

Adam S Darwich

List of Publications by Year in descending order

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39
papers

1,121
citations

393982

19
h-index

414034

32
g-index

41
all docs

41
docs citations

41
times ranked

1333
citing authors

#	ARTICLE	IF	CITATIONS
1	Why Has Model-Informed Precision Dosing Not Yet Become Common Clinical Reality? Lessons From the Past and a Roadmap for the Future. <i>Clinical Pharmacology and Therapeutics</i> , 2017, 101, 646-656.	2.3	169
2	Meta-Analysis of the Turnover of Intestinal Epithelia in Preclinical Animal Species and Humans. <i>Drug Metabolism and Disposition</i> , 2014, 42, 2016-2022.	1.7	146
3	Model-Informed Precision Dosing: Background, Requirements, Validation, Implementation, and Forward Trajectory of Individualizing Drug Therapy. <i>Annual Review of Pharmacology and Toxicology</i> , 2021, 61, 225-245.	4.2	74
4	The Pharmacokinetics of the CYP3A Substrate Midazolam in Morbidly Obese Patients Before and One Year After Bariatric Surgery. <i>Pharmaceutical Research</i> , 2015, 32, 3927-3936.	1.7	58
5	IMI "Oral biopharmaceutics tools project" Evaluation of bottom-up PBPK prediction success part 2: An introduction to the simulation exercise and overview of results. <i>European Journal of Pharmaceutical Sciences</i> , 2017, 96, 610-625.	1.9	58
6	A mechanistic pharmacokinetic model to assess modified oral drug bioavailability post bariatric surgery in morbidly obese patients: interplay between CYP3A gut wall metabolism, permeability and dissolution. <i>Journal of Pharmacy and Pharmacology</i> , 2012, 64, 1008-1024.	1.2	47
7	Virtual bioequivalence for achlorhydric subjects: The use of PBPK modelling to assess the formulation-dependent effect of achlorhydria. <i>European Journal of Pharmaceutical Sciences</i> , 2017, 109, 111-120.	1.9	47
8	Trends in oral drug bioavailability following bariatric surgery: examining the variable extent of impact on exposure of different drug classes. <i>British Journal of Clinical Pharmacology</i> , 2012, 74, 774-787.	1.1	45
9	IMI "Oral biopharmaceutics tools project" Evaluation of bottom-up PBPK prediction success part 3: Identifying gaps in system parameters by analysing In Silico performance across different compound classes. <i>European Journal of Pharmaceutical Sciences</i> , 2017, 96, 626-642.	1.9	41
10	IMI "oral biopharmaceutics tools project" evaluation of bottom-up PBPK prediction success part 1: Characterisation of the OrBiTo database of compounds. <i>European Journal of Pharmaceutical Sciences</i> , 2017, 96, 598-609.	1.9	34
11	Semiphysiologically based pharmacokinetic model for midazolam and CYP3A mediated metabolite 1-OH-midazolam in morbidly obese and weight loss surgery patients. <i>CPT: Pharmacometrics and Systems Pharmacology</i> , 2016, 5, 20-30.	1.3	30
12	Deconvolution and IVIVC: Exploring the Role of Rate-Limiting Conditions. <i>AAPS Journal</i> , 2016, 18, 321-332.	2.2	30
13	Analysis of the impact of controlled release formulations on oral drug absorption, gut wall metabolism and relative bioavailability of CYP3A substrates using a physiologically-based pharmacokinetic model. <i>European Journal of Pharmaceutical Sciences</i> , 2015, 67, 32-44.	1.9	29
14	What Does it Take to Make Model-Informed Precision Dosing Common Practice? Report from the 1st Asian Symposium on Precision Dosing. <i>AAPS Journal</i> , 2019, 21, 17.	2.2	29
15	Forecasting oral absorption across biopharmaceutics classification system classes with physiologically based pharmacokinetic models. <i>Journal of Pharmacy and Pharmacology</i> , 2016, 68, 1501-1515.	1.2	28
16	IMI "Oral biopharmaceutics tools project" Evaluation of bottom-up PBPK prediction success part 4: Prediction accuracy and software comparisons with improved data and modelling strategies. <i>European Journal of Pharmaceutics and Biopharmaceutics</i> , 2020, 156, 50-63.	2.0	27
17	Role of pharmacokinetic modeling and simulation in precision dosing of anticancer drugs. <i>Translational Cancer Research</i> , 2017, 6, S1512-S1529.	0.4	26
18	Drug disposition and modelling before and after gastric bypass: immediate and controlled-release metoprolol formulations. <i>British Journal of Clinical Pharmacology</i> , 2015, 80, 1021-1030.	1.1	25

