

# Hans Lennerns

## List of Publications by Citations

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178  
papers

12,542  
citations

49  
h-index

110  
g-index

189  
ext. papers

13,535  
ext. citations

4.2  
avg, IF

6.34  
L-index

| #   | Paper  | IF  | Citations |
|-----|--|-----|-----------|
| 178 | A theoretical basis for a biopharmaceutic drug classification: the correlation of in vitro drug product dissolution and in vivo bioavailability. <i>Pharmaceutical Research</i> , <b>1995</b> , 12, 413-20   | 4.5 | 3518      |
| 177 | Molecular properties of WHO essential drugs and provisional biopharmaceutical classification. <i>Molecular Pharmaceutics</i> , <b>2004</b> , 1, 85-96  | 5.6 | 597       |
| 176 | Clinical pharmacokinetics of atorvastatin. <i>Clinical Pharmacokinetics</i> , <b>2003</b> , 42, 1141-60  | 6.2 | 391       |
| 175 | Correlation of human jejunal permeability (in vivo) of drugs with experimentally and theoretically derived parameters. A multivariate data analysis approach. <i>Journal of Medicinal Chemistry</i> , <b>1998</b> , 41, 4939-49  | 8.3 | 380       |
| 174 | Pharmacodynamics and pharmacokinetics of the HMG-CoA reductase inhibitors. Similarities and differences. <i>Clinical Pharmacokinetics</i> , <b>1997</b> , 32, 403-25   | 6.2 | 379       |
| 173 | Comparison between permeability coefficients in rat and human jejunum. <i>Pharmaceutical Research</i> , <b>1996</b> , 13, 1336-42  | 4.5 | 338       |
| 172 | Characterization of fluids from the stomach and proximal jejunum in men and women. <i>Pharmaceutical Research</i> , <b>1997</b> , 14, 497-502  | 4.5 | 308       |
| 171 | Comparison of human duodenum and Caco-2 gene expression profiles for 12,000 gene sequences tags and correlation with permeability of 26 drugs. <i>Pharmaceutical Research</i> , <b>2002</b> , 19, 1400-16  | 4.5 | 305       |
| 170 | Why is it challenging to predict intestinal drug absorption and oral bioavailability in human using rat model. <i>Pharmaceutical Research</i> , <b>2006</b> , 23, 1675-86  | 4.5 | 293       |
| 169 | The effects of food on the dissolution of poorly soluble drugs in human and in model small intestinal fluids. <i>Pharmaceutical Research</i> , <b>2005</b> , 22, 2141-51   | 4.5 | 226       |
| 168 | In vivo methods for drug absorption - comparative physiologies, model selection, correlations with in vitro methods (IVIVC), and applications for formulation/API/excipient characterization including food effects. <i>European Journal of Pharmaceutical Sciences</i> , <b>2014</b> , 57, 99-151 | 5.1 | 196       |
| 167 | Absorption/metabolism of sulforaphane and quercetin, and regulation of phase II enzymes, in human jejunum in vivo. <i>Drug Metabolism and Disposition</i> , <b>2003</b> , 31, 805-13   | 4   | 183       |
| 166 | Regional jejunal perfusion, a new in vivo approach to study oral drug absorption in man. <i>Pharmaceutical Research</i> , <b>1992</b> , 9, 1243-51   | 4.5 | 178       |
| 165 | Optimizing levodopa pharmacokinetics: intestinal infusion versus oral sustained-release tablets. <i>Clinical Neuropharmacology</i> , <b>2003</b> , 26, 156-63  | 1.4 | 172       |
| 164 | Human jejunal effective permeability and its correlation with preclinical drug absorption models. <i>Journal of Pharmacy and Pharmacology</i> , <b>1997</b> , 49, 627-38   | 4.8 | 143       |
| 163 | The use of biopharmaceutic classification of drugs in drug discovery and development: current status and future extension. <i>Journal of Pharmacy and Pharmacology</i> , <b>2005</b> , 57, 273-85  | 4.8 | 134       |
| 162 | Improving glucocorticoid replacement therapy using a novel modified-release hydrocortisone tablet: a pharmacokinetic study. <i>European Journal of Endocrinology</i> , <b>2009</b> , 161, 119-30   | 6.5 | 129       |

