

Etsuo Yonemochi

List of Publications by Year in descending order

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215
papers

3,881
citations

109321

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197818

49
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222
all docs

222
docs citations

222
times ranked

3336
citing authors

#	ARTICLE	IF	CITATIONS
1	Effect of Water-Soluble Carriers on Dissolution Characteristics of Nifedipine Solid Dispersions. <i>Drug Development and Industrial Pharmacy</i> , 2000, 26, 1141-1150.	2.0	110
2	Cocrystallization and amorphization induced by drug-excipient interaction improves the physical properties of acyclovir. <i>International Journal of Pharmaceutics</i> , 2012, 422, 160-169.	5.2	108
3	Formulation design of a novel fast-disintegrating tablet. <i>International Journal of Pharmaceutics</i> , 2005, 306, 83-90.	5.2	105
4	The use of near infra-red spectroscopy to detect changes in the form of amorphous and crystalline lactose. <i>International Journal of Pharmaceutics</i> , 1998, 168, 231-241.	5.2	98
5	Evaluation of rapidly disintegrating tablets containing glycine and carboxymethylcellulose. <i>International Journal of Pharmaceutics</i> , 2006, 310, 101-109.	5.2	92
6	Improved dissolution of ofloxacin via solid dispersion. <i>International Journal of Pharmaceutics</i> , 1997, 156, 175-180.	5.2	85
7	Physicochemical properties of amorphous clarithromycin obtained by grinding and spray drying. <i>European Journal of Pharmaceutical Sciences</i> , 1999, 7, 331-338.	4.0	80
8	Self-Degradable Lipid-Like Materials Based on Hydrolysis accelerated by the intra-Particle Enrichment of Reactant (HyPER) for Messenger RNA Delivery. <i>Advanced Functional Materials</i> , 2020, 30, 1910575.	14.9	65
9	Evaluation of solid dispersions on a molecular level by the Raman mapping technique. <i>International Journal of Pharmaceutics</i> , 2008, 361, 12-18.	5.2	61
10	Freeze-drying of proteins with glass-forming oligosaccharide-derived sugar alcohols. <i>International Journal of Pharmaceutics</i> , 2010, 389, 107-113.	5.2	61
11	Novel Approach to DPI Carrier Lactose with Mechanofusion Process with Additives and Evaluation by IGC. <i>Chemical and Pharmaceutical Bulletin</i> , 2006, 54, 1508-1514.	1.3	58
12	Drug-Drug Multicomponent Crystals as an Effective Technique to Overcome Weaknesses in Parent Drugs. <i>Crystal Growth and Design</i> , 2016, 16, 3577-3581.	3.0	52
13	Correlation between Glass-Forming Ability and Fragility of Pharmaceutical Compounds. <i>Journal of Physical Chemistry B</i> , 2015, 119, 4873-4880.	2.6	51
14	Solid-state ¹³ C NMR study of indomethacin polymorphism. <i>International Journal of Pharmaceutics</i> , 2006, 318, 146-153.	5.2	50
15	Relationship between Crystallization Tendencies during Cooling from Melt and Isothermal Storage: Toward a General Understanding of Physical Stability of Pharmaceutical Glasses. <i>Molecular Pharmaceutics</i> , 2014, 11, 1835-1843.	4.6	48
16	Freeze-Drying of Proteins in Glass Solids Formed by Basic Amino Acids and Dicarboxylic Acids. <i>Chemical and Pharmaceutical Bulletin</i> , 2009, 57, 43-48.	1.3	46
17	Characterization and Quality Control of Pharmaceutical Cocrystals. <i>Chemical and Pharmaceutical Bulletin</i> , 2016, 64, 1421-1430.	1.3	46
18	Solubility improvement of epalrestat by layered structure formation via cocrystallization. <i>CrystEngComm</i> , 2017, 19, 2614-2622.	2.6	45

