Jukka Rantanen

List of Publications by Year in descending order

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6254 24,019 514 80 citations h-index papers

g-index 525 525 525 15792 docs citations times ranked citing authors all docs

18647

119

#	Article	IF	CITATIONS
1	Emerging trends in the stabilization of amorphous drugs. International Journal of Pharmaceutics, 2013, 453, 65-79.	5.2	360
2	Recent advances in co-amorphous drug formulations. Advanced Drug Delivery Reviews, 2016, 100, 116-125.	13.7	350
3	Terahertz pulsed spectroscopy and imaging in the pharmaceutical setting - a review. Journal of Pharmacy and Pharmacology, 2010, 59, 209-223.	2.4	330
4	Using Terahertz Pulsed Spectroscopy to Quantify Pharmaceutical Polymorphism and Crystallinity. Journal of Pharmaceutical Sciences, 2005, 94, 837-846.	3.3	326
5	The Future of Pharmaceutical Manufacturing Sciences. Journal of Pharmaceutical Sciences, 2015, 104, 3612-3638.	3.3	303
6	Coamorphous Drug Systems: Enhanced Physical Stability and Dissolution Rate of Indomethacin and Naproxen. Molecular Pharmaceutics, 2011, 8, 1919-1928.	4.6	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.	2.4	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.	4.3	246
9	Solid form screening – A review. European Journal of Pharmaceutics and Biopharmaceutics, 2009, 71, 23-37.	4.3	237
10	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.	9.9	236
11	An overview of recent studies on the analysis of pharmaceutical polymorphs. Journal of Pharmaceutical and Biomedical Analysis, 2011, 55, 618-644.	2.8	233
12	Raman spectroscopy in pharmaceutical product design. Advanced Drug Delivery Reviews, 2015, 89, 3-20.	13.7	221
13	Using terahertz pulsed spectroscopy to study crystallinity of pharmaceutical materials. Chemical Physics Letters, 2004, 390, 20-24.	2.6	217
14	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.	2.4	205
15	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.	4.3	197
16	Raman spectroscopy for quantitative analysis of pharmaceutical solids. Journal of Pharmacy and Pharmacology, 2010, 59, 179-192.	2.4	196
17	High loading efficiency and sustained release of siRNA encapsulated in PLGA nanoparticles: Quality by design optimization and characterization. European Journal of Pharmaceutics and Biopharmaceutics, 2011, 77, 26-35.	4.3	191
18	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.	3.3	179

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19	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.	4.3	179
20	Analysis of solid-state transformations of pharmaceutical compounds using vibrational spectroscopy. Journal of Pharmacy and Pharmacology, 2010, 61, 971-988.	2.4	179
21	Cell-penetrating peptides for drug delivery across membrane barriers. Expert Opinion on Drug Delivery, 2008, 5, 105-117.	5.0	177
22	Spray drying of siRNA-containing PLGA nanoparticles intended for inhalation. Journal of Controlled Release, 2010, 142, 138-145.	9.9	176
23	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.	5.2	170
24	The adjuvant mechanism of cationic dimethyldioctadecylammonium liposomes. Immunology, 2007, 121, 216-226.	4.4	167
25	Design of an inhalable dry powder formulation of DOTAP-modified PLGA nanoparticles loaded with siRNA. Journal of Controlled Release, 2012, 157, 141-148.	9.9	162
26	Anti-tuberculosis drug combination for controlled oral delivery using 3D printed compartmental dosage forms: From drug product design to in vivo testing. Journal of Controlled Release, 2017, 268, 40-48.	9.9	154
27	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.	4.3	153
28	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.	5.2	149
29	Rheology as a tool for evaluation of melt processability of innovative dosage forms. International Journal of Pharmaceutics, 2015, 494, 623-642.	5.2	147
30	Design space approach in the optimization of the spray-drying process. European Journal of Pharmaceutics and Biopharmaceutics, 2012, 80, 226-234.	4.3	138
31	Intestinal mucosa permeability following oral insulin delivery using core shell corona nanolipoparticles. Biomaterials, 2013, 34, 9678-9687.	11.4	137
32	Drug hydrate systems and dehydration processes studied by terahertz pulsed spectroscopy. International Journal of Pharmaceutics, 2007, 334, 78-84.	5.2	134
33	Use of In-Line Near-Infrared Spectroscopy in Combination with Chemometrics for Improved Understanding of Pharmaceutical Processes. Analytical Chemistry, 2005, 77, 556-563.	6.5	132
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.	3.3	131
35	Three-Dimensional Printing of Drug-Eluting Implants: Preparation of an Antimicrobial Polylactide Feedstock Material. Journal of Pharmaceutical Sciences, 2015, 104, 1099-1107.	3.3	131
36	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.	3.3	126