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19	Review article: time to revisit Childâ€Pugh score as the basis for predicting drug clearance in hepatic impairment. <i>Alimentary Pharmacology and Therapeutics</i> , 2021, 54, 388-401.	1.9	25
20	Implications of intercorrelation between hepatic CYP3A4â€CYP2C8 enzymes for the evaluation of drugâ€drug interactions: a case study with repaglinide. <i>British Journal of Clinical Pharmacology</i> , 2018, 84, 972-986.	1.1	19
21	Towards Further Verification of Physiologically-Based Kidney Models: Predictability of the Effects of Urine-Flow and Urine-pH on Renal Clearance. <i>Journal of Pharmacology and Experimental Therapeutics</i> , 2019, 368, 157-168.	1.3	17
22	Accounting for inter-correlation between enzyme abundance: a simulation study to assess implications on global sensitivity analysis within physiologically-based pharmacokinetics. <i>Journal of Pharmacokinetics and Pharmacodynamics</i> , 2019, 46, 137-154.	0.8	16
23	Global Sensitivity Analysis of the Rodgers and Rowland Model for Prediction of Tissue: Plasma Partitioning Coefficients: Assessment of the Key Physiological and Physicochemical Factors That Determine Small-Molecule Tissue Distribution. <i>AAPS Journal</i> , 2020, 22, 41.	2.2	15
24	Simultaneous Assessment In Vitro of Transporter and Metabolic Processes in Hepatic Drug Clearance: Use of a Media Loss Approach. <i>Drug Metabolism and Disposition</i> , 2018, 46, 405-414.	1.7	13
25	Integration of advanced methods and models to study drug absorption and related processes: An UNGAP perspective. <i>European Journal of Pharmaceutical Sciences</i> , 2022, 172, 106100.	1.9	12
26	Variance based global sensitivity analysis of physiologically based pharmacokinetic absorption models for BCS Iâ€IV drugs. <i>Journal of Pharmacokinetics and Pharmacodynamics</i> , 2019, 46, 27-42.	0.8	10
27	Physiologically Based Pharmacokinetic Modeling of Transporter-Mediated Hepatic Disposition of Imaging Biomarker Gadoxetate in Rats. <i>Molecular Pharmaceutics</i> , 2021, 18, 2997-3009.	2.3	10
28	A study of the dosage and duration for levobupivacaine infusion by the caudalâ€epidural route in infants aged 3â€6 months. <i>Paediatric Anaesthesia</i> , 2019, 29, 161-168.	0.6	8
29	Developing Clinically Relevant Dissolution Specifications (CRDSs) for Oral Drug Products: Virtual Webinar Series. <i>Pharmaceutics</i> , 2022, 14, 1010.	2.0	7
30	The Impact of Formulation, Delivery, and Dosing Regimen on the Risk of Drugâ€Drug Interactions. <i>Clinical Pharmacology and Therapeutics</i> , 2019, 105, 1329-1331.	2.3	6
31	The nested enzyme-within-enterocyte (NEWE) turnover model for predicting dynamic drug and disease effects on the gut wall. <i>European Journal of Pharmaceutical Sciences</i> , 2019, 131, 195-207.	1.9	5
32	Physiologically-Based Pharmacokinetics. , 2011, , 361-386.		4
33	USING PAGERANK AND SOCIAL NETWORK ANALYSIS TO SPECIFY MENTAL HEALTH FACTORS. <i>Proceedings of the Design Society</i> , 2021, 1, 3379-3388.	0.5	4
34	Serious Gaming of Logistics Management in Pediatric Emergency Medicine. <i>International Journal of Serious Games</i> , 2020, 7, 47-77.	0.8	4
35	A latent variable approach to account for correlated inputs in global sensitivity analysis. <i>Journal of Pharmacokinetics and Pharmacodynamics</i> , 2021, 48, 671-686.	0.8	2
36	Application of the Nested Enzymeâ€Withinâ€Enterocyte (NEWE) Turnover Model for Predicting the Time Course of Pharmacodynamic Effects. <i>CPT: Pharmacometrics and Systems Pharmacology</i> , 2020, 9, 617-627.	1.3	1

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37	Can We Rationalize Oral Drug Exposure Following Bariatric Surgery to Meet the Pharmacotherapeutic Needs of a Growing Patient Population? Commentary on: "Lithium Toxicity Following Roux-en-Y Gastric Bypass" Bariatric Surgical Patient Care, 2014, 9, 81-83.	0.1	0
38	Ratifying the dosage and duration of levo-bupivacaine infusion by the caudal-epidural route in infants aged three to six months. British Journal of Anaesthesia, 2018, 120, e11-e12.	1.5	0
39	Simulation and Model Validation for Mental Health Factors Using a Multi-Methodology Hybrid Approach. , 2021, , .		0