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19	Effects of sugar ester and hydroxypropyl methylcellulose on the physicochemical stability of amorphous cefditoren pivoxil in aqueous suspension. <i>International Journal of Pharmaceutics</i> , 2005, 290, 91-99.	5.2	44
20	Evaluation of antitumor effect of zoledronic acid entrapped in folate-linked liposome for targeting to tumor-associated macrophages. <i>Journal of Liposome Research</i> , 2015, 25, 131-140.	3.3	44
21	Development of Fast Disintegrating Compressed Tablets Using Amino Acid as Disintegration Accelerator: Evaluation of Wetting and Disintegration of Tablet on the Basis of Surface Free Energy. <i>Chemical and Pharmaceutical Bulletin</i> , 2005, 53, 1536-1539.	1.3	43
22	Application of Eudragit RS to thermo-sensitive drug delivery systems. <i>Journal of Controlled Release</i> , 2005, 102, 49-57.	9.9	43
23	Structural change and complexation of strictly linear amylose induced by sealed-heating with salicylic acid. <i>Journal of the Chemical Society, Faraday Transactions</i> , 1998, 94, 923-927.	1.7	42
24	Estimation of physical stability of amorphous solid dispersion using differential scanning calorimetry. <i>Journal of Thermal Analysis and Calorimetry</i> , 2006, 85, 689-692.	3.6	42
25	Physicochemical Understanding of Polymorphism and Solid-State Dehydration/Rehydration Processes for the Pharmaceutical Material Acrinol, by Ab Initio Powder X-ray Diffraction Analysis and Other Techniques. <i>Journal of Physical Chemistry C</i> , 2010, 114, 580-586.	3.1	42
26	Water sorption and near IR spectroscopy to study the differences between microcrystalline cellulose and silicified microcrystalline cellulose before and after wet granulation. <i>International Journal of Pharmaceutics</i> , 1999, 181, 41-47.	5.2	41
27	Swelling kinetics of spray-dried chitosan acetate assessed by magnetic resonance imaging and their relation to drug release kinetics of chitosan matrix tablets. <i>European Journal of Pharmaceutics and Biopharmaceutics</i> , 2011, 77, 320-326.	4.3	41
28	Process analytical technology applied for end-point detection of pharmaceutical blending by combining two calibration-free methods: Simultaneously monitoring specific near-infrared peak intensity and moving block standard deviation. <i>Powder Technology</i> , 2011, 210, 122-131.	4.2	41
29	Amorphous ultrafine particle preparation for improvement of bioavailability of insoluble drugs: grinding characteristics of fine grinding mills. <i>International Journal of Mineral Processing</i> , 2004, 74, S165-S172.	2.6	40
30	Comparison of Molecular Mobility in the Glassy State Between Amorphous Indomethacin and Salicin Based on Spin-Lattice Relaxation Times. <i>Pharmaceutical Research</i> , 2005, 22, 797-805.	3.5	40
31	Investigation of the dynamic process during spray-drying to improve aerodynamic performance of inhalation particles. <i>International Journal of Pharmaceutics</i> , 2010, 390, 250-259.	5.2	40
32	Application and Mechanism of Inhalation Profile Improvement of DPI Formulations by Mechanofusion with Magnesium Stearate. <i>Chemical and Pharmaceutical Bulletin</i> , 2008, 56, 617-625.	1.3	38
33	Evaluation of amorphous ursodeoxycholic acid by thermal methods. <i>Pharmaceutical Research</i> , 1997, 14, 798-803.	3.5	36
34	Differences in crystallization behavior between quenched and ground amorphous ursodeoxycholic acid. <i>Pharmaceutical Research</i> , 1999, 16, 835-840.	3.5	36
35	Isostructural Multicomponent Gliclazide Crystals with Improved Solubility. <i>Crystal Growth and Design</i> , 2016, 16, 6568-6573.	3.0	36
36	Solubility Parameter and Dissolution Behavior of Cefalexin Powders with Different Crystallinity.. <i>Chemical and Pharmaceutical Bulletin</i> , 1992, 40, 819-820.	1.3	35