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| 161 | In silico predictions of gastrointestinal drug absorption in pharmaceutical product development: application of the mechanistic absorption model GI-Sim. <i>European Journal of Pharmaceutical Sciences</i> , <b>2013</b> , 49, 679-98 | 5.1  | 120 |
| 160 | SPR biosensor studies of the direct interaction between 27 drugs and a liposome surface: correlation with fraction absorbed in humans. <i>Journal of Medicinal Chemistry</i> , <b>2000</b> , 43, 2083-6                                | 8.3  | 118 |
| 159 | Animal data: the contributions of the Ussing Chamber and perfusion systems to predicting human oral drug delivery in vivo. <i>Advanced Drug Delivery Reviews</i> , <b>2007</b> , 59, 1103-20   | 18.5 | 117 |
| 158 | The use of BDDCS in classifying the permeability of marketed drugs. <i>Pharmaceutical Research</i> , <b>2008</b> , 25, 483-8   | 4.5  | 111 |
| 157 | Pulmonary absorption rate and bioavailability of drugs in vivo in rats: structure-absorption relationships and physicochemical profiling of inhaled drugs. <i>Journal of Pharmaceutical Sciences</i> , <b>2003</b> , 92, 1216-33       | 3.9  | 106 |
| 156 | Transport characteristics of fexofenadine in the Caco-2 cell model. <i>Pharmaceutical Research</i> , <b>2004</b> , 21, 1398-404  | 4.5  | 105 |
| 155 | Toward an increased understanding of the barriers to colonic drug absorption in humans: implications for early controlled release candidate assessment. <i>Molecular Pharmaceutics</i> , <b>2009</b> , 6, 60-73                        | 5.6  | 97  |
| 154 | St John's wort decreases the bioavailability of R- and S-verapamil through induction of the first-pass metabolism. <i>Clinical Pharmacology and Therapeutics</i> , <b>2004</b> , 75, 298-309   | 6.1  | 96  |
| 153 | Passive lipoidal diffusion and carrier-mediated cell uptake are both important mechanisms of membrane permeation in drug disposition. <i>Molecular Pharmaceutics</i> , <b>2014</b> , 11, 1727-38                                       | 5.6  | 87  |
| 152 | High-permeability criterion for BCS classification: segmental/pH dependent permeability considerations. <i>Molecular Pharmaceutics</i> , <b>2010</b> , 7, 1827-34  | 5.6  | 84  |
| 151 | Jejunal absorption and metabolism of R/S-verapamil in humans. <i>Pharmaceutical Research</i> , <b>1998</b> , 15, 856-62  | 4.5  | 81  |
| 150 | Does fluid flow across the intestinal mucosa affect quantitative oral drug absorption? Is it time for a reevaluation?. <i>Pharmaceutical Research</i> , <b>1995</b> , 12, 1573-82  | 4.5  | 80  |
| 149 | Experimental estimation of the effective unstirred water layer thickness in the human jejunum, and its importance in oral drug absorption. <i>European Journal of Pharmaceutical Sciences</i> , <b>1995</b> , 3, 247-253               | 5.1  | 79  |
| 148 | Drug absorption from the isolated perfused rat lung--correlations with drug physicochemical properties and epithelial permeability. <i>Journal of Drug Targeting</i> , <b>2003</b> , 11, 61-74   | 5.4  | 77  |
| 147 | Regional intestinal permeability in rats of compounds with different physicochemical properties and transport mechanisms. <i>Journal of Pharmacy and Pharmacology</i> , <b>1997</b> , 49, 687-90                                       | 4.8  | 75  |
| 146 | Modeling gastrointestinal drug absorption requires more in vivo biopharmaceutical data: experience from in vivo dissolution and permeability studies in humans. <i>Current Drug Metabolism</i> , <b>2007</b> , 8, 645-57               | 3.5  | 75  |
| 145 | Jejunal permeability and hepatic extraction of fluvastatin in humans. <i>Clinical Pharmacology and Therapeutics</i> , <b>1996</b> , 60, 493-503  | 6.1  | 75  |
| 144 | Multiple transport mechanisms involved in the intestinal absorption and first-pass extraction of fexofenadine. <i>Clinical Pharmacology and Therapeutics</i> , <b>2003</b> , 74, 423-36  | 6.1  | 71  |

