Dongyang Liu

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Characterizing the Physicochemical Properties of Two Weakly Basic Drugs and the Precipitates Obtained from Biorelevant Media. Pharmaceutics, 2022, 14, 330.	4.5	2
2	Reduction effect of oral pravastatin on the acute phase response to intravenous zoledronic acid: protocol for a real-world prospective, placebo-controlled trial. BMJ Open, 2022, 12, e060703.	1.9	1
3	Reply to Wolowich and Kwon. Clinical Infectious Diseases, 2021, 72, 1678-1680.	5.8	4
4	Development of a physiologically based pharmacokinetic (PBPK) population model for Chinese elderly subjects. British Journal of Clinical Pharmacology, 2021, 87, 2711-2722.	2.4	19
5	Mechanistic prediction of first-in-human dose for bispecific CD3/EpCAM T-cell engager antibody M701, using an integrated PK/PD modeling method. European Journal of Pharmaceutical Sciences, 2021, 158, 105584.	4.0	12
6	Physiologically based pharmacokinetic model of renally cleared antibacterial drugs in Chinese renal impairment patients. Biopharmaceutics and Drug Disposition, 2021, 42, 24-34.	1.9	4
7	Population-based meta-analysis of chloroquine: informing chloroquine pharmacokinetics in COVID-19 patients. European Journal of Clinical Pharmacology, 2021, 77, 583-593.	1.9	2
8	Evaluation of the efficacy and safety of hydroxychloroquine in comparison with chloroquine in moderate and severe patients with COVID-19. Science China Life Sciences, 2021, 64, 660-663.	4.9	3
9	Pharmacokinetics analysis based on target-mediated drug distribution for RC18, a novel BLyS/APRIL fusion protein to treat systemic lupus erythematosus and rheumatoid arthritis. European Journal of Pharmaceutical Sciences, 2021, 159, 105704.	4.0	6
10	Development of a Virtual Chinese Pediatric Population Physiological Model Targeting Specific Metabolism and Kidney Elimination Pathways. Frontiers in Pharmacology, 2021, 12, 648697.	3.5	2
11	Comprehensive PBPK model to predict drug interaction potential of Zanubrutinib as a victim or perpetrator. CPT: Pharmacometrics and Systems Pharmacology, 2021, 10, 441-454.	2.5	15
12	Dipeptidyl-Peptidase-IV Inhibitors, Imigliptin and Alogliptin, Improve Beta-Cell Function in Type 2 Diabetes. Frontiers in Endocrinology, 2021, 12, 694390.	3.5	3
13	Safety, tolerability, pharmacokinetics, and pharmacodynamics of the glucokinase activator PB-201 and its effects on the glucose excursion profile in drug-naA ve Chinese patients with type 2 diabetes: a randomised controlled, crossover, single-centre phase 1 trial. EClinicalMedicine, 2021, 42, 101185.	7.1	5
14	Application of LC–MS/MS method for determination of dihydroartemisin in human plasma in a pharmacokinetic study. Bioanalysis, 2020, 12, 1635-1646.	1.5	0
15	Response to Jia and Wang. Clinical Infectious Diseases, 2020, 73, 352-353.	5.8	6
16	Preliminary physiologically based pharmacokinetic modeling of renally cleared drugs in Chinese pregnant women. Biopharmaceutics and Drug Disposition, 2020, 41, 248-267.	1.9	10
17	In Vitro Antiviral Activity and Projection of Optimized Dosing Design of Hydroxychloroquine for the Treatment of Severe Acute Respiratory Syndrome Coronavirus 2 (SARS-CoV-2). Clinical Infectious Diseases, 2020, 71, 732-739.	5.8	2,111
18	Translational prediction of first-in-human pharmacokinetics and pharmacodynamics of janagliflozin, a selective SGLT2 inhibitor, using allometric scaling, dedrick and PK/PD modeling methods. European Journal of Pharmaceutical Sciences, 2020, 147, 105281.	4.0	9

