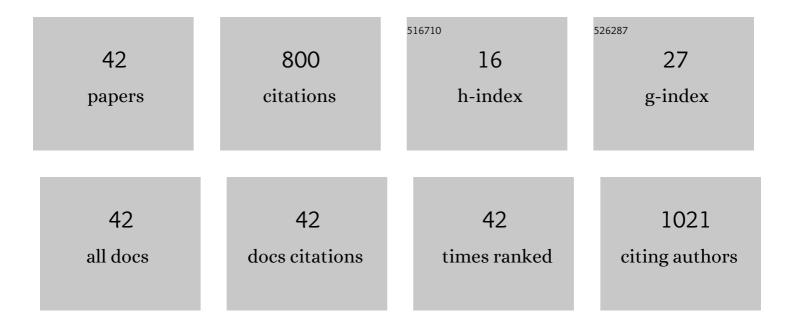
Michel Tod

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Drug interactions between emergency contraceptive drugs and cytochrome inducers: literature review and quantitative prediction. Fundamental and Clinical Pharmacology, 2021, 35, 208-216.	1.9	2
2	Does DDI-Predictor Help Pharmacists to Detect Drug-Drug Interactions and Resolve Medication Issues More Effectively?. Metabolites, 2021, 11, 173.	2.9	8
3	Model-Based Comparative Analysis of Rifampicin and Rifabutin Drug-Drug Interaction Profile. Antimicrobial Agents and Chemotherapy, 2021, 65, e0104321.	3.2	13
4	Quantitative Prediction of Interactions Mediated by Transporters and Cytochromes: Application to Organic Anion Transporting Polypeptides, Breast Cancer Resistance Protein and Cytochrome 2C8. Clinical Pharmacokinetics, 2020, 59, 757-770.	3.5	5
5	Potential drug–drug interactions associated with drugs currently proposed for COVIDâ€19 treatment in patients receiving other treatments. Fundamental and Clinical Pharmacology, 2020, 34, 530-547.	1.9	33
6	A Generic Model for Quantitative Prediction of Interactions Mediated by Efflux Transporters and Cytochromes: Application to P-Glycoprotein and Cytochrome 3A4. Clinical Pharmacokinetics, 2019, 58, 503-523.	3.5	10
7	<i>In Silico</i> Evaluation of Pharmacokinetic Optimization for Antimitogram-Based Clinical Trials. Cancer Research, 2018, 78, 1873-1882.	0.9	0
8	Identification of Cytochrome P450-Mediated Drug–Drug Interactions at Risk in Cases of Gene Polymorphisms by Using a Quantitative Prediction Model. Clinical Pharmacokinetics, 2018, 57, 1581-1591.	3.5	8
9	Semi-Mechanistic Model for Predicting the Dosing Rate in Children and Neonates for Drugs Mainly Eliminated by Cytochrome Metabolism. Clinical Pharmacokinetics, 2018, 57, 831-841.	3.5	4
10	Over-adherence to capecitabine: a potential safety issue in breast and colorectal cancer patients. Cancer Chemotherapy and Pharmacology, 2018, 82, 319-327.	2.3	16
11	Pharmacokinetic interactions in mice between irinotecan and MBLâ€Iâ€I41, an ABCG2 inhibitor. Biopharmaceutics and Drug Disposition, 2017, 38, 351-362.	1.9	2
12	Quantitative Prediction of Drug-Drug Interactions Involving Inhibitory Metabolites by Physiologically Based Pharmacokinetic Models: Is it Worth It?. CPT: Pharmacometrics and Systems Pharmacology, 2017, 6, 226-226.	2.5	2
13	Respiratory depression related to multiple drug–drug interactions precipitated by a fluconazole loading dose in a patient treated with oxycodone. European Journal of Clinical Pharmacology, 2017, 73, 787-788.	1.9	5
14	A Model for Predicting the Interindividual Variability of Drug-Drug Interactions. AAPS Journal, 2017, 19, 497-509.	4.4	7
15	Comparison of the static <i>in vivo</i> approach to a physiologically based pharmacokinetic approach for metabolic drug–drug interactions prediction. International Journal of Pharmacokinetics, 2016, 1, 25-34.	0.5	10
16	Feasibility and Utility of the Individualized Hydrocodone Therapy Based on Phenotype, Pharmacogenetics, and Pharmacokinetic Dosing. Clinical Journal of Pain, 2016, 32, 1106-1107.	1.9	1
17	Response: Is It Truly the Answer? Personalized Oxycodone Dosing Based on Pharmacogenetic Testing and Corresponding Pharmacokinetics. Pain Medicine, 2016, 17, pnv092.	1.9	1
18	Quantitative Prediction of Drug Interactions Caused by CYP1A2 Inhibitors and Inducers. Clinical Pharmacokinetics, 2016, 55, 977-990.	3.5	23