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| 143 | The BCS, BDDCS, and regulatory guidances. <i>Pharmaceutical Research</i> , <b>2011</b> , 28, 1774-8   | 4.5 | 70 |
| 142 | Intestinal Permeability and Drug Absorption: Predictive Experimental, Computational and In Vivo Approaches. <i>Pharmaceutics</i> , <b>2019</b> , 11,  | 6.4 | 66 |
| 141 | Direct In Vivo Human Intestinal Permeability (Peff ) Determined with Different Clinical Perfusion and Intubation Methods. <i>Journal of Pharmaceutical Sciences</i> , <b>2015</b> , 104, 2702-26  | 3.9 | 66 |
| 140 | Regional intestinal drug permeation: biopharmaceutics and drug development. <i>European Journal of Pharmaceutical Sciences</i> , <b>2014</b> , 57, 333-41   | 5.1 | 66 |
| 139 | A clinical single-pass perfusion investigation of the dynamic in vivo secretory response to a dietary meal in human proximal small intestine. <i>Pharmaceutical Research</i> , <b>2006</b> , 23, 742-51                                   | 4.5 | 66 |
| 138 | Dissolution of hydrocortisone in human and simulated intestinal fluids. <i>Pharmaceutical Research</i> , <b>2000</b> , 17, 183-9  | 4.5 | 66 |
| 137 | The fraction dose absorbed, in humans, and high jejunal human permeability relationship. <i>Molecular Pharmaceutics</i> , <b>2012</b> , 9, 1847-51  | 5.6 | 64 |
| 136 | Characterization of jejunal absorption and apical efflux of ropivacaine, lidocaine and bupivacaine in the rat using in situ and in vitro absorption models. <i>European Journal of Pharmaceutical Sciences</i> , <b>2004</b> , 21, 553-60 | 5.1 | 62 |
| 135 | Human in vivo regional intestinal permeability: importance for pharmaceutical drug development. <i>Molecular Pharmaceutics</i> , <b>2014</b> , 11, 12-23  | 5.6 | 60 |
| 134 | The effect of ketoconazole on the in vivo intestinal permeability of fexofenadine using a regional perfusion technique. <i>British Journal of Clinical Pharmacology</i> , <b>2003</b> , 55, 182-90  | 3.8 | 60 |
| 133 | The effect of ketoconazole on the jejunal permeability and CYP3A metabolism of (R/S)-verapamil in humans. <i>British Journal of Clinical Pharmacology</i> , <b>1999</b> , 48, 180-9   | 3.8 | 59 |
| 132 | Chemotherapy and antiangiogenesis--drug-specific, dose-related effects. <i>Acta Oncologica</i> , <b>2003</b> , 42, 294-303  | 3.0 | 54 |
| 131 | Pharmacokinetics of gefitinib in humans: the influence of gastrointestinal factors. <i>International Journal of Pharmaceutics</i> , <b>2007</b> , 341, 134-42   | 6.5 | 50 |
| 130 | Biliary secretion of rosuvastatin and bile acids in humans during the absorption phase. <i>European Journal of Pharmaceutical Sciences</i> , <b>2006</b> , 29, 205-14   | 5.1 | 50 |
| 129 | Human jejunal permeability of two polar drugs: cimetidine and ranitidine. <i>Pharmaceutical Research</i> , <b>2001</b> , 18, 742-4  | 4.5 | 47 |
| 128 | Human in vivo regional intestinal permeability: quantitation using site-specific drug absorption data. <i>Molecular Pharmaceutics</i> , <b>2015</b> , 12, 2026-39   | 5.6 | 46 |
| 127 | The biopharmaceutics risk assessment roadmap for optimizing clinical drug product performance. <i>Journal of Pharmaceutical Sciences</i> , <b>2014</b> , 103, 3377-3397   | 3.9 | 45 |
| 126 | Human intestinal permeability of piroxicam, propranolol, phenylalanine, and PEG 400 determined by jejunal perfusion. <i>Pharmaceutical Research</i> , <b>1997</b> , 14, 1127-32   | 4.5 | 45 |

