## **Thomas Rades**

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Emerging trends in the stabilization of amorphous drugs. International Journal of Pharmaceutics, 2013, 453, 65-79.	2.6	360
2	Recent advances in co-amorphous drug formulations. Advanced Drug Delivery Reviews, 2016, 100, 116-125.	6.6	350
3	Selection of excipients for melt extrusion with two poorly water-soluble drugs by solubility parameter calculation and thermal analysis. International Journal of Pharmaceutics, 2001, 226, 147-161.	2.6	345
4	Terahertz pulsed spectroscopy and imaging in the pharmaceutical setting - a review. Journal of Pharmacy and Pharmacology, 2010, 59, 209-223.	1.2	330
5	Using Terahertz Pulsed Spectroscopy to Quantify Pharmaceutical Polymorphism and Crystallinity. Journal of Pharmaceutical Sciences, 2005, 94, 837-846.	1.6	326
6	Coamorphous Drug Systems: Enhanced Physical Stability and Dissolution Rate of Indomethacin and Naproxen. Molecular Pharmaceutics, 2011, 8, 1919-1928.	2.3	302
7	New perspectives on lipid and surfactant based drug delivery systems for oral delivery of poorly soluble drugs. Journal of Pharmacy and Pharmacology, 2010, 62, 1622-1636.	1.2	246
8	Amino acids as co-amorphous stabilizers for poorly water soluble drugs – Part 1: Preparation, stability and dissolution enhancement. European Journal of Pharmaceutics and Biopharmaceutics, 2013, 85, 873-881.	2.0	246
9	Enhanced dissolution rate and synchronized release of drugs in binary systems through formulation: Amorphous naproxen–cimetidine mixtures prepared by mechanical activation. Journal of Controlled Release, 2009, 136, 45-53.	4.8	236
10	An overview of recent studies on the analysis of pharmaceutical polymorphs. Journal of Pharmaceutical and Biomedical Analysis, 2011, 55, 618-644.	1.4	233
11	Using terahertz pulsed spectroscopy to study crystallinity of pharmaceutical materials. Chemical Physics Letters, 2004, 390, 20-24.	1.2	217
12	Characterization of glass solutions of poorly water-soluble drugs produced by melt extrusion with hydrophilic amorphous polymers. Journal of Pharmacy and Pharmacology, 2010, 53, 303-315.	1.2	205
13	Co-amorphous simvastatin and glipizide combinations show improved physical stability without evidence of intermolecular interactions. European Journal of Pharmaceutics and Biopharmaceutics, 2012, 81, 159-169.	2.0	197
14	Effects of intraduodenal fatty acids on appetite, antropyloroduodenal motility, and plasma CCK and GLP-1 in humans vary with their chain length. American Journal of Physiology - Regulatory Integrative and Comparative Physiology, 2004, 287, R524-R533.	0.9	196
15	Raman spectroscopy for quantitative analysis of pharmaceutical solidsâ€. Journal of Pharmacy and Pharmacology, 2010, 59, 179-192.	1.2	196
16	Analysis of Coating Structures and Interfaces in Solid Oral Dosage Forms by Three Dimensional Terahertz Pulsed Imaging. Journal of Pharmaceutical Sciences, 2007, 96, 330-340.	1.6	179
17	Physical characterization and stability of amorphous indomethacin and ranitidine hydrochloride binary systems prepared by mechanical activation. European Journal of Pharmaceutics and Biopharmaceutics, 2009, 71, 47-54.	2.0	179
18	Analysis of solid-state transformations of pharmaceutical compounds using vibrational spectroscopy. Journal of Pharmacy and Pharmacology, 2010, 61, 971-988.	1.2	179

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19	In vitro and in vivo performance of novel supersaturated self-nanoemulsifying drug delivery systems (super-SNEDDS). Journal of Controlled Release, 2012, 160, 25-32.	4.8	178
20	Structural investigations on nanoemulsions, solid lipid nanoparticles and nanostructured lipid carriers by cryo-field emission scanning electron microscopy and Raman spectroscopy. International Journal of Pharmaceutics, 2006, 314, 56-62.	2.6	170
