Sandra Klein

List of Publications by Year in Descending Order

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

73	2, 009 citations	22	43
papers		h-index	g-index
77	2,287 ext. citations	4.9	5.31
ext. papers		avg, IF	L-index

#	Paper	IF	Citations
73	Vehicles for Drug Administration to Children: Results and Learnings from an In-Depth Screening of FDA-Recommended Liquids and Soft Foods for Product Quality Assessment <i>Pharmaceutical Research</i> , 2022 , 39, 497	4.5	O
72	Dissolution Equipment and Hydrodynamic Considerations for Evaluating Modified-Release Behavior 2022 , 253-271		
71	A systematic approach for assessing the suitability of enteral feeding tubes for the administration of controlled-release pellet formulations. <i>International Journal of Pharmaceutics</i> , 2021 , 612, 121286	6.5	O
70	Review of paediatric gastrointestinal physiology relevant to the absorption of orally administered medicines <i>Advanced Drug Delivery Reviews</i> , 2021 , 181, 114084	18.5	4
69	Prediction of subcutaneous drug absorption - do we have reliable data to design a simulated interstitial fluid?. <i>International Journal of Pharmaceutics</i> , 2021 , 610, 121257	6.5	3
68	A Toolbox for Mimicking Gastrointestinal Conditions in Children: Simulated Paediatric Breakfast Media (SPBM) for Addressing the Variability of Gastric Contents After Typical Paediatric Breakfasts. <i>Journal of Pharmaceutical Sciences</i> , 2021 ,	3.9	1
67	Pediatric formulation development - Challenges of today and strategies for tomorrow: Summary report from M-CERSI workshop 2019. <i>European Journal of Pharmaceutics and Biopharmaceutics</i> , 2021 , 164, 54-65	5.7	3
66	Path towards efficient paediatric formulation development based on partnering with clinical pharmacologists and clinicians, a conect4children expert group white paper. <i>British Journal of Clinical Pharmacology</i> , 2021 ,	3.8	1
65	A Customized Screening Tool Approach for the Development of a Self-Nanoemulsifying Drug Delivery System (SNEDDS) <i>AAPS PharmSciTech</i> , 2021 , 23, 39	3.9	O
64	Age-appropriate solid oral formulations for pediatric applications with a focus on multiparticulates and minitablets: Summary of September 2019 EuPFI workshop. <i>European Journal of Pharmaceutics and Biopharmaceutics</i> , 2020 , 153, 222-225	5.7	6
63	Impact of gastrointestinal physiology on drug absorption in special populationsAn UNGAP review. <i>European Journal of Pharmaceutical Sciences</i> , 2020 , 147, 105280	5.1	63
62	Dissolution testing of modified release products with biorelevant media: An OrBiTo ring study using the USP apparatus III and IV. <i>European Journal of Pharmaceutics and Biopharmaceutics</i> , 2020 , 156, 40-49	5.7	3
61	Safe, swallowable and palatable paediatric mini-tablet formulations for a WHO model list of essential medicines for children compound - A promising starting point for future PUMA applications. <i>European Journal of Pharmaceutics and Biopharmaceutics</i> , 2020 , 156, 11-19	5.7	8
60	A Biopredictive In Vitro Approach for Assessing Compatibility of a Novel Pediatric Hydrocortisone Drug Product within Common Pediatric Dosing Vehicles. <i>Pharmaceutical Research</i> , 2020 , 37, 203	4.5	8
59	A review of GI conditions critical to oral drug absorption in malnourished children. <i>European Journal of Pharmaceutics and Biopharmaceutics</i> , 2019 , 137, 9-22	5.7	7
58	In Vitro Methods for Evaluating Drug Release of Vaginal Ring Formulations-A Critical Review. <i>Pharmaceutics</i> , 2019 , 11,	6.4	8
57	Regulatory Considerations 2019 , 253-284		