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| 125 | Regional Intestinal Permeability of Three Model Drugs in Human. <i>Molecular Pharmaceutics</i> , <b>2016</b> , 13, 3013-21  | 5.6 | 45 |
| 124 | Drug metabolism of CYP3A4, CYP2C9 and CYP2D6 substrates in pigs and humans. <i>European Journal of Pharmaceutical Sciences</i> , <b>2011</b> , 43, 89-98  | 5.1 | 44 |
| 123 | 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> , <b>2017</b> , 96, 610-625  | 5.1 | 43 |
| 122 | Simultaneous assessment of lipid classes and bile acids in human intestinal fluid by solid-phase extraction and HPLC methods. <i>Journal of Lipid Research</i> , <b>2007</b> , 48, 242-51   | 6.3 | 43 |
| 121 | The lack of effect of induced net fluid absorption on the in vivo permeability of terbutaline in the human jejunum. <i>Journal of Drug Targeting</i> , <b>1995</b> , 3, 191-200   | 5.4 | 43 |
| 120 | Presentation of a structurally diverse and commercially available drug data set for correlation and benchmarking studies. <i>Journal of Medicinal Chemistry</i> , <b>2006</b> , 49, 6660-71   | 8.3 | 40 |
| 119 | The influence of net water absorption on the permeability of antipyrine and levodopa in the human jejunum. <i>Pharmaceutical Research</i> , <b>1994</b> , 11, 1540-7  | 4.5 | 40 |
| 118 | In Vivo Mechanisms of Intestinal Drug Absorption from Aprepitant Nanoformulations. <i>Molecular Pharmaceutics</i> , <b>2017</b> , 14, 4233-4242   | 5.6 | 36 |
| 117 | In vivo dog intestinal precipitation of mebendazole: a basic BCS class II drug. <i>Molecular Pharmaceutics</i> , <b>2012</b> , 9, 2903-11   | 5.6 | 36 |
| 116 | Ethanol-drug absorption interaction: potential for a significant effect on the plasma pharmacokinetics of ethanol vulnerable formulations. <i>Molecular Pharmaceutics</i> , <b>2009</b> , 6, 1429-40  | 5.6 | 36 |
| 115 | The multiple depletion curves method provides accurate estimates of intrinsic clearance (CL <sub>int</sub> ), maximum velocity of the metabolic reaction (V <sub>max</sub> ), and Michaelis constant (K <sub>m</sub> ): accuracy and robustness evaluated through experimental data and Monte Carlo simulations. <i>Drug Metabolism and Disposition</i> , <b>2009</b> , 37, 47-58 | 4   | 35 |
| 114 | Regional rectal perfusion: a new in vivo approach to study rectal drug absorption in man. <i>Pharmaceutical Research</i> , <b>1995</b> , 12, 426-32   | 4.5 | 35 |
| 113 | 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> , <b>2017</b> , 96, 626-642  | 5.1 | 34 |
| 112 | A residence-time distribution analysis of the hydrodynamics within the intestine in man during a regional single-pass perfusion with Loc-I-Gut: in-vivo permeability estimation. <i>Journal of Pharmacy and Pharmacology</i> , <b>1997</b> , 49, 682-6  | 4.8 | 34 |
| 111 | First-pass effects of verapamil on the intestinal absorption and liver disposition of fexofenadine in the porcine model. <i>Drug Metabolism and Disposition</i> , <b>2006</b> , 34, 1182-9  | 4   | 34 |
| 110 | Dose-dependent intestinal absorption and significant intestinal excretion (exsorption) of the beta-blocker pafenolol in the rat. <i>Pharmaceutical Research</i> , <b>1993</b> , 10, 727-31  | 4.5 | 34 |
| 109 | Direct estimation of the in vivo dissolution of spironolactone, in two particle size ranges, using the single-pass perfusion technique (Loc-I-Gut) in humans. <i>European Journal of Pharmaceutical Sciences</i> , <b>2001</b> , 12, 239-50   | 5.1 | 33 |
| 108 | Regional gastrointestinal absorption of the beta-blocker pafenolol in the rat and intestinal transit rate determined by movement of <sup>14</sup> C-polyethylene glycol (PEG) 4000. <i>Pharmaceutical Research</i> , <b>1993</b> , 10, 130-5  | 4.5 | 33 |

