Geny M M Groothuis

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Significance of the Vitamin D Receptor on Crosstalk with Nuclear Receptors and Regulation of Enzymes and Transporters. AAPS Journal, 2022, 24, .	2.2	10
2	Ex vivo toxicological evaluation of experimental anticancer gold(i) complexes with lansoprazole-type ligands. Toxicology Research, 2019, 8, 885-895.	0.9	10
3	Differentiation of human-induced pluripotent stem cell under flow conditions to mature hepatocytes for liver tissue engineering. Journal of Tissue Engineering and Regenerative Medicine, 2018, 12, 1273-1284.	1.3	26
4	Development of a mechanistic biokinetic model for hepatic bile acid handling to predict possible cholestatic effects of drugs. European Journal of Pharmaceutical Sciences, 2018, 115, 175-184.	1.9	12
5	Challenges on the road to a multicellular bioartificial liver. Journal of Tissue Engineering and Regenerative Medicine, 2018, 12, e227-e236.	1.3	12
6	Alterations in gene expression in vitamin Dâ€deficiency: Downâ€regulation of liver Cyp7a1 and renal Oat3 in mice. Biopharmaceutics and Drug Disposition, 2018, 39, 99-115.	1.1	11
7	Bioconjugation of Supramolecular Metallacages to Integrin Ligands for Targeted Delivery of Cisplatin. Bioconjugate Chemistry, 2018, 29, 3856-3865.	1.8	41
8	Human and rat precision-cut intestinal slices as ex vivo models to study bile acid uptake by the apical sodium-dependent bile acid transporter. European Journal of Pharmaceutical Sciences, 2018, 121, 65-73.	1.9	7
9	Rat precision-cut liver slices predict drug-induced cholestatic injury. Archives of Toxicology, 2017, 91, 3403-3413.	1.9	14
10	The consequence of regional gradients of P-gp and CYP3A4 for drug-drug interactions by P-gp inhibitors and the P-gp/CYP3A4 interplay in the human intestine ex vivo. Toxicology in Vitro, 2017, 40, 26-33.	1.1	22
11	On the toxicity and transport mechanisms of cisplatin in kidney tissues in comparison to a gold-based cytotoxic agent. Metallomics, 2017, 9, 1786-1795.	1.0	20
12	Pâ€gp activity and inhibition in the different regions of human intestine <i>ex vivo</i> . Biopharmaceutics and Drug Disposition, 2017, 38, 127-138.	1.1	18
13	Judging the value of â€~liver-on-a-chip' devices for prediction of toxicity. Expert Opinion on Drug Metabolism and Toxicology, 2017, 13, 125-128.	1.5	27
14	Validation of precision-cut liver slices to study drug-induced cholestasis: a transcriptomics approach. Archives of Toxicology, 2017, 91, 1401-1412.	1.9	32
15	Maintenance of drug metabolism and transport functions in human precision-cut liver slices during prolonged incubation for 5Âdays. Archives of Toxicology, 2017, 91, 2079-2092.	1.9	33
16	Anticancer Gold <i>N</i> â€Heterocyclic Carbene Complexes: A Comparative inâ€vitro and exâ€vivo Study. ChemMedChem, 2017, 12, 1429-1435.	1.6	52
17	Human precision-cut liver slices as a model to test antifibrotic drugs in the early onset of liver fibrosis. Toxicology in Vitro, 2016, 35, 77-85.	1.1	44
18	Classification of Cholestatic and Necrotic Hepatotoxicants Using Transcriptomics on Human Precision-Cut Liver Slices. Chemical Research in Toxicology, 2016, 29, 342-351.	1.7	21

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19	The Consequence of Drug-Drug Interactions Influencing the Interplay between P-Glycoprotein and Cytochrome P450 3a: An Ex Vivo Study with Rat Precision-Cut Intestinal Slices. Drug Metabolism and Disposition, 2016, 44, 683-691.	1.7	16
20	Translational Modeling in Schizophrenia: Predicting Human Dopamine D2 Receptor Occupancy. Pharmaceutical Research, 2016, 33, 1003-1017.	1.7	14
21	Precision-cut intestinal slices: alternative model for drug transport, metabolism, and toxicology research. Expert Opinion on Drug Metabolism and Toxicology, 2016, 12, 175-190.	1.5	56
22	Rat precision-cut intestinal slices to study P-gp activity and the potency of its inhibitors ex vivo. Toxicology in Vitro, 2015, 29, 1070-1078.	1.1	16
23	Viability, function and morphological integrity of precision-cut liver slices during prolonged incubation: Effects of culture medium. Toxicology in Vitro, 2015, 30, 288-299.	1.1	21