21	Preparation of biodegradable insulin nanocapsules from biocompatible microemulsions. Pharmaceutical Research, 2000, 17, 684-689.	1.7	158
22	Liposomal delivery of antigen to human dendritic cells. Vaccine, 2003, 21, 883-890.	1.7	157
23	Understanding the Influence of Polymorphism on Phonon Spectra:Â Lattice Dynamics Calculations and Terahertz Spectroscopy of Carbamazepine. Journal of Physical Chemistry B, 2006, 110, 447-456.	1.2	157
24	Silica-lipid hybrid (SLH) microcapsules: A novel oral delivery system for poorly soluble drugs. Journal of Controlled Release, 2009, 134, 62-70.	4.8	154
25	Amino acids as co-amorphous stabilizers for poorly water-soluble drugs – Part 2: Molecular interactions. European Journal of Pharmaceutics and Biopharmaceutics, 2013, 85, 882-888.	2.0	153
26	Preparation of glass solutions of three poorly water soluble drugs by spray drying, melt extrusion and ball milling. International Journal of Pharmaceutics, 2007, 336, 22-34.	2.6	149
27	Liposome-Based Adjuvants for Subunit Vaccines: Formulation Strategies for Subunit Antigens and Immunostimulators. Pharmaceutics, 2016, 8, 7.	2.0	147
28	Analysis of sustained-release tablet film coats using terahertz pulsed imaging. Journal of Controlled Release, 2007, 119, 253-261.	4.8	145
29	Preparation of phytantriol cubosomes by solvent precursor dilution for the delivery of protein vaccines. European Journal of Pharmaceutics and Biopharmaceutics, 2011, 79, 15-22.	2.0	145
30	Fat digestion modulates gastrointestinal sensations induced by gastric distention and duodenal lipid in humans. Gastroenterology, 2001, 120, 1100-1107.	0.6	142
31	W/O microemulsions for ocular delivery: Evaluation of ocular irritation and precorneal retention. Journal of Controlled Release, 2006, 111, 145-152.	4.8	138
32	Drug hydrate systems and dehydration processes studied by terahertz pulsed spectroscopy. International Journal of Pharmaceutics, 2007, 334, 78-84.	2.6	134
33	Characterisation of bicontinuous cubic liquid crystalline systems of phytantriol and water using cryo field emission scanning electron microscopy (cryo FESEM). Micron, 2007, 38, 478-485.	1.1	131
34	Precipitation of a Poorly Soluble Model Drug during In Vitro Lipolysis: Characterization and Dissolution of the Precipitate. Journal of Pharmaceutical Sciences, 2010, 99, 4982-4991.	1.6	131
35	Characterization of Temperature-Induced Phase Transitions in Five Polymorphic Forms of Sulfathiazole by Terahertz Pulsed Spectroscopy and Differential Scanning Calorimetry. Journal of Pharmaceutical Sciences, 2006, 95, 2486-2498.	1.6	126
36	The Influence of Thermal and Mechanical Preparative Techniques on the Amorphous State of Four Poorly Soluble Compounds. Journal of Pharmaceutical Sciences, 2005, 94, 1998-2012.	1.6	124

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37	Investigation of properties and recrystallisation behaviour of amorphous indomethacin samples prepared by different methods. International Journal of Pharmaceutics, 2011, 417, 94-100.	2.6	124
38	Correlating thermodynamic and kinetic parameters with amorphous stability. European Journal of Pharmaceutical Sciences, 2009, 37, 492-498.	1.9	123
39	Non-destructive quantification of pharmaceutical tablet coatings using terahertz pulsed imaging and optical coherence tomography. Optics and Lasers in Engineering, 2011, 49, 361-365.	2.0	120
40	Refining stability and dissolution rate of amorphous drug formulations. Expert Opinion on Drug Delivery, 2014, 11, 977-989.	2.4	119
41	Quantifying ternary mixtures of different solid-state forms of indomethacin by Raman and near-infrared spectroscopy. European Journal of Pharmaceutical Sciences, 2007, 32, 182-192.	1.9	115