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37	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.	3.3	124
38	Investigation of properties and recrystallisation behaviour of amorphous indomethacin samples prepared by different methods. International Journal of Pharmaceutics, 2011, 417, 94-100.	5.2	124
39	Correlating thermodynamic and kinetic parameters with amorphous stability. European Journal of Pharmaceutical Sciences, 2009, 37, 492-498.	4.0	123
40	Non-destructive quantification of pharmaceutical tablet coatings using terahertz pulsed imaging and optical coherence tomography. Optics and Lasers in Engineering, 2011, 49, 361-365.	3.8	120
41	Refining stability and dissolution rate of amorphous drug formulations. Expert Opinion on Drug Delivery, 2014, 11, 977-989.	5.0	119
42	Modifying release characteristics from 3D printed drug-eluting products. European Journal of Pharmaceutical Sciences, 2016, 90, 47-52.	4.0	118
43	Subunit vaccines of the future: the need for safe, customized and optimized particulate delivery systems. Therapeutic Delivery, 2011, 2, 1057-1077.	2.2	116
44	Chitosan-Based Nano-Embedded Microparticles: Impact of Nanogel Composition on Physicochemical Properties. Pharmaceutics, 2017, 9, 1.	4.5	116
45	Implementation of a Process Analytical Technology System in a Freeze-Drying Process Using Raman Spectroscopy for In-Line Process Monitoring. Analytical Chemistry, 2007, 79, 7992-8003.	6.5	115
46	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.	4.0	115
47	Stabilization of liposomes during drying. Expert Opinion on Drug Delivery, 2011, 8, 375-388.	5.0	114
48	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.	4.4	114
49	Comparative Study of Different Methods for the Prediction of Drug–Polymer Solubility. Molecular Pharmaceutics, 2015, 12, 3408-3419.	4.6	111
50	On-line monitoring of moisture content in an instrumented fluidized bed granulator with a multi-channel NIR moisture sensor. Powder Technology, 1998, 99, 163-170.	4.2	110
51	Solvent-Mediated Phase Transformation Kinetics of an Anhydrate/Hydrate System. Crystal Growth and Design, 2006, 6, 2053-2060.	3.0	106
52	Roadmap to 3D-Printed Oral Pharmaceutical Dosage Forms: Feedstock Filament Properties and Characterization for Fused Deposition Modeling. Journal of Pharmaceutical Sciences, 2019, 108, 26-35.	3.3	106
53	Improving Co-Amorphous Drug Formulations by the Addition of the Highly Water Soluble Amino Acid, Proline. Pharmaceutics, 2014, 6, 416-435.	4.5	105
54	License to kill: Formulation requirements for optimal priming of CD8+ CTL responses with particulate vaccine delivery systems. European Journal of Pharmaceutical Sciences, 2012, 45, 482-491.	4.0	103