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19	Dose selection of chloroquine phosphate for treatment of COVID-19 based on a physiologically based pharmacokinetic model. Acta Pharmaceutica Sinica B, 2020, 10, 1216-1227.	12.0	40
20	Time-Dependent Distribution of Hydroxychloroquine in Cynomolgus Macaques Using Population Pharmacokinetic Modeling Method. Frontiers in Pharmacology, 2020, 11, 602880.	3.5	4
21	Development of a Physiologically Based Pharmacokinetic Model for Hydroxychloroquine and Its Application in Dose Optimization in Specific COVID-19 Patients. Frontiers in Pharmacology, 2020, 11, 585021.	3.5	6
22	Cytotoxicity Evaluation of Chloroquine and Hydroxychloroquine in Multiple Cell Lines and Tissues by Dynamic Imaging System and Physiologically Based Pharmacokinetic Model. Frontiers in Pharmacology, 2020, 11, 574720.	3.5	16
23	Toward Greater Insights on Applications of Modeling and Simulation in Pregnancy. Current Drug Metabolism, 2020, 21, 722-741.	1.2	0
24	Review on the Clinical Pharmacology of Hydroxychloroquine Sulfate for the Treatment of COVID-19. Current Drug Metabolism, 2020, 21, 427-435.	1.2	1
25	Updates on the Pharmacology of Chloroquine against Coronavirus Disease 2019 (COVID-19): A Perspective on its Use in the General and Geriatric Population. Current Drug Metabolism, 2020, 21, 534-540.	1.2	3
26	Current trends in drug metabolism and pharmacokinetics. Acta Pharmaceutica Sinica B, 2019, 9, 1113-1144.	12.0	147
27	A population pharmacokinetic study to accelerate early phase clinical development for a novel drug, teriflunomide sodium, to treat systemic lupus erythematosus. European Journal of Pharmaceutical Sciences, 2019, 136, 104942.	4.0	7
28	Safety, Pharmacokinetics, and Pharmacogenetics of Single-Dose Teriflunomide Sodium and Leflunomide in Healthy Chinese Subjects. Clinical Drug Investigation, 2019, 39, 643-651.	2.2	9
29	A high-performance liquid chromatography-tandem mass spectrometry method for the determination of lifrafenib, a novel RAF kinase and EGFR inhibitor, in human plasma and urine and its application in clinical pharmacokinetic study. Journal of Pharmaceutical and Biomedical Analysis, 2019, 166, 20-29.	2.8	1
30	Development of a simple HPLC–MS/MS method to simultaneously determine teriflunomide and its metabolite in human plasma and urine: Application to clinical pharmacokinetic study of teriflunomide sodium and leflunomide. Biomedical Chromatography, 2019, 33, e4420.	1.7	6
31	Simultaneous determination of TPN729 and its five metabolites in human plasma and urine by liquid chromatography coupled to tandem mass spectrometry. Journal of Pharmaceutical and Biomedical Analysis, 2018, 151, 91-105.	2.8	1
32	Development of an HPLC–MS/MS method to determine janagliflozin in human plasma and urine: application in clinical study. Bioanalysis, 2018, 10, 1439-1454.	1.5	6
33	Metabolites characterization of a novel DPP-4 inhibitor, imigliptin in humans and rats using ultra-high performance liquid chromatography coupled with synapt high-resolution mass spectrometry. Journal of Pharmaceutical and Biomedical Analysis, 2018, 157, 189-200.	2.8	2
34	A highâ€performance liquid chromatography–tandem mass spectrometry method for simultaneous determination of imigliptin, its five metabolites and alogliptin in human plasma and urine and its application to a multipleâ€dose pharmacokinetic study. Biomedical Chromatography, 2018, 32, e4324.	1.7	7
35	Translational Modeling and Simulation in Supporting Early-Phase Clinical Development of New Drug: A Learn–Research–Confirm Process. Clinical Pharmacokinetics, 2017, 56, 925-939.	3.5	7
36	A single-dose study investigating the pharmacokinetics and pharmacodynamics of edoxaban at 30–90 mg in healthy Chinese volunteers. Xenobiotica, 2017, 47, 592-599.	1.1	3

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37	A unified strategy in selection of the best allometric scaling methods to predict human clearance based on drug disposition pathway. Xenobiotica, 2016, 46, 1105-1111.	1.1	10
38	Simultaneous determination of imigliptin and its three metabolites in human plasma and urine by liquid chromatography coupled to tandem mass spectrometry. Journal of Chromatography B: Analytical Technologies in the Biomedical and Life Sciences, 2015, 1002, 300-312.	2.3	2
39	Clinical pharmacokinetics of Icotinib, an anti-cancer drug: evaluation of dose proportionality, food effect, and tolerability in healthy subjects. Cancer Chemotherapy and Pharmacology, 2014, 73, 721-727.	2.3	17