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19	Quantitative Methods for Prediction of the Effect of Cytochrome P450 Gene Polymorphisms on Substrate Drug Exposure. Clinical Pharmacokinetics, 2015, 54, 319-320.	3.5	0
20	A Prediction Model of Drug Exposure in Cirrhotic Patients According to Child–Pugh Classification. Clinical Pharmacokinetics, 2015, 54, 1245-1258.	3.5	14
21	Adherence to oral anticancer chemotherapy: What influences patients' over or non-adherence? Analysis of the OCTO study through quantitative–qualitative methods. BMC Research Notes, 2015, 8, 291.	1.4	20
22	Etoposide pharmacokinetics impact the outcomes of lymphoma patients treated with BEAM regimen and ASCT: a multicenter study of the LYmphoma Study Association (LYSA). Cancer Chemotherapy and Pharmacology, 2015, 76, 939-948.	2.3	7
23	Determination of the Most Influential Sources of Variability in Tacrolimus Trough Blood Concentrations in Adult Liver Transplant Recipients: A Bottom-Up Approach. AAPS Journal, 2014, 16, 379-391.	4.4	37
24	Fractionation of daily dose increases the predicted risk of severe sorafenib-induced hand–foot syndrome (HFS). Cancer Chemotherapy and Pharmacology, 2014, 73, 287-297.	2.3	12
25	Reliability and Extension of Quantitative Prediction of CYP3A4-Mediated Drug Interactions Based on Clinical Data. AAPS Journal, 2014, 16, 1309-1320.	4.4	31
26	Quantitative Prediction of the Impact of Drug Interactions and Genetic Polymorphisms on Cytochrome P450 2C9 Substrate Exposure. Clinical Pharmacokinetics, 2013, 52, 199-209.	3.5	31
27	Impact of Genetic Polymorphism on Drug-Drug Interactions Mediated by Cytochromes: A General Approach. AAPS Journal, 2013, 15, 1242-1252.	4.4	42
28	Theoretical investigation of the efficacy of antiangiogenic drugs combined to chemotherapy in xenografted mice. Journal of Theoretical Biology, 2013, 320, 86-99.	1.7	21
29	In Vivo Quantitative Prediction of the Effect of Gene Polymorphisms and Drug Interactions on Drug Exposure for CYP2C19 Substrates. AAPS Journal, 2013, 15, 415-426.	4.4	39
30	Pharmacodynamic Models for Discrete Data. Clinical Pharmacokinetics, 2012, 51, 767-786.	3.5	6
31	Dose adaptation of capecitabine based on individual prediction of limiting toxicity grade: evaluation by clinical trial simulation. Cancer Chemotherapy and Pharmacology, 2012, 69, 447-455.	2.3	10
32	Quantitative Prediction of Cytochrome P450 (CYP) 2D6-Mediated Drug Interactions. Clinical Pharmacokinetics, 2011, 50, 519-530.	3.5	43
33	Genotype-Based Quantitative Prediction of Drug Exposure for Drugs Metabolized by CYP2D6. Clinical Pharmacology and Therapeutics, 2011, 90, 582-587.	4.7	22
34	Extreme bradycardia due to multiple drug–drug interactions in a patient with HIV postâ€exposure prophylaxis containing lopinavir–ritonavir. British Journal of Clinical Pharmacology, 2011, 71, 621-623.	2.4	23
35	Empirical Bayes estimation of random effects of a mixed-effects proportional odds Markov model for ordinal data. Computer Methods and Programs in Biomedicine, 2011, 104, 505-513.	4.7	2
36	Links Between Cyclosporin Exposure in Tissues and Graft-Versus-Host Disease in Pediatric Bone Marrow Transplantation: Analysis by a PBPK Model. Pharmaceutical Research, 2011, 28, 531-539.	3.5	13

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37	Influence of Dosing Schedule on Organ Exposure to Cyclosporin in Pediatric Hematopoietic Stem Cell Transplantation: Analysis with a PBPK Model. Pharmaceutical Research, 2010, 27, 2602-2613.	3.5	13
38	Etoposide pharmacokinetics and survival in patients with small cell lung cancer: A multicentre study. Lung Cancer, 2008, 62, 261-272.	2.0	37
39	Facilitation of Drug Evaluation in Children byÂPopulation Methods and Modellingâ€. Clinical Pharmacokinetics, 2008, 47, 231-243.	3.5	170
40	Pharmacokinetic-Pharmacodynamic Assessment of Tacrolimus in Liver-Transplant Recipients during the Early Post-Transplantation Period. Therapeutic Drug Monitoring, 2008, 30, 412-418.	2.0	33
41	Pharmacokinetic/Pharmacodynamic and Time-to-Event Models of Ribavirin-Induced Anaemia in Chronic Hepatitis C. Clinical Pharmacokinetics, 2005, 44, 417-428.	3.5	24
42	Quantitative Prediction of Adverse Event Probability Due to Pharmacokinetic Interactions. Drug Safety, 0, , .	3.2	0