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55	Ciprofloxacin-loaded sodium alginate/poly (lactic-co-glycolic acid) electrospun fibrous mats for wound healing. European Journal of Pharmaceutics and Biopharmaceutics, 2018, 123, 42-49.	4.3	103
56	Crystallization of glycine with ultrasound. International Journal of Pharmaceutics, 2006, 320, 23-29.	5.2	102
57	In-line moisture measurement during granulation with a four-wavelength near infrared sensor: an evaluation of particle size and binder effects. European Journal of Pharmaceutics and Biopharmaceutics, 2000, 50, 271-276.	4.3	101
58	Screening for differences in the amorphous state of indomethacin using multivariate visualization. European Journal of Pharmaceutical Sciences, 2007, 30, 113-123.	4.0	101
59	Hydrate formation during wet granulation studied by spectroscopic methods and multivariate analysis. Pharmaceutical Research, 2002, 19, 1285-1291.	3 . 5	99
60	Role of Water in the Physical Stability of Solid Dosage Formulations. Journal of Pharmaceutical Sciences, 2005, 94, 2147-2165.	3.3	99
61	Predicting Crystallization of Amorphous Drugs with Terahertz Spectroscopy. Molecular Pharmaceutics, 2015, 12, 3062-3068.	4.6	97
62	A theoretical and spectroscopic study of co-amorphous naproxen and indomethacin. International Journal of Pharmaceutics, 2013, 453, 80-87.	5.2	95
63	Preparation and characterization of spray-dried co-amorphous drug–amino acid salts. Journal of Pharmacy and Pharmacology, 2016, 68, 615-624.	2.4	95
64	Novel Identification of Pseudopolymorphic Changes of Theophylline During Wet Granulation Using Near Infrared Spectroscopy. Journal of Pharmaceutical Sciences, 2001, 90, 389-396.	3.3	94
65	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.	4.3	94
66	Supersaturating drug delivery systems: The potential of co-amorphous drug formulations. International Journal of Pharmaceutics, 2017, 532, 1-12.	5.2	93
67	Glass-Transition Temperature of the \hat{l}^2 -Relaxation as the Major Predictive Parameter for Recrystallization of Neat Amorphous Drugs. Journal of Physical Chemistry B, 2018, 122, 2803-2808.	2.6	93
68	Perspectives in the use of spectroscopy to characterise pharmaceutical solids. International Journal of Pharmaceutics, 2008, 364, 159-169.	5. 2	90
69	Characterisation of pore structures of pharmaceutical tablets: A review. International Journal of Pharmaceutics, 2018, 538, 188-214.	5. 2	90
70	Unidirectional drug release from 3D printed mucoadhesive buccal films using FDM technology: In vitro and ex vivo evaluation. European Journal of Pharmaceutics and Biopharmaceutics, 2019, 144, 180-192.	4.3	90
71	QR encoded smart oral dosage forms by inkjet printing. International Journal of Pharmaceutics, 2018, 536, 138-145.	5. 2	89
72	Amino Acids as Co-amorphous Excipients for Simvastatin and Glibenclamide: Physical Properties and Stability. Molecular Pharmaceutics, 2014, 11, 2381-2389.	4.6	88