Vaginal and Intrauterine Delivery Systems 2019, 177-209 56 О Individualized in vitro and in silico methods for predicting in vivo performance of enteric-coated tablets containing a narrow therapeutic index drug. European Journal of Pharmaceutics and 55 13 5.7 Biopharmaceutics, **2019**, 135, 13-24 A Biopredictive In[Vitro Comparison of Oral Locally Acting Mesalazine Formulations by a Novel Dissolution Model for Assessing Intraluminal Drug Release in Individual Subjects. Journal of 54 3.9 14 Pharmaceutical Sciences, 2018, 107, 1680-1689 Robustness of barrier membrane coated metoprolol tartrate matrix tablets: Drug release evaluation under physiologically relevant in vitro conditions. International Journal of Pharmaceutics, 6.5 6 53 2018, 543, 368-375 Enzymatically Modified Shea Butter and Palm Kernel Oil as Potential Lipid Drug Delivery Matrices. 52 3 3 European Journal of Lipid Science and Technology, 2018, 120, 1700332 Food effects in paediatric medicines development for products Co-administered with food. 6.5 51 19 International Journal of Pharmaceutics, 2018, 536, 530-535 Simulated Genital Tract Fluids and Their Applicability in Drug Release/Dissolution Testing of 16 50 1.7 Vaginal Dosage Forms. Dissolution Technologies, 2018, 25, 40-51 Bioequivalence of locally acting lozenges: Evaluation of critical in vivo parameters and first steps towards a bio-predictive in vitro test method. European Journal of Pharmaceutics and 49 5.7 Biopharmaceutics, 2018, 123, 71-83 Predicting local drug availability of locally acting lozenges: From method design to a linear level A 48 5.7 4 IVIVC. European Journal of Pharmaceutics and Biopharmaceutics, 2018, 133, 269-276 Biorelevant in vitro assessment of dissolution and compatibility properties of a novel paediatric hydrocortisone drug product following exposure of the drug product to child-appropriate 47 5.7 13 administration fluids. European Journal of Pharmaceutics and Biopharmaceutics, 2018, 133, 277-284 Simulating Different Dosing Scenarios for a Child-Appropriate Valproate ER Formulation in a New 46 3.9 16 Pediatric Two-Stage Dissolution Model. AAPS PharmSciTech, 2017, 18, 309-316 A review of patient-specific gastrointestinal parameters as a platform for developing in vitro models for predicting the in vivo performance of oral dosage forms in patients with Parkinsons 6.5 10 45 disease. International Journal of Pharmaceutics, 2017, 533, 298-314 In vitro dissolution testing of parenteral aqueous solutions and oily suspensions of paracetamol 6.5 44 4 and prednisolone. International Journal of Pharmaceutics, 2017, 532, 519-527 Assessing the influence of media composition and ionic strength on drug release from commercial immediate-release and enteric-coated aspirin tablets. Journal of Pharmacy and Pharmacology, 2017, 4.8 43 19 69, 1327-1340 Development and evaluation of accelerated drug release testing methods for a matrix-type 42 13 5.7 intravaginal ring. European Journal of Pharmaceutics and Biopharmaceutics, 2017, 110, 1-12 Development and Validation of a Robust and Efficient HPLC Method for the Simultaneous 41 Quantification of Levodopa, Carbidopa, Benserazide and Entacapone in Complex Matrices. Journal 6 3.4 of Pharmacy and Pharmaceutical Sciences, 2017, 20, 258-269 Dissolution Test Considerations for Oral Multiparticulate Systems. Advances in Delivery Science and 40 O Technology, 2017, 169-212 Towards the development of a paediatric biopharmaceutics classification system: Results of a 6.5 39 13 survey of experts. International Journal of Pharmaceutics, 2016, 511, 1151-7

38	Impact of vibration and agitation speed on dissolution of USP prednisone tablets RS and various IR tablet formulations. <i>AAPS PharmSciTech</i> , 2015 , 16, 759-66	3.9	6
37	Paediatric oral biopharmaceutics: key considerations and current challenges. <i>Advanced Drug Delivery Reviews</i> , 2014 , 73, 102-26	18.5	92
36	A dynamic system for the simulation of fasting luminal pH-gradients using hydrogen carbonate buffers for dissolution testing of ionisable compounds. <i>European Journal of Pharmaceutical Sciences</i> , 2014 , 51, 224-31	5.1	52
35	In vitro models for the prediction of in vivo performance of oral dosage forms. <i>European Journal of Pharmaceutical Sciences</i> , 2014 , 57, 342-66	5.1	240
34	An automated system for monitoring and regulating the pH of bicarbonate buffers. <i>AAPS PharmSciTech</i> , 2013 , 14, 517-22	3.9	34
33	In vitro tools for evaluating novel dosage forms of poorly soluble, weakly basic drugs: case example ketoconazole. <i>Journal of Pharmaceutical Sciences</i> , 2013 , 102, 3645-52	3.9	22
32	Mechanistic understanding of the effect of PPIs and acidic carbonated beverages on the oral absorption of itraconazole based on absorption modeling with appropriate in vitro data. <i>Molecular Pharmaceutics</i> , 2013 , 10, 4016-23	5.6	14
31	Investigating the feasibility of temperature-controlled accelerated drug release testing for an intravaginal ring. <i>European Journal of Pharmaceutics and Biopharmaceutics</i> , 2013 , 85, 966-73	5.7	17
30	The role of individual gastric emptying of pellets in the prediction of diclofenac in vivo dissolution. Journal of Controlled Release, 2013 , 166, 286-93	11.7	15
29	New formulation approaches to improve solubility and drug release from fixed dose combinations: case examples pioglitazone/glimepiride and ezetimibe/simvastatin. <i>European Journal of Pharmaceutics and Biopharmaceutics</i> , 2013 , 84, 208-18	5.7	51
28	Design of biorelevant test setups for the prediction of diclofenac in vivo features after oral administration. <i>Pharmaceutical Research</i> , 2013 , 30, 1483-501	4.5	21
27	Cyclodextrin-water soluble polymer ternary complexes enhance the solubility and dissolution behaviour of poorly soluble drugs. Case example: itraconazole. <i>European Journal of Pharmaceutics and Biopharmaceutics</i> , 2013 , 83, 378-87	5.7	79
26	Application of a ternary HP-ECD-complex approach to improve the dissolution performance of a poorly soluble weak acid under biorelevant conditions. <i>International Journal of Pharmaceutics</i> , 2012 , 430, 176-83	6.5	36
25	Dissolution testing of oral modified-release dosage forms. <i>Journal of Pharmacy and Pharmacology</i> , 2012 , 64, 944-68	4.8	54
24	Miniaturized transfer models to predict the precipitation of poorly soluble weak bases upon entry into the small intestine. <i>AAPS PharmSciTech</i> , 2012 , 13, 1230-5	3.9	17
23	A novel liquefied gas based oral controlled release drug delivery system for liquid drug formulations. <i>European Journal of Pharmaceutics and Biopharmaceutics</i> , 2012 , 81, 334-8	5.7	7
22	An oral-controlled release drug delivery system for liquid and semisolid drug formulations. <i>AAPS PharmSciTech</i> , 2011 , 12, 1183-5	3.9	4
21	Polysaccharides in Oral Drug Delivery Recent Applications and Future Perspectives. <i>ACS Symposium Series</i> , 2010 , 13-30	0.4	8

