Franco Lombardo

List of Publications by Year in descending order

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Version: 2024-02-01

17	10,582 citations	16 h-index	8	18 g-index
papers	Citations	II-IIIQEX		g-macx
18 all docs	18 docs citations	18 times ranked		12392 citing authors

#	Article	IF	CITATIONS
1	In Silico Models of Human PK Parameters. Prediction of Volume of Distribution Using an Extensive Data Set and a Reduced Number of Parameters. Journal of Pharmaceutical Sciences, 2021, 110, 500-509.	1.6	23
2	An Accurate In Vitro Prediction of Human VD _{ss} Based on the Ã~ie–Tozer Equation and Primary Physicochemical Descriptors. 3. Analysis and Assessment of Predictivity on a Large Dataset. Drug Metabolism and Disposition, 2019, 47, 1380-1387.	1.7	13
3	Trend Analysis of a Database of Intravenous Pharmacokinetic Parameters in Humans for 1352 Drug Compounds. Drug Metabolism and Disposition, 2018, 46, 1466-1477.	1.7	91
4	<i>In Silico</i> Absorption, Distribution, Metabolism, Excretion, and Pharmacokinetics (ADME-PK): Utility and Best Practices. An Industry Perspective from the International Consortium for Innovation through Quality in Pharmaceutical Development. Journal of Medicinal Chemistry, 2017, 60, 9097-9113.	2.9	102
5	In Silico Prediction of Volume of Distribution in Humans. Extensive Data Set and the Exploration of Linear and Nonlinear Methods Coupled with Molecular Interaction Fields Descriptors. Journal of Chemical Information and Modeling, 2016, 56, 2042-2052.	2.5	36
6	Clearance Mechanism Assignment and Total Clearance Prediction in Human Based upon in Silico Models. Journal of Medicinal Chemistry, 2014, 57, 4397-4405.	2.9	51
7	Comprehensive Assessment of Human Pharmacokinetic Prediction Based on In Vivo Animal Pharmacokinetic Data, Part 2: Clearance. Journal of Clinical Pharmacology, 2013, 53, 178-191.	1.0	76
8	Comprehensive Assessment of Human Pharmacokinetic Prediction Based on In Vivo Animal Pharmacokinetic Data, Part 1: Volume of Distribution at Steady State. Journal of Clinical Pharmacology, 2013, 53, 167-177.	1.0	60
9	In silico Prediction of Total Human Plasma Clearance. Journal of Chemical Information and Modeling, 2012, 52, 2069-2078.	2.5	49
10	Use of the \tilde{A} ie-Tozer Model in Understanding Mechanisms and Determinants of Drug Distribution. Drug Metabolism and Disposition, 2010, 38, 1159-1165.	1.7	26
11	In Silico Prediction of Volume of Distribution in Human Using Linear and Nonlinear Models on a 669 Compound Data Set. Journal of Medicinal Chemistry, 2009, 52, 4488-4495.	2.9	69
12	Trend Analysis of a Database of Intravenous Pharmacokinetic Parameters in Humans for 670 Drug Compounds. Drug Metabolism and Disposition, 2008, 36, 1385-1405.	1.7	345
13	A Hybrid Mixture Discriminant Analysisâ 'Random Forest Computational Model for the Prediction of Volume of Distribution of Drugs in Human. Journal of Medicinal Chemistry, 2006, 49, 2262-2267.	2.9	101
14	Prediction of Human Volume of Distribution Values for Neutral and Basic Drugs. 2. Extended Data Set and Leave-Class-Out Statistics. Journal of Medicinal Chemistry, 2004, 47, 1242-1250.	2.9	161
15	Prediction of Volume of Distribution Values in Humans for Neutral and Basic Drugs Using Physicochemical Measurements and Plasma Protein Binding Data. Journal of Medicinal Chemistry, 2002, 45, 2867-2876.	2.9	194
16	ElogDoct:Â A Tool for Lipophilicity Determination in Drug Discovery. 2. Basic and Neutral Compounds. Journal of Medicinal Chemistry, 2001, 44, 2490-2497.	2.9	302
17	Experimental and computational approaches to estimate solubility and permeability in drug discovery and development settings. Advanced Drug Delivery Reviews, 1997, 23, 3-25.	6.6	8,880