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73	In Situ Measurement of Solvent-Mediated Phase Transformations During Dissolution Testing. Journal of Pharmaceutical Sciences, 2006, 95, 2730-2737.	3.3	87
74	Temperature dependent terahertz pulsed spectroscopy of carbamazepine. Thermochimica Acta, 2005, 436, 71-77.	2.7	85
75	Physicochemical Properties and Stability of Two Differently Prepared Amorphous Forms of Simvastatin. Crystal Growth and Design, 2008, 8, 128-135.	3.0	85
76	Understanding the solid-state forms of fenofibrate $\hat{a} \in \text{``} A$ spectroscopic and computational study. European Journal of Pharmaceutics and Biopharmaceutics, 2009, 71, 100-108.	4.3	85
77	A Step Toward Development of Printable Dosage Forms for Poorly Soluble Drugs. Journal of Pharmaceutical Sciences, 2013, 102, 3694-3704.	3.3	85
78	The role of mucus as an invisible cloak to transepithelial drug delivery by nanoparticles. Advanced Drug Delivery Reviews, 2018, 124, 107-124.	13.7	85
79	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.	3.3	84
80	Process analytical applications of Raman spectroscopy. Journal of Pharmacy and Pharmacology, 2010, 59, 171-177.	2.4	83
81	Comparison of the effects of two drying methods on polymorphism of theophylline. International Journal of Pharmaceutics, 2004, 276, 129-141.	5.2	82
82	Factors affecting crystallization of hydrates. Journal of Pharmacy and Pharmacology, 2010, 62, 1534-1546.	2.4	82
83	Physical stability and moisture sorption of aqueous chitosan–amylose starch films plasticized with polyols. European Journal of Pharmaceutics and Biopharmaceutics, 2004, 58, 69-76.	4.3	81
84	Applications of terahertz pulsed imaging to sustained-release tablet film coating quality assessment and dissolution performance. Journal of Controlled Release, 2008, 127, 79-87.	9.9	81
85	Status and future prospects of lipid-based particulate delivery systems as vaccine adjuvants and their combination with immunostimulators. Expert Opinion on Drug Delivery, 2009, 6, 657-672.	5.0	81
86	Solid-state properties and dissolution behaviour of tablets containing co-amorphous indomethacin–arginine. European Journal of Pharmaceutics and Biopharmaceutics, 2015, 96, 44-52.	4.3	80
87	Trehalose preserves DDA/TDB liposomes and their adjuvant effect during freeze-drying. Biochimica Et Biophysica Acta - Biomembranes, 2007, 1768, 2120-2129.	2.6	79
88	Formation Kinetics and Stability of Carbamazepineâ^Nicotinamide Cocrystals Prepared by Mechanical Activation. Crystal Growth and Design, 2009, 9, 2377-2386.	3.0	79
89	Characterising Lipid Lipolysis and Its Implication in Lipid-Based Formulation Development. AAPS Journal, 2012, 14, 860-871.	4.4	79
90	Development of a screening method for co-amorphous formulations of drugs and amino acids. European Journal of Pharmaceutical Sciences, 2016, 95, 28-35.	4.0	78

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91	Elucidating the molecular mechanism of PAMAM–siRNA dendriplex self-assembly: Effect of dendrimer charge density. International Journal of Pharmaceutics, 2011, 416, 410-418.	5. 2	77
92	Co-former selection for co-amorphous drug-amino acid formulations. International Journal of Pharmaceutics, 2019, 557, 366-373.	5.2	76
93	Recent pharmaceutical applications of raman and terahertz spectroscopies. Journal of Pharmaceutical Sciences, 2008, 97, 4598-4621.	3.3	75
94	Molecular Characterization of the Interaction between siRNA and PAMAM G7 Dendrimers by SAXS, ITC, and Molecular Dynamics Simulations. Biomacromolecules, 2010, 11, 3571-3577.	5.4	75
95	Design of PLGA-based depot delivery systems for biopharmaceuticals prepared by spray drying. International Journal of Pharmaceutics, 2016, 498, 82-95.	5.2	75
96	Transforming nanomedicine manufacturing toward Quality by Design and microfluidics. Advanced Drug Delivery Reviews, 2018, 128, 115-131.	13.7	75
97	Toward an Understanding of the Factors Influencing Anhydrate-to-Hydrate Transformation Kinetics in Aqueous Environments. Crystal Growth and Design, 2008, 8, 2684-2693.	3.0	72
98	Formation Mechanism of Coamorphous Drug–Amino Acid Mixtures. Molecular Pharmaceutics, 2015, 12, 2484-2492.	4.6	72
99	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.	4.5	71
100	Influence of raw material properties upon critical quality attributes of continuously produced granules and tablets. European Journal of Pharmaceutics and Biopharmaceutics, 2014, 87, 252-263.	4.3	70
101	Use of the Near-Infrared Reflectance Method for Measurement of Moisture Content During Granulation. Pharmaceutical Development and Technology, 2000, 5, 209-217.	2.4	69
102	Analysis of 3D Prints by X-ray Computed Microtomography andÂTerahertz Pulsed Imaging. Pharmaceutical Research, 2017, 34, 1037-1052.	3.5	69
103	Quantitative analysis of polymorphic mixtures of carbamazepine by Raman spectroscopy and principal components analysis. Journal of Raman Spectroscopy, 2004, 35, 347-352.	2.5	68
104	Improved Understanding of Factors Contributing to Quantification of Anhydrate/Hydrate Powder Mixtures. Applied Spectroscopy, 2005, 59, 942-951.	2.2	68
105	A theoretical and spectroscopic study of \hat{I}^3 -crystalline and amorphous indometacin. Journal of Pharmacy and Pharmacology, 2010, 59, 261-269.	2.4	68
106	Insights into the Early Dissolution Events of Amlodipine Using UV Imaging and Raman Spectroscopy. Molecular Pharmaceutics, 2011, 8, 1372-1380.	4.6	68
107	Influence of solvent evaporation rate and formulation factors on solid dispersion physical stability. European Journal of Pharmaceutical Sciences, 2011, 44, 610-620.	4.0	68
108	Excipient selection can significantly affect solid-state phase transformation in formulation during wet granulation. AAPS PharmSciTech, 2005, 6, E311-E322.	3.3	67