(2004-2010)

20	A biorelevant dissolution stress test device - background and experiences. <i>Expert Opinion on Drug Delivery</i> , 2010 , 7, 1251-61	8	51
19	Enhanced dissolution of poorly soluble drugs from solid dispersions in carboxymethylcellulose acetate butyrate matrices. <i>ACS Symposium Series</i> , 2010 , 93-113	0.4	4
18	The use of biorelevant dissolution media to forecast the in vivo performance of a drug. <i>AAPS Journal</i> , 2010 , 12, 397-406	3.7	253
17	Improving glyburide solubility and dissolution by complexation with hydroxybutenyl-Ecyclodextrin. <i>Journal of Pharmacy and Pharmacology</i> , 2010 , 61, 23-30	4.8	16
16	Building new drug delivery systems: in vitro and in vivo studies of drug-hydroxybutenyl cyclodextrin complexes. <i>ACS Symposium Series</i> , 2010 , 31-64	0.4	
15	Can Biorelevant Media be Simplified by using SLS and Tween 80 to Replace Bile Compounds?~!2009-10-23~!2010-01-04~!2010-04-29~!. <i>Open Drug Delivery Journal</i> , 2010 , 4, 30-37		15
14	A comparative study of different release apparatus in generating in vitro-in vivo correlations for extended release formulations. <i>European Journal of Pharmaceutics and Biopharmaceutics</i> , 2009 , 73, 115-	-207	53
13	Improving glyburide solubility and dissolution by complexation with hydroxybutenyl-beta-cyclodextrin. <i>Journal of Pharmacy and Pharmacology</i> , 2009 , 61, 23-30	4.8	16
12	Predicting Food Effects on Drug Release from Extended-Release Oral Dosage Forms Containing a Narrow Therapeutic Index Drug. <i>Dissolution Technologies</i> , 2009 , 16, 28-40	1.7	18
11	Use of the BioDis to generate a physiologically relevant IVIVC. <i>Journal of Controlled Release</i> , 2008 , 130, 216-9	11.7	44
10	A standardized mini paddle apparatus as an alternative to the standard paddle. <i>AAPS PharmSciTech</i> , 2008 , 9, 1179-84	3.9	59
9	Pharmacokinetics of itraconazole after intravenous and oral dosing of itraconazole-cyclodextrin formulations. <i>Journal of Pharmaceutical Sciences</i> , 2007 , 96, 3100-16	3.9	42
8	Simplified Biorelevant Media for Screening Dissolution Performance of Poorly Soluble Drugs. <i>Dissolution Technologies</i> , 2007 , 14, 8-13	1.7	34
7	Comparison of Drug Release From Metoprolol Modified Release Dosage Forms in Single Buffer versus a pH-Gradient Dissolution Test. <i>Dissolution Technologies</i> , 2006 , 13, 6-12	1.7	13
6	The Mini Paddle Apparatus Useful Tool in the Early Developmental Stage? Experiences with Immediate-Release Dosage Forms. <i>Dissolution Technologies</i> , 2006 , 13, 6-11	1.7	32
5	Site-specific delivery of anti-inflammatory drugs in the gastrointestinal tract: an in-vitro release model. <i>Journal of Pharmacy and Pharmacology</i> , 2005 , 57, 709-19	4.8	53
4	Development of Dissolution Tests on the Basis of Gastrointestinal Physiology 2005 , 193-227		6
3	Media to simulate the postprandial stomach I. Matching the physicochemical characteristics of standard breakfasts. <i>Journal of Pharmacy and Pharmacology</i> , 2004 , 56, 605-10	4.8	88

2	Drug Release Characteristics of Different Mesalazine Products Using USP Apparatus 3 to Simulate
	Passage Through the GI Tract. <i>Dissolution Technologies</i> , 2002 , 9, 6-12

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A new 5-aminosalicylic acid multi-unit dosage form for the therapy of ulcerative colitis. *European Journal of Pharmaceutics and Biopharmaceutics*, **2001**, 51, 183-90

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