24	Improved Synthesis of <i>N</i> Benzylaminoferrocene-Based Prodrugs and Evaluation of Their Toxicity and Antileukemic Activity. Journal of Medicinal Chemistry, 2015, 58, 2015-2024.	2.9	73
25	Acute toxicity of CCl4 but not of paracetamol induces a transcriptomic signature of fibrosis in precision-cut liver slices. Toxicology in Vitro, 2015, 29, 1012-1020.	1.1	21
26	Consequences of Mrp2 deficiency for diclofenac toxicity in the rat intestine ex vivo. Toxicology in Vitro, 2015, 29, 168-175.	1.1	11
27	Diclofenac toxicity in human intestine ex vivo is not related to the formation of intestinal metabolites. Archives of Toxicology, 2015, 89, 107-119.	1.9	24
28	The Effect of Antifibrotic Drugs in Rat Precision-Cut Fibrotic Liver Slices. PLoS ONE, 2014, 9, e95462.	1.1	46
29	Precision-cut liver slices as a model for the early onset of liver fibrosis to test antifibrotic drugs. Toxicology and Applied Pharmacology, 2014, 274, 328-338.	1.3	65
30	Caffeine-Based Gold(I) <i>N</i> -Heterocyclic Carbenes as Possible Anticancer Agents: Synthesis and Biological Properties. Inorganic Chemistry, 2014, 53, 2296-2303.	1.9	196
31	Proteomic profiling in incubation medium of mouse, rat and human precisionâ€cut liver slices for biomarker detection regarding acute drugâ€induced liver injury. Journal of Applied Toxicology, 2014, 34, 993-1001.	1.4	9
32	Precision cut intestinal slices are an appropriate ex vivo model to study NSAID-induced intestinal toxicity in rats. Toxicology in Vitro, 2014, 28, 1296-1305.	1.1	31
33	Drug-Induced Endoplasmic Reticulum and Oxidative Stress Responses Independently Sensitize Toward TNFI±-Mediated Hepatotoxicity. Toxicological Sciences, 2014, 140, 144-159.	1.4	74
34	Human Precision-Cut Liver Slices as an <i>ex Vivo</i> Model to Study Idiosyncratic Drug-Induced Liver Injury. Chemical Research in Toxicology, 2013, 26, 710-720.	1.7	41
35	Recent advances in 2D and 3D in vitro systems using primary hepatocytes, alternative hepatocyte sources and non-parenchymal liver cells and their use in investigating mechanisms of hepatotoxicity, cell signaling and ADME. Archives of Toxicology, 2013, 87, 1315-1530.	1.9	1,089
36	AMAP, the alleged non-toxic isomer of acetaminophen, is toxic in rat and human liver. Archives of Toxicology, 2013, 87, 155-165.	1.9	46

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37	Evaluation of the intestinal toxicity and transport of xenobiotics utilizing precision-cut slices. Xenobiotica, 2013, 43, 73-83.	0.5	18
38	Nanoparticle Formulation of a Poorly Soluble Cannabinoid Receptor 1 Antagonist Improves Absorption by Rat and Human Intestine. Drug Metabolism and Disposition, 2013, 41, 1557-1565.	1.7	10
39	Bronchoconstriction Induces TGF-β Release and Airway Remodelling in Guinea Pig Lung Slices. PLoS ONE, 2013, 8, e65580.	1.1	58
40	Precision-cut intestinal slices as in vitro tool for studies on drug metabolism. Current Drug Metabolism, 2013, 14, 112-9.	0.7	9
41	Mouse Precision-Cut Liver Slices as an ex Vivo Model To Study Idiosyncratic Drug-Induced Liver Injury. Chemical Research in Toxicology, 2012, 25, 1938-1947.	1.7	19
42	Precision-cut Intestinal Slices as In Vitro Tool for Studies on Drug Metabolism. Current Drug Metabolism, 2012, 14, 112-119.	0.7	11
43	On-line HPLC Analysis System for Metabolism and Inhibition Studies in Precision-Cut Liver Slices. Analytical Chemistry, 2011, 83, 84-91.	3.2	38
44	The role of lithocholic acid in the regulation of bile acid detoxication, synthesis, and transport proteins in rat and human intestine and liver slices. Toxicology in Vitro, 2011, 25, 80-90.	1.1	30
45	Hydrogel embedding of precisionâ€cut liver slices in a microfluidic device improves drug metabolic activity. Biotechnology and Bioengineering, 2011, 108, 1404-1412.	1.7	38
46	Microfluidics Enables Small-Scale Tissue-Based Drug Metabolism Studies with Scarce Human Tissue. Journal of the Association for Laboratory Automation, 2011, 16, 468-476.	2.8	27
47	Reduced Ischemia-Reoxygenation Injury in Rat Intestine After Luminal Preservation With a Tailored Solution. Transplantation, 2010, 90, 622-629.	0.5	20