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109	The Role of Configurational Entropy in Amorphous Systems. Pharmaceutics, 2010, 2, 224-244.	4.5	67
110	A novel method of producing a microcrystalline \hat{l}^2 -sitosterol suspension in oil. European Journal of Pharmaceutical Sciences, 2002, 15, 261-269.	4.0	66
111	Mechanistic profiling of the siRNA delivery dynamics of lipid–polymer hybrid nanoparticles. Journal of Controlled Release, 2015, 201, 22-31.	9.9	66
112	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.	5.2	65
113	Influence of variation in molar ratio on co-amorphous drug-amino acid systems. European Journal of Pharmaceutics and Biopharmaceutics, 2016, 107, 32-39.	4.3	64
114	Effects of Excipients on Hydrate Formation in Wet Masses Containing Theophylline. Journal of Pharmaceutical Sciences, 2003, 92, 516-528.	3.3	63
115	Prediction of aqueous solubility for a diverse set of organic compounds based on atom-type electrotopological state indices. European Journal of Medicinal Chemistry, 2000, 35, 1081-1088.	5.5	61
116	Monitoring tablet surface roughness during the film coating process. AAPS PharmSciTech, 2006, 7, E1-E6.	3.3	61
117	Hot Melt Extrusion and Spray Drying of Co-amorphous Indomethacin-Arginine With Polymers. Journal of Pharmaceutical Sciences, 2017, 106, 302-312.	3.3	61
118	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> < scp>-Arginine. Molecular Pharmaceutics, 2018, 15, 2036-2044.</scp>	4.6	61
119	Microcrystalline cellulose-water interaction—a novel approach using thermoporosimetry. Pharmaceutical Research, 2001, 18, 1562-1569.	3.5	60
120	Critical Solvent Properties Affecting the Particle Formation Process and Characteristics of Celecoxib-Loaded PLGA Microparticles via Spray-Drying. Pharmaceutical Research, 2013, 30, 1065-1076.	3.5	59
121	Budesonide nanocrystal-loaded hyaluronic acid microparticles for inhalation: In vitro and in vivo evaluation. Carbohydrate Polymers, 2018, 181, 1143-1152.	10.2	59
122	Polymorph Screening Using Near-Infrared Spectroscopy. Analytical Chemistry, 2003, 75, 5267-5273.	6.5	58
123	Influence of Polymorphic Form, Morphology, and Excipient Interactions on the Dissolution of Carbamazepine Compacts. Journal of Pharmaceutical Sciences, 2007, 96, 584-594.	3.3	57
124	Performance comparison between crystalline and co-amorphous salts of indomethacin-lysine. International Journal of Pharmaceutics, 2017, 533, 138-144.	5.2	57
125	In-line moisture measurement during granulation with a four-wavelength near-infrared sensor: an evaluation of process-related variables and a development of non-linear calibration model. Chemometrics and Intelligent Laboratory Systems, 2001, 56, 51-58.	3.5	56
126	Near-Infrared Spectroscopy for Cocrystal Screening. A Comparative Study with Raman Spectroscopy. Analytical Chemistry, 2008, 80, 7755-7764.	6.5	56