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| 107 | Miniaturized nebulization catheters: a new approach for delivery of defined aerosol doses to the rat lung. <i>Journal of Aerosol Medicine and Pulmonary Drug Delivery</i> , <b>2002</b> , 15, 283-96   |     | 32 |
| 106 | Regional Intestinal Permeability in Rats: A Comparison of Methods. <i>Molecular Pharmaceutics</i> , <b>2017</b> , 14, 4252-4261  | 5.6 | 30 |
| 105 | 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> , <b>2017</b> , 96, 598-609                                 | 5.1 | 30 |
| 104 | A comparison between direct determination of in vivo dissolution and the deconvolution technique in humans. <i>European Journal of Pharmaceutical Sciences</i> , <b>1999</b> , 8, 19-27  | 5.1 | 30 |
| 103 | In Vitro Release Mechanisms of Doxorubicin From a Clinical Bead Drug-Delivery System. <i>Journal of Pharmaceutical Sciences</i> , <b>2016</b> , 105, 3387-3398   | 3.9 | 30 |
| 102 | The effect of St. John's wort on the pharmacokinetics, metabolism and biliary excretion of finasteride and its metabolites in healthy men. <i>European Journal of Pharmaceutical Sciences</i> , <b>2009</b> , 36, 433-43   | 5.1 | 29 |
| 101 | Water-soluble beta-cyclodextrins in paediatric oral solutions of spironolactone: preclinical evaluation of spironolactone bioavailability from solutions of beta-cyclodextrin derivatives in rats. <i>Journal of Pharmacy and Pharmacology</i> , <b>1998</b> , 50, 611-9 | 4.8 | 28 |
| 100 | Enterohepatic disposition of rosuvastatin in pigs and the impact of concomitant dosing with cyclosporine and gemfibrozil. <i>Drug Metabolism and Disposition</i> , <b>2009</b> , 37, 2349-58   | 4   | 27 |
| 99  | Regional Intestinal Permeability in Dogs: Biopharmaceutical Aspects for Development of Oral Modified-Release Dosage Forms. <i>Molecular Pharmaceutics</i> , <b>2016</b> , 13, 3022-33  | 5.6 | 27 |
| 98  | Formulation predictive dissolution (fPD) testing to advance oral drug product development: An introduction to the US FDA funded Q1st Century BA/BE Qproject. <i>International Journal of Pharmaceutics</i> , <b>2018</b> , 548, 120-127                                  | 6.5 | 27 |
| 97  | Preclinical Effect of Absorption Modifying Excipients on Rat Intestinal Transport of Model Compounds and the Mucosal Barrier Marker Cr-EDTA. <i>Molecular Pharmaceutics</i> , <b>2017</b> , 14, 4243-4251  | 5.6 | 26 |
| 96  | Gastroparesis, metoclopramide, and tardive dyskinesia: Risk revisited. <i>Neurogastroenterology and Motility</i> , <b>2019</b> , 31, e13617  | 4   | 25 |
| 95  | Gastrointestinal metabolism of a vegetable-oil emulsion in healthy subjects. <i>American Journal of Clinical Nutrition</i> , <b>2010</b> , 92, 515-24  | 7   | 25 |
| 94  | Rat jejunal permeability and metabolism of mu-selective tetrapeptides in gastrointestinal fluids from humans and rats. <i>Pharmaceutical Research</i> , <b>1997</b> , 14, 1780-5   | 4.5 | 25 |
| 93  | Regional transport and metabolism of ropivacaine and its CYP3A4 metabolite PPX in human intestine. <i>Journal of Pharmacy and Pharmacology</i> , <b>2003</b> , 55, 963-72  | 4.8 | 24 |
| 92  | Liver Cancer Cell Lines Treated with Doxorubicin under Normoxia and Hypoxia: Cell Viability and Oncologic Protein Profile. <i>Cancers</i> , <b>2019</b> , 11,  | 6.6 | 23 |
| 91  | A new approach for direct in vivo dissolution studies of poorly soluble drugs. <i>Pharmaceutical Research</i> , <b>1997</b> , 14, 1490-2   | 4.5 | 23 |
| 90  | Treatment of intermediate stage hepatocellular carcinoma: a review of intrahepatic doxorubicin drug-delivery systems. <i>Therapeutic Delivery</i> , <b>2014</b> , 5, 447-66  | 3.8 | 22 |