48	Microfluidic biochip for the perifusion of precisionâ€eut rat liver slices for metabolism and toxicology studies. Biotechnology and Bioengineering, 2010, 105, 184-194.	1.7	118
49	Preparation and incubation of precision-cut liver and intestinal slices for application in drug metabolism and toxicity studies. Nature Protocols, 2010, 5, 1540-1551.	5.5	321
50	Regulation of VDR expression in rat and human intestine and liver – Consequences for CYP3A expression. Toxicology in Vitro, 2010, 24, 822-829.	1.1	23
51	A microfluidic approach for in vitro assessment of interorgan interactions in drug metabolism using intestinal and liver slices. Lab on A Chip, 2010, 10, 2778.	3.1	184
52	Expression and regulation of the bile acid transporter, OST <i>α</i> â€OST <i>β</i> in rat and human intestine and liver. Biopharmaceutics and Drug Disposition, 2009, 30, 241-258.	1.1	34
53	Comparison of effects of VDR versus PXR, FXR and GR ligands on the regulation of CYP3A isozymes in rat and human intestine and liver. European Journal of Pharmaceutical Sciences, 2009, 37, 115-125.	1.9	71
54	Induction of Metabolism and Transport in Human Intestine: Validation of Precision-Cut Slices as a Tool to Study Induction of Drug Metabolism in Human Intestine in Vitro. Drug Metabolism and Disposition, 2008, 36, 604-613.	1.7	80

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55	In Vitro Methods to Study Intestinal Drug Metabolism. Current Drug Metabolism, 2007, 8, 658-675.	0.7	116
56	Precision-cut tissue slices as a tool to predict metabolism of novel drugs. Expert Opinion on Drug Metabolism and Toxicology, 2007, 3, 879-898.	1.5	116
57	Analysis of bile acid-induced regulation of FXR target genes in human liver slices. Liver International, 2007, 27, 137-44.	1.9	45
58	Species differences between mouse, rat, dog, monkey and human CYP-mediated drug metabolism, inhibition and induction. Expert Opinion on Drug Metabolism and Toxicology, 2006, 2, 875-894.	1.5	1,122
59	Innovative Methods to Study Human Intestinal Drug Metabolism in Vitro: Precision-Cut Slices Compared with Ussing Chamber Preparations. Drug Metabolism and Disposition, 2006, 34, 1893-1902.	1.7	86
60	COMPARISON OF MOUSE AND RAT CYTOCHROME P450-MEDIATED METABOLISM IN LIVER AND INTESTINE. Drug Metabolism and Disposition, 2006, 34, 1047-1054.	1.7	91
61	EMPIRICAL VALIDATION OF A RAT IN VITRO ORGAN SLICE MODEL AS A TOOL FOR IN VIVO CLEARANCE PREDICTION. Drug Metabolism and Disposition, 2006, 34, 591-599.	1.7	43
62	A new technique for preparing precision-cut slices from small intestine and colon for drug biotransformation studies. Journal of Pharmacological and Toxicological Methods, 2005, 51, 65-72.	0.3	46
63	Precision-Cut Liver Slices as a New Model to Study Toxicity-Induced Hepatic Stellate Cell Activation in a Physiologic Milieu. Toxicological Sciences, 2005, 85, 632-638.	1.4	85
64	CHARACTERIZATION OF RAT SMALL INTESTINAL AND COLON PRECISION-CUT SLICES AS AN IN VITRO SYSTEM FOR DRUG METABOLISM AND INDUCTION STUDIES. Drug Metabolism and Disposition, 2005, 33, 1613-1620.	1.7	52
65	Rat liver slices as a tool to study LPS-induced inflammatory response in the liver. Journal of Hepatology, 2001, 35, 187-194.	1.8	86
66	Characteristics of the hepatic stellate cell-selective carrier mannose 6-phosphate modified albumin (M6P28-HSA). Liver, 2001, 21, 320-328.	0.1	69
67	The capability of isolated hepatocytes and liver slices of donor livers to predict graft function after liver transplantation. Liver International, 2000, 20, 374-380.	1.9	11
68	Human Liver Slices Express the Same Lidocaine Biotransformation Rate as Isolated Human Hepatocytes. ATLA Alternatives To Laboratory Animals, 1993, 21, 466-469.	0.7	19
69	Influence of albumin on the net sinusoidal efflux of the organic anion dibromosulfophthalein from rat liver. Hepatology, 1992, 15, 302-309.	3.6	12
70	Selective hepatobiliary transport defect for organic anions and neutral steroids in mutant rats with hereditary-conjugated hyperbilirubinemia. Hepatology, 1987, 7, 71-76.	3.6	141