmaceutics, 2014, 11, 2948-2953.	2.3	6
17	Solid phase synthesis and biological evaluation of probestin as an angiogenesis inhibitor. Bioorganic and Medicinal Chemistry Letters, 2013, 23, 3561-3564.	1.0	7
18	Synthesis and Evaluation of Novel Tc-99m Labeled Probestin Conjugates for Imaging APN/CD13 Expression In Vivo. Bioconjugate Chemistry, 2012, 23, 115-124.	1.8	17

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19	Synthesis and biodistribution studies of technetium-99m-labeled aminopeptidase N inhibitor conjugates. Bioorganic and Medicinal Chemistry Letters, 2012, 22, 4567-4570.	1.0	4
20	Single-step radiosynthesis and in vivo evaluation of a novel fluorine-18 labeled hippurate for use as a PET renal agent. Nuclear Medicine and Biology, 2012, 39, 1195-1201.	0.3	14
21	Design and Synthesis of a Bombesin Peptide-Conjugated Tripodal Phosphino Dithioether Ligand Topology for the Stabilization of the <i>fac-</i> [M(CO) ₃] ⁺ Core (M = ^{99) Tj ET}	Qq 1. 9 0.7	84 3 &4 rgBT /0
22	Renogram comparison of p-[18F]fluorohippurate with o-[125I]iodohippurate and [99mTc]MAG3 in normal rats. Nuclear Medicine Communications, 2011, 32, 908-912.	0.5	15
23	Synthesis and In Vivo Evaluation of <i>p</i> - ¹⁸ F-Fluorohippurate as a New Radiopharmaceutical for Assessment of Renal Function by PET. Journal of Nuclear Medicine, 2011, 52, 147-153.	2.8	34
24	Antimicrobial Photodynamic Therapy with Functionalized Fullerenes: Quantitative Structure-activity Relationships. Journal of Nanomedicine & Nanotechnology, 2011, 02, 1-9.	1.1	80
25	Synthesis and in vivo evaluation of Tc-99m-labeled cyclic CisoDGRC peptide conjugates for targeting αvβ3 integrin expression. Bioorganic and Medicinal Chemistry Letters, 2010, 20, 5969-5972.	1.0	13
26	Feasibility Evaluation of Detecting Hydroxymethylphosphine Oxide In Vivo by (31)P-MRS. International Journal of Biomedical Science, 2010, 6, 228-32.	0.5	1
27	Photodynamic therapy with fullerenes. Photochemical and Photobiological Sciences, 2007, 6, 1139-1149.	1.6	259
28	Functionalized fullerenes mediate photodynamic killing of cancer cells: Type I versus Type II photochemical mechanism. Free Radical Biology and Medicine, 2007, 43, 711-719.	1.3	225
29	In vitro and in vivo Evaluation of 111In-labeled E. coli Heat-Stable Enterotoxin Analogs for Specific Targeting of Human Breast Cancers. Breast Cancer Research and Treatment, 2006, 98, 7-15.	1.1	7
30	Cationic Fullerenes Are Effective and Selective Antimicrobial Photosensitizers. Chemistry and Biology, 2005, 12, 1127-1135.	6.2	231
31	In Vitro and in Vivo Comparison of HumanEscherichia coliHeat-Stable Peptide Analogues Incorporating the111In-DOTA Group and Distinct Linker Moieties. Bioconjugate Chemistry, 2004, 15, 872-880.	1.8	16
32	Radiochemical Investigations of99mTcâ^'N3S-X-BBN[7â^'14]NH2:Â An in Vitro/in Vivo Structureâ^'Activity Relationship Study Where X = 0-, 3-, 5-, 8-, and 11-Carbon Tethering Moieties. Bioconjugate Chemistry, 2003, 14, 93-102.	1.8	91
33	Radiochemical investigations of 177Lu-DOTA-8-Aoc-BBN[7-14]NH2: an in vitro/in vivo assessment of the targeting ability of this new radiopharmaceutical for PC-3 human prostate cancer cells. Nuclear Medicine and Biology, 2003, 30, 101-109.	0.3	97
34	Novel series of 1111n-labeled bombesin analogs as potential radiopharmaceuticals for specific targeting of gastrin-releasing peptide receptors expressed on human prostate cancer cells. Journal of Nuclear Medicine, 2003, 44, 823-31.	2.8	105
35	Chemical Synthesis ofEscherichia ColiSThAnalogues by Regioselective Disulfide Bond Formation:Â Biological Evaluation of an111In-DOTAâ [~] Phe19-SThAnalogue for Specific Targeting of Human Colon Cancers. Bioconjugate Chemistry, 2002, 13, 224-231.	1.8	27
36	Synthesis, Characterization, and Labeling with 99mTc/188Re of Peptide Conjugates Containing a Dithia-bisphosphine Chelating Agent. Bioconjugate Chemistry, 2001, 12, 354-363.	1.8	63

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37	In Vivo evaluation of an 111 In-labeled ST-peptide analog for specific-targeting of human colon cancers. Nuclear Medicine and Biology, 2001, 28, 903-909.	0.3	15
38	Facile Ring-Opening Reactions of Phthalimides as a New Strategy to Synthesize Amide-Functionalized Phosphonates, Primary Phosphines, and Bisphosphines. Journal of Organic Chemistry, 2000, 65, 676-680.	1.7	44
39	Unprecedented Selective Aminolysis:Â Aminopropyl Phosphine as a Building Block for a New Family of Air Stable Mono-, Bis-, and Tris-Primary Phosphines. Journal of the American Chemical Society, 2000, 122, 1554-1555.	6.6	47
40	Synthesis of Dithio-Diphosphine (P ₂ S ₂ COOH)-Based Bifunctional Chelating Agent. Its Coupling Reactions with Peptide Analogs and Steroids. Phosphorus, Sulfur and Silicon and the Related Elements, 1999, 147, 87-87.	0.8	0
41	Construction of Water-Soluble Phosphines, New Advances in Aqueous Organometallic Chemistry. Phosphorus, Sulfur and Silicon and the Related Elements, 1999, 144, 461-464.	0.8	4
42	99mTc-Labeling and in Vivo Studies of a Bombesin Analogue with a Novel Water-Soluble Dithiadiphosphine-Based Bifunctional Chelating Agent. Bioconjugate Chemistry, 1999, 10, 254-260.	1.8	100
43	Design and Development of Functionalized Water-Soluble Phosphines:  Catalytic and Biomedical Implications. Accounts of Chemical Research, 1999, 32, 9-17.	7.6	118