

# Michel Tod

## List of Publications by Year in descending order

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42  
papers

800  
citations

516215

16  
h-index

525886

27  
g-index

42  
all docs

42  
docs citations

42  
times ranked

1021  
citing authors

#	ARTICLE	IF	CITATIONS
1	Drug interactions between emergency contraceptive drugs and cytochrome inducers: literature review and quantitative prediction. <i>Fundamental and Clinical Pharmacology</i> , 2021, 35, 208-216.	1.0	2
2	Does DDI-Predictor Help Pharmacists to Detect Drug-Drug Interactions and Resolve Medication Issues More Effectively?. <i>Metabolites</i> , 2021, 11, 173.	1.3	8
3	Model-Based Comparative Analysis of Rifampicin and Rifabutin Drug-Drug Interaction Profile. <i>Antimicrobial Agents and Chemotherapy</i> , 2021, 65, e0104321.	1.4	13
4	Quantitative Prediction of Interactions Mediated by Transporters and Cytochromes: Application to Organic Anion Transporting Polypeptides, Breast Cancer Resistance Protein and Cytochrome 2C8. <i>Clinical Pharmacokinetics</i> , 2020, 59, 757-770.	1.6	5
5	Potential drug-drug interactions associated with drugs currently proposed for COVID-19 treatment in patients receiving other treatments. <i>Fundamental and Clinical Pharmacology</i> , 2020, 34, 530-547.	1.0	33
6	A Generic Model for Quantitative Prediction of Interactions Mediated by Efflux Transporters and Cytochromes: Application to P-Glycoprotein and Cytochrome 3A4. <i>Clinical Pharmacokinetics</i> , 2019, 58, 503-523.	1.6	10
7	<i>In Silico</i> Evaluation of Pharmacokinetic Optimization for Antimitogram-Based Clinical Trials. <i>Cancer Research</i> , 2018, 78, 1873-1882.	0.4	0
8	Identification of Cytochrome P450-Mediated Drug-Drug Interactions at Risk in Cases of Gene Polymorphisms by Using a Quantitative Prediction Model. <i>Clinical Pharmacokinetics</i> , 2018, 57, 1581-1591.	1.6	8
9	Semi-Mechanistic Model for Predicting the Dosing Rate in Children and Neonates for Drugs Mainly Eliminated by Cytochrome Metabolism. <i>Clinical Pharmacokinetics</i> , 2018, 57, 831-841.	1.6	4
10	Over-adherence to capecitabine: a potential safety issue in breast and colorectal cancer patients. <i>Cancer Chemotherapy and Pharmacology</i> , 2018, 82, 319-327.	1.1	16
11	Pharmacokinetic interactions in mice between irinotecan and MBL-141, an ABCG2 inhibitor. <i>Biopharmaceutics and Drug Disposition</i> , 2017, 38, 351-362.	1.1	2
12	Quantitative Prediction of Drug-Drug Interactions Involving Inhibitory Metabolites by Physiologically Based Pharmacokinetic Models: Is it Worth It?. <i>CPT: Pharmacometrics and Systems Pharmacology</i> , 2017, 6, 226-226.	1.3	2
13	Respiratory depression related to multiple drug-drug interactions precipitated by a fluconazole loading dose in a patient treated with oxycodone. <i>European Journal of Clinical Pharmacology</i> , 2017, 73, 787-788.	0.8	5
14	A Model for Predicting the Interindividual Variability of Drug-Drug Interactions. <i>AAPS Journal</i> , 2017, 19, 497-509.	2.2	7
15	Comparison of the static <i>in vivo</i> approach to a physiologically based pharmacokinetic approach for metabolic drug-drug interactions prediction. <i>International Journal of Pharmacokinetics</i> , 2016, 1, 25-34.	0.5	10
16	Feasibility and Utility of the Individualized Hydrocodone Therapy Based on Phenotype, Pharmacogenetics, and Pharmacokinetic Dosing. <i>Clinical Journal of Pain</i> , 2016, 32, 1106-1107.	0.8	1
17	Response: Is It Truly the Answer? Personalized Oxycodone Dosing Based on Pharmacogenetic Testing and Corresponding Pharmacokinetics. <i>Pain Medicine</i> , 2016, 17, pnv092.	0.9	1
18	Quantitative Prediction of Drug Interactions Caused by CYP1A2 Inhibitors and Inducers. <i>Clinical Pharmacokinetics</i> , 2016, 55, 977-990.	1.6	23

