

Barbara WiÅ›niowska

List of Publications by Year in descending order

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Version: 2024-02-01

40
papers

546
citations

623734

14
h-index

677142

22
g-index

51
all docs

51
docs citations

51
times ranked

729
citing authors

#	ARTICLE	IF	CITATIONS
1	Open-access database of literature derived drug-related Torsade de Pointes cases. <i>BMC Pharmacology & Toxicology</i> , 2022, 23, 7.	2.4	3
2	Development and Performance Verification of the PBPK Model for Antazoline and Its Metabolite and Its Utilization for Pharmacological Hypotheses Formulating. <i>Pharmaceuticals</i> , 2022, 15, 379.	3.8	1
3	How circadian variability of the heart rate and plasma electrolytes concentration influence the cardiac electrophysiology – model-based case study. <i>Journal of Pharmacokinetics and Pharmacodynamics</i> , 2021, 48, 387-399.	1.8	1
4	Magnetic Core-Shell Molecularly Imprinted Nano-Conjugates for Extraction of Antazoline and Hydroxyantazoline from Human Plasma – Material Characterization, Theoretical Analysis and Pharmacokinetics. <i>International Journal of Molecular Sciences</i> , 2021, 22, 3665.	4.1	5
5	Development of physiologically based pharmacokinetic model for the immediate release ropinirole tablets. <i>Acta Poloniae Pharmaceutica</i> , 2021, 78, 317-328.	0.1	0
6	Evolutionary Algorithms in Modeling Aerodynamic Properties of Spray-Dried Microparticulate Systems. <i>Applied Sciences (Switzerland)</i> , 2020, 10, 7109.	2.5	0
7	Characterization of In Vitro and In Vivo Metabolism of Antazoline Using Liquid Chromatography-Tandem Mass Spectrometry. <i>International Journal of Molecular Sciences</i> , 2020, 21, 9693.	4.1	2
8	An Open-Access Dataset of Thorough QT Studies Results. <i>Data</i> , 2020, 5, 10.	2.3	4
9	Better prediction of the local concentration-effect relationship: the role of physiologically based pharmacokinetics and quantitative systems pharmacology and toxicology in the evolution of model-informed drug discovery and development. <i>Drug Discovery Today</i> , 2019, 24, 1344-1354.	6.4	17
10	Towards Bridging Translational Gap in Cardiotoxicity Prediction: an Application of Progressive Cardiac Risk Assessment Strategy in TdP Risk Assessment of Moxifloxacin. <i>AAPS Journal</i> , 2018, 20, 47.	4.4	10
11	Drug-physiology interaction and its influence on the QT prolongation-mechanistic modeling study. <i>Journal of Pharmacokinetics and Pharmacodynamics</i> , 2018, 45, 483-490.	1.8	0
12	Real Patient and its Virtual Twin: Application of Quantitative Systems Toxicology Modelling in the Cardiac Safety Assessment of Citalopram. <i>AAPS Journal</i> , 2018, 20, 6.	4.4	23
13	Drug interaction at hERG channel: In vitro assessment of the electrophysiological consequences of drug combinations and comparison against theoretical models. <i>Journal of Applied Toxicology</i> , 2018, 38, 450-458.	2.8	7
14	Effect of multiple drugs interacting with the hERG channel – In vitro study. <i>Journal of Pharmacological and Toxicological Methods</i> , 2018, 93, 118.	0.7	0
15	Quantitative Assessment of the Physiological Parameters Influencing QT Interval Response to Medication: Application of Computational Intelligence Tools. <i>Computational and Mathematical Methods in Medicine</i> , 2018, 2018, 1-11.	1.3	2
16	Virtual Thorough QT (TQT) Trial – Extrapolation of In Vitro Cardiac Safety Data to In Vivo Situation Using Multi-Scale Physiologically Based Ventricular Cell-wall Model Exemplified with Tolterodine and Fesoterodine. <i>AAPS Journal</i> , 2018, 20, 83.	4.4	10
17	Thorough QT (TQT) studies: concordance with torsadogenesis and an evolving cardiac safety testing paradigm. <i>Drug Discovery Today</i> , 2017, 22, 1460-1465.	6.4	4
18	Am I or am I not proarrhythmic? Comparison of various classifications of drug TdP propensity. <i>Drug Discovery Today</i> , 2017, 22, 10-16.	6.4	19