42	Supersaturated Self-Nanoemulsifying Drug Delivery Systems (Super-SNEDDS) Enhance the Bioavailability of the Poorly Water-Soluble Drug Simvastatin in Dogs. AAPS Journal, 2013, 15, 219-227.	2.2	114
43	Bicontinuous cubic liquid crystals as sustained delivery systems for peptides and proteins. Expert Opinion on Drug Delivery, 2010, 7, 1133-1144.	2.4	112
44	Comparative Study of Different Methods for the Prediction of Drug–Polymer Solubility. Molecular Pharmaceutics, 2015, 12, 3408-3419.	2.3	111
45	Comparative study of liposomes, transfersomes, ethosomes and cubosomes for transcutaneous immunisation: characterisation and in vitro skin penetration. Journal of Pharmacy and Pharmacology, 2012, 64, 1560-1569.	1.2	110
46	Determination of Solubility Parameters of Ibuprofen and Ibuprofen Lysinate. Molecules, 2015, 20, 21549-21568.	1.7	110
47	Liquid Crystalline Systems of Phytantriol and Glyceryl Monooleate Containing a Hydrophilic Protein: Characterisation, Swelling and Release Kinetics. Journal of Pharmaceutical Sciences, 2009, 98, 4191-4204.	1.6	107
48	Effects of alcohols and diols on the phase behaviour of quaternary systems. International Journal of Pharmaceutics, 2000, 196, 141-145.	2.6	105
49	Improving Co-Amorphous Drug Formulations by the Addition of the Highly Water Soluble Amino Acid, Proline. Pharmaceutics, 2014, 6, 416-435.	2.0	105
50	Screening for differences in the amorphous state of indomethacin using multivariate visualization. European Journal of Pharmaceutical Sciences, 2007, 30, 113-123.	1.9	101
51	Lipid based particulate formulations for the delivery of antigen. Immunology and Cell Biology, 2005, 83, 97-105.	1.0	100
52	Drug nanocrystallisation within liposomes. Journal of Controlled Release, 2018, 288, 96-110.	4.8	100
53	Characterizing the conversion kinetics of carbamazepine polymorphs to the dihydrate in aqueous suspension using Raman spectroscopy. Journal of Pharmaceutical and Biomedical Analysis, 2006, 40, 271-280.	1.4	99
54	Cubosomes containing the adjuvants imiquimod and monophosphoryl lipid A stimulate robust cellular and humoral immune responses. Journal of Controlled Release, 2013, 165, 16-21.	4.8	98

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55	In Vitro Lipolysis Data Does Not Adequately Predict the In Vivo Performance of Lipid-Based Drug Delivery Systems Containing Fenofibrate. AAPS Journal, 2014, 16, 539-549.	2.2	98
56	Predicting Crystallization of Amorphous Drugs with Terahertz Spectroscopy. Molecular Pharmaceutics, 2015, 12, 3062-3068.	2.3	97
57	A theoretical and spectroscopic study of co-amorphous naproxen and indomethacin. International Journal of Pharmaceutics, 2013, 453, 80-87.	2.6	95
58	Preparation and characterization of spray-dried co-amorphous drug–amino acid salts. Journal of Pharmacy and Pharmacology, 2016, 68, 615-624.	1.2	95
59	Quantitative analysis of polymorphic mixtures of ranitidine hydrochloride by Raman spectroscopy and principal components analysis. European Journal of Pharmaceutics and Biopharmaceutics, 2002, 54, 337-341.	2.0	94
60	Supersaturating drug delivery systems: The potential of co-amorphous drug formulations. International Journal of Pharmaceutics, 2017, 532, 1-12.	2.6	93
61	Glass-Transition Temperature of the β-Relaxation as the Major Predictive Parameter for Recrystallization of Neat Amorphous Drugs. Journal of Physical Chemistry B, 2018, 122, 2803-2808.	1.2	93
62	Characterizing Colloidal Structures of Pseudoternary Phase Diagrams Formed by Oil/Water/Amphiphile Systems. Drug Development and Industrial Pharmacy, 2001, 27, 31-38.	0.9	92
63	Perspectives in the use of spectroscopy to characterise pharmaceutical solids. International Journal of Pharmaceutics, 2008, 364, 159-169.	2.6	90