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127	Nanoparticle-mediated delivery of the antimicrobial peptide plectasin against Staphylococcus aureus in infected epithelial cells. European Journal of Pharmaceutics and Biopharmaceutics, 2015, 92, 65-73.	4.3	56
128	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.	4.3	56
129	Quality by design thinking in the development of long-acting injectable PLGA/PLA-based microspheres for peptide and protein drug delivery. International Journal of Pharmaceutics, 2020, 585, 119441.	5.2	56
130	IR spectroscopy together with multivariate data analysis as a process analytical tool for in-line monitoring of crystallization process and solid-state analysis of crystalline product. Journal of Pharmaceutical and Biomedical Analysis, 2005, 38, 275-284.	2.8	55
131	Cellular uptake and membrane-destabilising properties of α-peptide/β-peptoid chimeras: lessons for the design of new cell-penetrating peptides. Biochimica Et Biophysica Acta - Biomembranes, 2008, 1778, 2487-2495.	2.6	55
132	Improvement of dissolution rate of indomethacin by inkjet printing. European Journal of Pharmaceutical Sciences, 2015, 75, 91-100.	4.0	55
133	Establishing quantitative in-line analysis of multiple solid-state transformations during dehydration. Journal of Pharmaceutical Sciences, 2008, 97, 4983-4999.	3.3	54
134	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.	4.3	54
135	Spatial confinement can lead to increased stability of amorphous indomethacin. European Journal of Pharmaceutics and Biopharmaceutics, 2012, 81, 418-425.	4.3	54
136	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.	5.2	53
137	The influence of various excipients on the conversion kinetics of carbamazepine polymorphs in aqueous suspension. Journal of Pharmacy and Pharmacology, 2010, 59, 193-201.	2.4	53
138	Investigation of the Formation Process of Two Piracetam Cocrystals during Grinding. Pharmaceutics, 2011, 3, 706-722.	4.5	53
139	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.	2.4	53
140	Engineering of small interfering RNA-loaded lipidoid-poly(DL -lactic-co-glycolic acid) hybrid nanoparticles for highly efficient and safe gene silencing: A quality by design-based approach. European Journal of Pharmaceutics and Biopharmaceutics, 2017, 120, 22-33.	4.3	53
141	Co-delivery of resveratrol and docetaxel via polymeric micelles to improve the treatment of drug-resistant tumors. Asian Journal of Pharmaceutical Sciences, 2019, 14, 78-85.	9.1	52
142	Fabrication of Mucoadhesive Buccal Films for Local Administration of Ketoprofen and Lidocaine Hydrochloride by Combining Fused Deposition Modeling and Inkjet Printing. Journal of Pharmaceutical Sciences, 2020, 109, 2757-2766.	3.3	52
143	Qualitative in situ analysis of multiple solidâ€state forms using spectroscopy and partial least squares discriminant modeling. Journal of Pharmaceutical Sciences, 2007, 96, 1802-1820.	3.3	51
144	Solvent Diversity in Polymorph Screening. Journal of Pharmaceutical Sciences, 2008, 97, 2145-2159.	3.3	51