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19	Humans Vary, So Cardiac Models Should Account for That Too!. <i>Frontiers in Physiology</i> , 2017, 8, 700.	2.8	4
20	The Role of Interaction Model in Simulation of Drug Interactions and QT Prolongation. <i>Current Pharmacology Reports</i> , 2016, 2, 339-344.	3.0	13
21	Virtual Clinical Trial Toward Polytherapy Safety Assessment: Combination of Physiologically Based Pharmacokinetic/Pharmacodynamic-Based Modeling and Simulation Approach With Drug-Drug Interactions Involving Terfenadine as an Example. <i>Journal of Pharmaceutical Sciences</i> , 2016, 105, 3415-3424.	3.3	15
22	Top-down, Bottom-up and Middle-out Strategies for Drug Cardiac Safety Assessment via Modeling and Simulations. <i>Current Pharmacology Reports</i> , 2016, 2, 171-177.	3.0	20
23	Drug-drug interactions and QT prolongation as a commonly assessed cardiac effect - comprehensive overview of clinical trials. <i>BMC Pharmacology & Toxicology</i> , 2016, 17, 12.	2.4	54
24	From Heuristic to Mathematical Modeling of Drugs Dissolution Profiles: Application of Artificial Neural Networks and Genetic Programming. <i>Computational and Mathematical Methods in Medicine</i> , 2015, 2015, 1-9.	1.3	17
25	Early Drug Discovery Prediction of Proarrhythmia Potential and Its Covariates. <i>AAPS Journal</i> , 2015, 17, 1025-1032.	4.4	22
26	Enhanced QSAR models for drug-triggered inhibition of the main cardiac ion currents. <i>Journal of Applied Toxicology</i> , 2015, 35, 1030-1039.	2.8	12
27	How in vitro influences in silico utilized for the prediction of in vivo - pilot study of the drug-induced pro-arrhythmic potency prediction. <i>Folia Medica Cracoviensia</i> , 2015, 55, 5-19.	0.3	1
28	In vitro-in vivo extrapolation of drug-induced proarrhythmia predictions at the population level. <i>Drug Discovery Today</i> , 2014, 19, 275-281.	6.4	22
29	Computer-based prediction of the drug proarrhythmic effect: problems, issues, known and suspected challenges. <i>Europace</i> , 2014, 16, 724-735.	1.7	14
30	Slow delayed rectifying potassium current (I_{Kr}) analysis of the in vitro inhibition data and predictive model development. <i>Journal of Applied Toxicology</i> , 2013, 33, 723-739.	2.8	9
31	Virtual population generator for human cardiomyocytes parameters: in silico drug cardiotoxicity assessment. <i>Toxicology Mechanisms and Methods</i> , 2012, 22, 31-40.	2.7	27
32	Tox-Database.net: a curated resource for data describing chemical triggered in vitro cardiac ion channels inhibition. <i>BMC Pharmacology & Toxicology</i> , 2012, 13, 6.	2.4	12
33	Predictive model for L-type channel inhibition: multichannel block in QT prolongation risk assessment. <i>Journal of Applied Toxicology</i> , 2012, 32, 858-866.	2.8	12
34	Evaluation of patients' adherence to statins in Poland. <i>Current Medical Research and Opinion</i> , 2011, 27, 99-105.	1.9	9
35	Prediction of the hERG potassium channel inhibition potential with use of artificial neural networks. <i>Applied Soft Computing Journal</i> , 2011, 11, 2611-2617.	7.2	15
36	BDT-comparator: a program for comparing binary classifiers. <i>Bioinformatics</i> , 2011, 27, 3439-3440.	4.1	22

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37	The open-access dataset for insilico cardiotoxicity prediction system. Bioinformation, 2011, 6, 244-245.	0.5	8
38	Prediction of the hERG Potassium Channel Inhibition Potential with Use of the Artificial Neural Networks. Advances in Intelligent and Soft Computing, 2010, , 91-99.	0.2	1
39	hERG in vitro interchange factorsâ€™ development and verification. Toxicology Mechanisms and Methods, 2009, 19, 278-284.	2.7	22
40	Collation, assessment and analysis of literature <i>in vitro</i> data on hERG receptor blocking potency for subsequent modeling of drugs' cardiotoxic properties. Journal of Applied Toxicology, 2009, 29, 183-206.	2.8	103