64	Dry Hybrid Lipidâ^'Silica Microcapsules Engineered from Submicron Lipid Droplets and Nanoparticles as a Novel Delivery System for Poorly Soluble Drugs. Molecular Pharmaceutics, 2009, 6, 861-872.	2.3	90
65	Characterisation of pore structures of pharmaceutical tablets: A review. International Journal of Pharmaceutics, 2018, 538, 188-214.	2.6	90
66	The Potential of Small-Scale Fusion Experiments and the Gordon-Taylor Equation to Predict the Suitability of Drug/Polymer Blends for Melt Extrusion. Drug Development and Industrial Pharmacy, 2001, 27, 549-560.	0.9	88
67	Amino Acids as Co-amorphous Excipients for Simvastatin and Glibenclamide: Physical Properties and Stability. Molecular Pharmaceutics, 2014, 11, 2381-2389.	2.3	88
68	Temperature dependent terahertz pulsed spectroscopy of carbamazepine. Thermochimica Acta, 2005, 436, 71-77.	1.2	85
69	Physicochemical Properties and Stability of Two Differently Prepared Amorphous Forms of Simvastatin. Crystal Growth and Design, 2008, 8, 128-135.	1.4	85
70	Understanding the solid-state forms of fenofibrate – A spectroscopic and computational study. European Journal of Pharmaceutics and Biopharmaceutics, 2009, 71, 100-108.	2.0	85
71	Characterization of microemulsion structures in the pseudoternary phase diagram of isopropyl palmitate/water/Brij 97:1-butanol. AAPS PharmSciTech, 2006, 7, E99-E104.	1.5	84
72	Influence of Polymer Molecular Weight on Drug–polymer Solubility: A Comparison between Experimentally Determined Solubility in PVP and Prediction Derived from Solubility in Monomer. Journal of Pharmaceutical Sciences, 2015, 104, 2905-2912.	1.6	84

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73	Polymer-Based Prodrugs: Improving Tumor Targeting and the Solubility of Small Molecule Drugs in Cancer Therapy. Molecules, 2015, 20, 21750-21769.	1.7	84
74	Adsorption of bovine serum albumin (BSA) onto lecithin studied by attenuated total reflectance Fourier transform infrared (ATR-FTIR) spectroscopy. International Journal of Pharmaceutics, 2007, 337, 40-47.	2.6	81
75	Applications of terahertz pulsed imaging to sustained-release tablet film coating quality assessment and dissolution performance. Journal of Controlled Release, 2008, 127, 79-87.	4.8	81
76	Solid-state properties and dissolution behaviour of tablets containing co-amorphous indomethacin–arginine. European Journal of Pharmaceutics and Biopharmaceutics, 2015, 96, 44-52.	2.0	80
77	Formation Kinetics and Stability of Carbamazepineâ^'Nicotinamide Cocrystals Prepared by Mechanical Activation. Crystal Growth and Design, 2009, 9, 2377-2386.	1.4	79
78	Characterising Lipid Lipolysis and Its Implication in Lipid-Based Formulation Development. AAPS Journal, 2012, 14, 860-871.	2.2	79
79	Nonlamellar liquid crystalline nanostructured particles: advances in materials and structure determination. Journal of Liposome Research, 2009, 19, 12-28.	1.5	78
80	Mannosylated liposomes as antigen delivery vehicles for targeting to dendritic cellsâ€. Journal of Pharmacy and Pharmacology, 2010, 58, 729-737.	1.2	78
81	Development of a screening method for co-amorphous formulations of drugs and amino acids. European Journal of Pharmaceutical Sciences, 2016, 95, 28-35.	1.9	78
82	Silica Nanoparticles To Control the Lipase-Mediated Digestion of Lipid-Based Oral Delivery Systems. Molecular Pharmaceutics, 2010, 7, 522-532.	2.3	76
83	In vitro digestion models to evaluate lipid based drug delivery systems; present status and current trends. Advanced Drug Delivery Reviews, 2019, 142, 35-49.	6.6	76
84	Co-former selection for co-amorphous drug-amino acid formulations. International Journal of Pharmaceutics, 2019, 557, 366-373.	2.6	76
85	Recent pharmaceutical applications of raman and terahertz spectroscopies. Journal of Pharmaceutical Sciences, 2008, 97, 4598-4621.	1.6	75