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|----|---|-----|----|
| 89 | The pharmacokinetics and hepatic disposition of repaglinide in pigs: mechanistic modeling of metabolism and transport. <i>Molecular Pharmaceutics</i> , <b>2012</b> , 9, 823-41   | 5.6 | 22 |
| 88 | Pulmonary absorption - estimation of effective pulmonary permeability and tissue retention of ten drugs using an ex vivo rat model and computational analysis. <i>European Journal of Pharmaceutics and Biopharmaceutics</i> , <b>2018</b> , 124, 1-12  | 5.7 | 22 |
| 87 | Concentration- and region-dependent intestinal permeability of fluvastatin in the rat. <i>Journal of Pharmacy and Pharmacology</i> , <b>1998</b> , 50, 737-44   | 4.8 | 21 |
| 86 | Effects of cholesterol and model transmembrane proteins on drug partitioning into lipid bilayers as analysed by immobilized-liposome chromatography. <i>Journal of Pharmacy and Pharmacology</i> , <b>2001</b> , 53, 1477-87  | 4.8 | 21 |
| 85 | Intestinal drug absorption and bioavailability: beyond involvement of single transport function. <i>Journal of Pharmacy and Pharmacology</i> , <b>2003</b> , 55, 429-33   | 4.8 | 21 |
| 84 | In Vivo Drug Delivery Performance of Lipiodol-Based Emulsion or Drug-Eluting Beads in Patients with Hepatocellular Carcinoma. <i>Molecular Pharmaceutics</i> , <b>2017</b> , 14, 448-458  | 5.6 | 20 |
| 83 | Investigation of hepatobiliary disposition of doxorubicin following intrahepatic delivery of different dosage forms. <i>Molecular Pharmaceutics</i> , <b>2014</b> , 11, 131-44  | 5.6 | 20 |
| 82 | Intestinal and hepatobiliary transport of ximelagatran and its metabolites in pigs. <i>Drug Metabolism and Disposition</i> , <b>2008</b> , 36, 1519-28  | 4   | 19 |
| 81 | Enantioselective transport and CYP3A4-mediated metabolism of R/S-verapamil in Caco-2 cell monolayers. <i>European Journal of Pharmaceutical Sciences</i> , <b>2003</b> , 19, 57-65  | 5.1 | 19 |
| 80 | Permeability and clearance views of drug absorption: a commentary. <i>Journal of Pharmacokinetics and Pharmacodynamics</i> , <b>1995</b> , 23, 333-43   |     | 19 |
| 79 | High airway-to-blood transport of an opioid tetrapeptide in the isolated rat lung after aerosol delivery. <i>Peptides</i> , <b>2002</b> , 23, 469-78  | 3.8 | 18 |
| 78 | Simultaneous quantification of the enantiomers of verapamil and its N-demethylated metabolite in human plasma using liquid chromatography-tandem mass spectrometry. <i>Journal of Chromatography B: Analytical Technologies in the Biomedical and Life Sciences</i> , <b>2004</b> , 804, 303-11 | 3.2 | 17 |
| 77 | Pulmonary Dissolution of Poorly Soluble Compounds Studied in an ex Vivo Rat Lung Model. <i>Molecular Pharmaceutics</i> , <b>2019</b> , 16, 3053-3064  | 5.6 | 16 |
| 76 | Evaluation of the use of Classical Nucleation Theory for predicting intestinal crystalline precipitation of two weakly basic BSC class II drugs. <i>European Journal of Pharmaceutical Sciences</i> , <b>2014</b> , 53, 17-27   | 5.1 | 16 |
| 75 | Different effects of ketoconazole on the stereoselective first-pass metabolism of R/S-verapamil in the intestine and the liver: important for the mechanistic understanding of first-pass drug-drug interactions. <i>Drug Metabolism and Disposition</i> , <b>2009</b> , 37, 2186-96            | 4   | 16 |
| 74 | Identification of finasteride metabolites in human bile and urine by high-performance liquid chromatography/tandem mass spectrometry. <i>Drug Metabolism and Disposition</i> , <b>2009</b> , 37, 2008-17  | 4   | 16 |
| 73 | Physiologically Based Pharmacokinetic Model of Itraconazole and Two of Its Metabolites to Improve the Predictions and the Mechanistic Understanding of CYP3A4 Drug-Drug Interactions. <i>Drug Metabolism and Disposition</i> , <b>2018</b> , 46, 1420-1433                                      | 4   | 15 |
| 72 | Pharmacokinetics of an injectable modified-release 2-hydroxyflutamide formulation in the human prostate gland using a semiphysiologically based biopharmaceutical model. <i>Molecular Pharmaceutics</i> , <b>2014</b> , 11, 3097-111  | 5.6 | 15 |