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145	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.	2.8	51
146	Particle size dependence of polymorphism in spray-dried mannitol. European Journal of Pharmaceutical Sciences, 2011, 44, 41-48.	4.0	51
147	Incorporation of the TLR4 Agonist Monophosphoryl Lipid A Into the Bilayer of DDA/TDB Liposomes: Physico-Chemical Characterization and Induction of CD8+ T-Cell Responses In Vivo. Pharmaceutical Research, 2011, 28, 553-562.	3 . 5	51
148	siRNA Delivery with Lipid-based Systems: Promises and Pitfalls. Current Topics in Medicinal Chemistry, 2012, 12, 97-107.	2.1	51
149	Recent advances and potential applications of modulated differential scanning calorimetry (mDSC) in drug development. European Journal of Pharmaceutical Sciences, 2016, 87, 164-173.	4.0	51
150	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.	2.8	51
151	Cryptopharmaceuticals: Increasing the Safety of Medication by a Blockchain of Pharmaceutical Products. Journal of Pharmaceutical Sciences, 2019, 108, 2838-2841.	3.3	51
152	Quality by design approach in the optimization of the spray-drying process. Pharmaceutical Development and Technology, 2012, 17, 389-397.	2.4	50
153	Solvent-Mediated Solid Phase Transformations of cArbamazepine: Effects of Simulated Intestinal Fluid and Fasted State Simulated Intestinal Fluid. Journal of Pharmaceutical Sciences, 2009, 98, 985-996.	3.3	49
154	Particle formation and characteristics of Celecoxib-loaded poly(lactic-co-glycolic acid) microparticles prepared in different solvents using electrospraying. Polymer, 2012, 53, 3220-3229.	3.8	49
155	Strategic funding priorities in the pharmaceutical sciences allied to Quality by Design (QbD) and Process Analytical Technology (PAT). European Journal of Pharmaceutical Sciences, 2012, 47, 402-405.	4.0	49
156	The Role of Glass Transition Temperatures in Coamorphous Drug–Amino Acid Formulations. Molecular Pharmaceutics, 2018, 15, 4247-4256.	4.6	49
157	Characterization of polymorphic solid-state changes using variable temperature X-ray powder diffraction. Journal of Pharmaceutical and Biomedical Analysis, 2005, 39, 27-32.	2.8	48
158	Solidâ€State Transition Mechanism in Carbamazepine Polymorphs by Timeâ€Resolved Terahertz Spectroscopy. ChemPhysChem, 2007, 8, 1924-1927.	2.1	48
159	In-line monitoring of solid-state transitions during fluidisation. Chemical Engineering Science, 2007, 62, 408-415.	3.8	48
160	Comparison of two DSC-based methods to predict drug-polymer solubility. International Journal of Pharmaceutics, 2018, 540, 98-105.	5.2	48
161	Tablet surface characterisation by various imaging techniques. International Journal of Pharmaceutics, 2003, 254, 281-286.	5. 2	47
162	Exploring the Solid-Form Landscape of Pharmaceutical Hydrates: Transformation Pathways of the Sodium Naproxen Anhydrate-Hydrate System. Pharmaceutical Research, 2013, 30, 280-289.	3.5	47

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163	Commentary: Towards Physico-Relevant Dissolution Testing: The Importance of Solid-State Analysis in Dissolution. Dissolution Technologies, 2009, 16, 47-54.	0.6	47
164	New Perspectives for Visual Characterization of Pharmaceutical Solids. Journal of Pharmaceutical Sciences, 2004, 93, 165-176.	3.3	46
165	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.	3.3	46
166	Lipidoid-polymer hybrid nanoparticles loaded with TNF siRNA suppress inflammation after intra-articular administration in a murine experimental arthritis model. European Journal of Pharmaceutics and Biopharmaceutics, 2019, 142, 38-48.	4.3	46
167	An insight into water of crystallization during processing using vibrational spectroscopy. Journal of Pharmaceutical Sciences, 2009, 98, 3903-3932.	3.3	45
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