86	Solubilisation of soybean oil in microemulsions using various surfactants. Food Hydrocolloids, 2006, 20, 253-260.	5.6	72
87	Formation Mechanism of Coamorphous Drug–Amino Acid Mixtures. Molecular Pharmaceutics, 2015, 12, 2484-2492.	2.3	72
88	Co-Amorphous Drug Formulations in Numbers: Recent Advances in Co-Amorphous Drug Formulations with Focus on Co-Formability, Molar Ratio, Preparation Methods, Physical Stability, In Vitro and In Vivo Performance, and New Formulation Strategies. Pharmaceutics, 2021, 13, 389.	2.0	71
89	Using different structure types of microemulsions for the preparation of poly(alkylcyanoacrylate) nanoparticles by interfacial polymerization. Journal of Controlled Release, 2005, 106, 76-87.	4.8	70
90	Self-Assembled Geometric Liquid-Crystalline Nanoparticles Imaged in Three Dimensions:  Hexosomes Are Not Necessarily Flat Hexagonal Prisms. Langmuir, 2007, 23, 12461-12464.	1.6	70

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91	Influence of Lipid Composition and Drug Load on the In Vitro Performance of Self-Nanoemulsifying Drug Delivery Systems. Journal of Pharmaceutical Sciences, 2012, 101, 1721-1731.	1.6	70
92	Analysis of 3D Prints by X-ray Computed Microtomography andÂTerahertz Pulsed Imaging. Pharmaceutical Research, 2017, 34, 1037-1052.	1.7	69
93	Quantitative analysis of polymorphic mixtures of carbamazepine by Raman spectroscopy and principal components analysis. Journal of Raman Spectroscopy, 2004, 35, 347-352.	1.2	68
94	Effect of milling conditions on the solid-state conversion of ranitidine hydrochloride form 1. International Journal of Pharmaceutics, 2006, 327, 36-44.	2.6	68
95	A theoretical and spectroscopic study of γ-crystalline and amorphous indometacin. Journal of Pharmacy and Pharmacology, 2010, 59, 261-269.	1.2	68
96	Insights into the Early Dissolution Events of Amlodipine Using UV Imaging and Raman Spectroscopy. Molecular Pharmaceutics, 2011, 8, 1372-1380.	2.3	68
97	Influence of solvent evaporation rate and formulation factors on solid dispersion physical stability. European Journal of Pharmaceutical Sciences, 2011, 44, 610-620.	1.9	68
98	Microemulsions containing lecithin and sugar-based surfactants: Nanoparticle templates for delivery of proteins and peptides. International Journal of Pharmaceutics, 2008, 350, 351-360.	2.6	67
99	The Role of Configurational Entropy in Amorphous Systems. Pharmaceutics, 2010, 2, 224-244.	2.0	67
100	Development and characterisation of modified poloxamer 407 thermoresponsive depot systems containing cubosomes. International Journal of Pharmaceutics, 2011, 408, 20-26.	2.6	66
101	An oral delivery system for indomethicin engineered from cationic lipid emulsions and silica nanoparticles. Journal of Controlled Release, 2010, 143, 367-373.	4.8	65
102	On the role of salt formation and structural similarity of co-formers in co-amorphous drug delivery systems. International Journal of Pharmaceutics, 2018, 535, 86-94.	2.6	65
103	Terahertz pulsed imaging as an analytical tool for sustained-release tablet film coating. European Journal of Pharmaceutics and Biopharmaceutics, 2009, 71, 117-123.	2.0	64
104	In vivo evaluation of chitosan as an adjuvant in subcutaneous vaccine formulations. Vaccine, 2013, 31, 4812-4819.	1.7	64
105	Activation of the NLRP3 inflammasome is not a feature of all particulate vaccine adjuvants. Immunology and Cell Biology, 2014, 92, 535-542.	1.0	64
106	Influence of variation in molar ratio on co-amorphous drug-amino acid systems. European Journal of Pharmaceutics and Biopharmaceutics, 2016, 107, 32-39.	2.0	64
107	The administration route is decisive for the ability of the vaccine adjuvant CAF09 to induce antigen-specific CD8 + T-cell responses: The immunological consequences of the biodistribution profile. Journal of Controlled Release, 2016, 239, 107-117.	4.8	62