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19	Quantitative Methods for Prediction of the Effect of Cytochrome P450 Gene Polymorphisms on Substrate Drug Exposure. <i>Clinical Pharmacokinetics</i> , 2015, 54, 319-320.	1.6	0
20	A Prediction Model of Drug Exposure in Cirrhotic Patients According to Child-Pugh Classification. <i>Clinical Pharmacokinetics</i> , 2015, 54, 1245-1258.	1.6	14
21	Adherence to oral anticancer chemotherapy: What influences patients' over or non-adherence? Analysis of the OCTO study through quantitative and qualitative methods. <i>BMC Research Notes</i> , 2015, 8, 291.	0.6	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). <i>Cancer Chemotherapy and Pharmacology</i> , 2015, 76, 939-948.	1.1	7
23	Determination of the Most Influential Sources of Variability in Tacrolimus Trough Blood Concentrations in Adult Liver Transplant Recipients: A Bottom-Up Approach. <i>AAPS Journal</i> , 2014, 16, 379-391.	2.2	37
24	Fractionation of daily dose increases the predicted risk of severe sorafenib-induced hand-foot syndrome (HFS). <i>Cancer Chemotherapy and Pharmacology</i> , 2014, 73, 287-297.	1.1	12
25	Reliability and Extension of Quantitative Prediction of CYP3A4-Mediated Drug Interactions Based on Clinical Data. <i>AAPS Journal</i> , 2014, 16, 1309-1320.	2.2	31
26	Quantitative Prediction of the Impact of Drug Interactions and Genetic Polymorphisms on Cytochrome P450 2C9 Substrate Exposure. <i>Clinical Pharmacokinetics</i> , 2013, 52, 199-209.	1.6	31
27	Impact of Genetic Polymorphism on Drug-Drug Interactions Mediated by Cytochromes: A General Approach. <i>AAPS Journal</i> , 2013, 15, 1242-1252.	2.2	42
28	Theoretical investigation of the efficacy of antiangiogenic drugs combined to chemotherapy in xenografted mice. <i>Journal of Theoretical Biology</i> , 2013, 320, 86-99.	0.8	21
29	In Vivo Quantitative Prediction of the Effect of Gene Polymorphisms and Drug Interactions on Drug Exposure for CYP2C19 Substrates. <i>AAPS Journal</i> , 2013, 15, 415-426.	2.2	39
30	Pharmacodynamic Models for Discrete Data. <i>Clinical Pharmacokinetics</i> , 2012, 51, 767-786.	1.6	6
31	Dose adaptation of capecitabine based on individual prediction of limiting toxicity grade: evaluation by clinical trial simulation. <i>Cancer Chemotherapy and Pharmacology</i> , 2012, 69, 447-455.	1.1	10
32	Quantitative Prediction of Cytochrome P450 (CYP) 2D6-Mediated Drug Interactions. <i>Clinical Pharmacokinetics</i> , 2011, 50, 519-530.	1.6	43
33	Genotype-Based Quantitative Prediction of Drug Exposure for Drugs Metabolized by CYP2D6. <i>Clinical Pharmacology and Therapeutics</i> , 2011, 90, 582-587.	2.3	22
34	Extreme bradycardia due to multiple drug-drug interactions in a patient with HIV post-exposure prophylaxis containing lopinavir-ritonavir. <i>British Journal of Clinical Pharmacology</i> , 2011, 71, 621-623.	1.1	23
35	Empirical Bayes estimation of random effects of a mixed-effects proportional odds Markov model for ordinal data. <i>Computer Methods and Programs in Biomedicine</i> , 2011, 104, 505-513.	2.6	2
36	Links Between Cyclosporin Exposure in Tissues and Graft-Versus-Host Disease in Pediatric Bone Marrow Transplantation: Analysis by a PBPK Model. <i>Pharmaceutical Research</i> , 2011, 28, 531-539.	1.7	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. <i>Pharmaceutical Research</i> , 2010, 27, 2602-2613.	1.7	13
38	Etoposide pharmacokinetics and survival in patients with small cell lung cancer: A multicentre study. <i>Lung Cancer</i> , 2008, 62, 261-272.	0.9	37
39	Facilitation of Drug Evaluation in Children by Population Methods and Modelling. <i>Clinical Pharmacokinetics</i> , 2008, 47, 231-243.	1.6	170
40	Pharmacokinetic-Pharmacodynamic Assessment of Tacrolimus in Liver-Transplant Recipients during the Early Post-Transplantation Period. <i>Therapeutic Drug Monitoring</i> , 2008, 30, 412-418.	1.0	33
41	Pharmacokinetic/Pharmacodynamic and Time-to-Event Models of Ribavirin-Induced Anaemia in Chronic Hepatitis C. <i>Clinical Pharmacokinetics</i> , 2005, 44, 417-428.	1.6	24
42	Quantitative Prediction of Adverse Event Probability Due to Pharmacokinetic Interactions. <i>Drug Safety</i> , 0, , .	1.4	0