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|----|--|------|----|
| 71 | Effect of a single gemfibrozil dose on the pharmacokinetics of rosuvastatin in bile and plasma in healthy volunteers. <i>Journal of Clinical Pharmacology</i> , <b>2010</b> , 50, 1039-49  | 2.9  | 15 |
| 70 | In vivo investigation in pigs of intestinal absorption, hepatobiliary disposition, and metabolism of the 5 $\alpha$ -reductase inhibitor finasteride and the effects of coadministered ketoconazole. <i>Drug Metabolism and Disposition</i> , <b>2011</b> , 39, 847-57 | 4    | 15 |
| 69 | Replacement therapy of oral hydrocortisone in adrenal insufficiency: the influence of gastrointestinal factors. <i>Expert Opinion on Drug Metabolism and Toxicology</i> , <b>2008</b> , 4, 749-58  | 5.5  | 15 |
| 68 | VOLSURF: A Tool for Drug ADME-Properties Prediction. <i>Methods and Principles in Medicinal Chemistry</i> , <b>2003</b> , 406-419  | 0.4  | 15 |
| 67 | Regional differences in bioavailability of an opioid tetrapeptide in vivo in rats after administration to the respiratory tract. <i>Peptides</i> , <b>2002</b> , 23, 479-88  | 3.8  | 15 |
| 66 | The Critical Role of Passive Permeability in Designing Successful Drugs. <i>ChemMedChem</i> , <b>2020</b> , 15, 1862-1874  | 3.74 | 15 |
| 65 | Translating Human Effective Jejunal Intestinal Permeability to Surface-Dependent Intrinsic Permeability: a Pragmatic Method for a More Mechanistic Prediction of Regional Oral Drug Absorption. <i>AAPS Journal</i> , <b>2015</b> , 17, 1177-92                        | 3.7  | 14 |
| 64 | A Model-Based Approach To Assessing the Importance of Intracellular Binding Sites in Doxorubicin Disposition. <i>Molecular Pharmaceutics</i> , <b>2017</b> , 14, 686-698   | 5.6  | 13 |
| 63 | Combined in vitro-in vivo approach to assess the hepatobiliary disposition of a novel oral thrombin inhibitor. <i>Molecular Pharmaceutics</i> , <b>2013</b> , 10, 4252-62  | 5.6  | 13 |
| 62 | Aqueous Solubility in Discovery, Chemistry, and Assay Changes. <i>Methods and Principles in Medicinal Chemistry</i> , <b>2003</b> , 215-231  | 0.4  | 13 |
| 61 | High-Throughput Measurement of log D and pKa. <i>Methods and Principles in Medicinal Chemistry</i> , <b>2003</b> , 21-45   | 0.4  | 13 |
| 60 | Cell Cultures in Drug Discovery: An Industrial Perspective. <i>Methods and Principles in Medicinal Chemistry</i> , <b>2003</b> , 90-131  | 0.4  | 13 |
| 59 | Effects of a novel combination of orlistat and acarbose on tolerability, appetite, and glucose metabolism in persons with obesity. <i>Obesity Science and Practice</i> , <b>2020</b> , 6, 313-323  | 2.6  | 12 |
| 58 | Effect on the Gastrointestinal Absorption of Drugs from Different Classes in the Biopharmaceutics Classification System, When Treating with Liraglutide. <i>Molecular Pharmaceutics</i> , <b>2015</b> , 12, 4166-73  | 5.6  | 11 |
| 57 | Extensive intestinal glucuronidation of raloxifene in vivo in pigs and impact for oral drug delivery. <i>Xenobiotica</i> , <b>2012</b> , 42, 917-28  | 2    | 11 |
| 56 | Is the jejunal permeability in rats age-dependent?. <i>Pharmaceutical Research</i> , <b>1997</b> , 14, 1278-81   | 4.5  | 11 |
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