108	Influence of polymer molecular weight on in vitro dissolution behavior and in vivo performance of celecoxib:PVP amorphous solid dispersions. European Journal of Pharmaceutics and Biopharmaceutics, 2016, 101, 145-151.	2.0	62

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109	Characterisation of indomethacin and nifedipine using variable-temperature solid-state NMR. Magnetic Resonance in Chemistry, 2005, 43, 881-892.	1.1	61
110	Monitoring tablet surface roughness during the film coating process. AAPS PharmSciTech, 2006, 7, E1-E6.	1.5	61
111	Hot Melt Extrusion and Spray Drying of Co-amorphous Indomethacin-Arginine With Polymers. Journal of Pharmaceutical Sciences, 2017, 106, 302-312.	1.6	61
112	Application of a Salt Coformer in a Co-Amorphous Drug System Dramatically Enhances the Glass Transition Temperature: A Case Study of the Ternary System Carbamazepine, Citric Acid, and <scp>l</scp> -Arginine. Molecular Pharmaceutics, 2018, 15, 2036-2044.	2.3	61
113	Poly(alkycyanoacrylate) nanoparticles for enhanced delivery of therapeutics – is there real potential?. Expert Opinion on Drug Delivery, 2009, 6, 371-387.	2.4	60
114	Determination of polymorphic forms of ranitidine–HCl by DRIFTS and XRPD. Journal of Pharmaceutical and Biomedical Analysis, 2001, 25, 741-750.	1.4	59
115	Polymeric microcontainers improve oral bioavailability of furosemide. International Journal of Pharmaceutics, 2016, 504, 98-109.	2.6	59
116	Factors influencing the entrapment of hydrophilic compounds in nanocapsules prepared by interfacial polymerisation of water-in-oil microemulsions. European Journal of Pharmaceutics and Biopharmaceutics, 2002, 53, 335-342.	2.0	58
117	Comparative effects of intraduodenal infusions of lauric and oleic acids on antropyloroduodenal motility, plasma cholecystokinin and peptide YY, appetite, and energy intake in healthy men. American Journal of Clinical Nutrition, 2008, 87, 1181-1187.	2.2	58
118	Preparation of an amorphous sodium furosemide salt improves solubility and dissolution rate and leads to a faster Tmax after oral dosing to rats. European Journal of Pharmaceutics and Biopharmaceutics, 2013, 85, 942-951.	2.0	58
119	Effects of formulation variables on characteristics of poly (ethylcyanoacrylate) nanocapsules prepared from w/o microemulsions. International Journal of Pharmaceutics, 2002, 235, 237-246.	2.6	57
120	Influence of Polymorphic Form, Morphology, and Excipient Interactions on the Dissolution of Carbamazepine Compacts. Journal of Pharmaceutical Sciences, 2007, 96, 584-594.	1.6	57
121	Oral insulin delivery using nanoparticles based on microemulsions with different structure-types: Optimisation and in vivo evaluation. European Journal of Pharmaceutical Sciences, 2009, 37, 53-61.	1.9	57
122	Transcutaneous immunization using microneedles and cubosomes: Mechanistic investigations using Optical Coherence Tomography and Two-Photon Microscopy. Journal of Controlled Release, 2013, 172, 894-903.	4.8	57
123	Amorphous drugs and dosage forms. Journal of Drug Delivery Science and Technology, 2013, 23, 403-408.	1.4	57
124	Performance comparison between crystalline and co-amorphous salts of indomethacin-lysine. International Journal of Pharmaceutics, 2017, 533, 138-144.	2.6	57
125	Organic acids as co-formers for co-amorphous systems – Influence of variation in molar ratio on the physicochemical properties of the co-amorphous systems. European Journal of Pharmaceutics and Biopharmaceutics, 2018, 131, 25-32.	2.0	56
126	Relaxation and Crystallization of Amorphous Carbamazepine Studied by Terahertz Pulsed Spectroscopy. Journal of Pharmaceutical Sciences, 2007, 96, 2703-2709.	1.6	55

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127	In-vitro release and oral bioactivity of insulin in diabetic rats using nanocapsules dispersed in biocompatible microemulsion. Journal of Pharmacy and Pharmacology, 2010, 54, 473-480.	1.2	55
128	Improvement of dissolution rate of indomethacin by inkjet printing. European Journal of Pharmaceutical Sciences, 2015, 75, 91-100.	1.9	55
129	The Precipitation Behavior of Poorly Water-Soluble Drugs with an Emphasis on the Digestion of Lipid Based Formulations. Pharmaceutical Research, 2016, 33, 548-562.	1.7	55
130	Formation and physical stability of the amorphous phase of ranitidine hydrochloride polymorphs prepared by cryo-milling. European Journal of Pharmaceutics and Biopharmaceutics, 2008, 68, 771-780.	2.0	54
131	In vitro and in vivo investigation of thermosensitive chitosan hydrogels containing silica nanoparticles for vaccine delivery. European Journal of Pharmaceutical Sciences, 2010, 41, 360-368.	1.9	54
132	Spatial confinement can lead to increased stability of amorphous indomethacin. European Journal of Pharmaceutics and Biopharmaceutics, 2012, 81, 418-425.	2.0	54
133	Effects of film coating thickness and drug layer uniformity on in vitro drug release from sustained-release coated pellets: A case study using terahertz pulsed imaging. International Journal of Pharmaceutics, 2009, 382, 151-159.	2.6	53
134	The influence of various excipients on the conversion kinetics of carbamazepine polymorphs in aqueous suspension. Journal of Pharmacy and Pharmacology, 2010, 59, 193-201.	1.2	53
135	Investigation of the Formation Process of Two Piracetam Cocrystals during Grinding. Pharmaceutics, 2011, 3, 706-722.	2.0	53
136	Studying the Propensity of Compounds to Supersaturate: A Practical and Broadly Applicable Approach. Journal of Pharmaceutical Sciences, 2016, 105, 3021-3029.	1.6	53
137	Investigation of physical properties and stability of indomethacin–cimetidine and naproxen–cimetidine co-amorphous systems prepared by quench cooling, coprecipitation and ball milling. Journal of Pharmacy and Pharmacology, 2016, 68, 36-45.	1.2	53
138	Influence of PVP/VA copolymer composition on drug–polymer solubility. European Journal of Pharmaceutical Sciences, 2016, 85, 10-17.	1.9	53
139	Visualizing the conversion of carbamazepine in aqueous suspension with and without the presence of excipients: A single crystal study using SEM and Raman microscopy. European Journal of Pharmaceutics and Biopharmaceutics, 2006, 64, 326-335.	2.0	52
140	Chitosan–magnesium aluminum silicate composite dispersions: Characterization of rheology, flocculate size and zeta potential. International Journal of Pharmaceutics, 2008, 351, 227-235.	2.6	51
141	Quantitative solid-state analysis of three solid forms of ranitidine hydrochloride in ternary mixtures using Raman spectroscopy and X-ray powder diffraction. Journal of Pharmaceutical and Biomedical Analysis, 2009, 49, 18-25.	1.4	51
142	Recent advances and potential applications of modulated differential scanning calorimetry (mDSC) in drug development. European Journal of Pharmaceutical Sciences, 2016, 87, 164-173.	1.9	51
143	The significance of the amorphous potential energy landscape for dictating glassy dynamics and driving solid-state crystallisation. Physical Chemistry Chemical Physics, 2017, 19, 30039-30047.	1.3	51
144	On the preparation, microscopic investigation and application of ISCOMs. Micron, 2006, 37, 724-734.	1.1	50

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145	The application of MALDI TOF MS in biopharmaceutical research. International Journal of Pharmaceutics, 2011, 417, 70-82.	2.6	49
146	The Role of Glass Transition Temperatures in Coamorphous Drug–Amino Acid Formulations. Molecular Pharmaceutics, 2018, 15, 4247-4256.	2.3	49
147	Solidâ€State Transition Mechanism in Carbamazepine Polymorphs by Timeâ€Resolved Terahertz Spectroscopy. ChemPhysChem, 2007, 8, 1924-1927.	1.0	48
148	Influence of particle size and preparation methods on the physical and chemical stability of amorphous simvastatin. European Journal of Pharmaceutics and Biopharmaceutics, 2009, 71, 64-70.	2.0	48
149	Chitosan hydrogels containing liposomes and cubosomes as particulate sustained release vaccine delivery systems. Journal of Liposome Research, 2012, 22, 193-204.	1.5	48
150	Comparison of two DSC-based methods to predict drug-polymer solubility. International Journal of Pharmaceutics, 2018, 540, 98-105.	2.6	48
151	Pseudo-ternary phase diagrams of aqueous mixtures of Quil A, cholesterol and phospholipid prepared by the lipid-film hydration method. International Journal of Pharmaceutics, 2004, 270, 229-239.	2.6	47
152	Dose-related effects of lauric acid on antropyloroduodenal motility, gastrointestinal hormone release, appetite, and energy intake in healthy men. American Journal of Physiology - Regulatory Integrative and Comparative Physiology, 2005, 289, R1090-R1098.	0.9	47
153	Effects of lauric acid on upper gut motility, plasma cholecystokinin and peptide YY, and energy intake are load, but not concentration, dependent in humans. Journal of Physiology, 2007, 581, 767-777.	1.3	47
154	Melt Extrusion and Spray Drying of Carbamazepine and Dipyridamole with Polyvinylpyrrolidone/Vinyl Acetate Copolymers. Drug Development and Industrial Pharmacy, 2008, 34, 95-106.	0.9	47
155	Commentary: Towards Physico-Relevant Dissolution Testing: The Importance of Solid-State Analysis in Dissolution. Dissolution Technologies, 2009, 16, 47-54.	0.2	47
156	Comparison of chitosan nanoparticles and chitosan hydrogels for vaccine delivery. Journal of Pharmacy and Pharmacology, 2010, 60, 1591-1600.	1.2	46
157	In Situ Lipolysis and Synchrotron Small-Angle X-ray Scattering for the Direct Determination of the Precipitation and Solid-State Form of a Poorly Water-Soluble Drug During Digestion of a Lipid-Based Formulation. Journal of Pharmaceutical Sciences, 2016, 105, 2631-2639.	1.6	46
158	Terahertz pulsed imaging as an advanced characterisation tool for film coatings—A review. International Journal of Pharmaceutics, 2013, 457, 510-520.	2.6	45
159	Use of low-frequency Raman spectroscopy and chemometrics for the quantification of crystallinity in amorphous griseofulvin tablets. Vibrational Spectroscopy, 2015, 77, 10-16.	1.2	45
160	Comparison of lipases for in vitro models of gastric digestion: lipolysis using two infant formulas as model substrates. Food and Function, 2016, 7, 3989-3998.	2.1	45
161	Are phytosomes a superior nanodelivery system for the antioxidant rutin?. International Journal of Pharmaceutics, 2018, 548, 82-91.	2.6	45
162	Transformations between Co-Amorphous and Co-Crystal Systems and Their Influence on the Formation and Physical Stability of Co-Amorphous Systems. Molecular Pharmaceutics, 2019, 16, 1294-1304.	2.3	45

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163	Comparison of the Structure and Properties of Liposomes Prepared from Milk Fat Globule Membrane and Soy Phospholipids. Journal of Agricultural and Food Chemistry, 2006, 54, 3704-3711.	2.4	44
164	Immunostimulatory colloidal delivery systems for cancer vaccines. Expert Opinion on Drug Delivery, 2006, 3, 345-354.	2.4	44
165	Protein delivery using nanoparticles based on microemulsions with different structure-types. European Journal of Pharmaceutical Sciences, 2008, 33, 434-444.	1.9	44
166	Investigations on the effect of different cooling rates on the stability of amorphous indomethacin. European Journal of Pharmaceutical Sciences, 2011, 44, 341-350.	1.9	44
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