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37	Effects of water content in physical mixture and heating temperature on crystallinity of troglitazone-PVP K30 solid dispersions prepared by closed melting method. <i>International Journal of Pharmaceutics</i> , 2005, 302, 103-112.	5.2	35
38	siRNA delivery to lung-metastasized tumor by systemic injection with cationic liposomes. <i>Journal of Liposome Research</i> , 2015, 25, 279-286.	3.3	35
39	Physicochemical properties and surface free energy of ground talc. <i>Solid State Ionics</i> , 2004, 172, 459-462.	2.7	34
40	Development of a method for nondestructive NIR transmittance spectroscopic analysis of acetaminophen and caffeine anhydrate in intact bilayer tablets. <i>Journal of Pharmaceutical and Biomedical Analysis</i> , 2010, 53, 396-402.	2.8	34
41	Development of a method for the determination of caffeine anhydrate in various designed intact tables by near-infrared spectroscopy: A comparison between reflectance and transmittance technique. <i>Journal of Pharmaceutical and Biomedical Analysis</i> , 2008, 47, 819-827.	2.8	33
42	Crystal Structure Determination of Dimenhydrinate after More than 60 Years: Solving Saltâ€™Cocrystal Ambiguity via Solid-State Characterizations and Solubility Study. <i>Crystal Growth and Design</i> , 2016, 16, 5223-5229.	3.0	33
43	Reevaluation of solubility of tolbutamide and polymorphic transformation from Form I to unknown crystal form. <i>International Journal of Pharmaceutics</i> , 2009, 369, 12-18.	5.2	32
44	Mechanism of Dehydrationâ€™Hydration Processes of Lisinopril Dihydrate Investigated by ab Initio Powder X-ray Diffraction Analysis. <i>Crystal Growth and Design</i> , 2012, 12, 6165-6172.	3.0	32
45	Enhanced Dissolution of Ursodeoxycholic Acid from the Solid Dispersion. <i>Drug Development and Industrial Pharmacy</i> , 1997, 23, 1115-1121.	2.0	31
46	In vivo siRNA delivery system for targeting to the liver by poly-l-glutamic acid-coated lipoplex. <i>Results in Pharma Sciences</i> , 2014, 4, 1-7.	4.2	30
47	Quantitative correlation between initial dissolution rate and heat of solution of drug. <i>Pharmaceutical Research</i> , 2000, 17, 920-924.	3.5	28
48	Characterization of Amorphous Ursodeoxycholic Acid Prepared by Spray-drying. <i>Journal of Pharmacy and Pharmacology</i> , 2011, 50, 1213-1219.	2.4	28
49	Simultaneous Improvement of Epalrestat Photostability and Solubility via Cocrystallization: A Case Study. <i>Crystal Growth and Design</i> , 2018, 18, 373-379.	3.0	28
50	New methods for preparing cyclodextrin inclusion compounds. III. Preparation of heptakis-(2,6-di-O-methyl)-BETA.-cyclodextrin-benzoic acid inclusion compound by sealed heating.. <i>Chemical and Pharmaceutical Bulletin</i> , 1990, 38, 1345-1348.	1.3	27
51	Effect of the type of lubricant on the characteristics of orally disintegrating tablets manufactured using the phase transition of sugar alcohol. <i>European Journal of Pharmaceutics and Biopharmaceutics</i> , 2008, 69, 986-992.	4.3	27
52	Solubility Improvement of Benexate through Salt Formation Using Artificial Sweetener. <i>Pharmaceutics</i> , 2018, 10, 64.	4.5	27
53	Improving mechanical properties of desloratadine via multicomponent crystal formation. <i>European Journal of Pharmaceutical Sciences</i> , 2018, 111, 65-72.	4.0	26
54	Polymorphism of Tegafur: Physico-chemical Properties of Four Polymorphs.. <i>Chemical and Pharmaceutical Bulletin</i> , 1993, 41, 1632-1635.	1.3	24

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55	Quantitative correlation between initial dissolution rate and heat of fusion of drug substance. <i>International Journal of Pharmaceutics</i> , 2000, 204, 1-6.	5.2	24
56	Evaluation of the physical stability and local crystallization of amorphous terfenadine using XRD and DSC and micro-TA. <i>Thermochimica Acta</i> , 2005, 432, 70-75.	2.7	24
57	siRNA Delivery into Tumor Cells by Cationic Cholesterol Derivative-Based Nanoparticles and Liposomes. <i>Biological and Pharmaceutical Bulletin</i> , 2015, 38, 30-38.	1.4	24
58	Effect of grinding on dehydration of crystal water of theophylline.. <i>Chemical and Pharmaceutical Bulletin</i> , 1990, 38, 2233-2236.	1.3	23
59	New Methods for Preparing Cyclodextrin Inclusion Compounds. IV. Enhancement of Combining Molar Ratio by Using a Ground Mixture in Heptakis-(2,6-di-O-methyl)-.BETA.-cyclodextrin and Benzoic Acid System.. <i>Chemical and Pharmaceutical Bulletin</i> , 1991, 39, 1532-1535.	1.3	23
60	Physicochemical characteristics of porous crystalline cellulose and formation of an amorphous state of ethenzamide by mixing. <i>International Journal of Pharmaceutics</i> , 1994, 108, 167-172.	5.2	23
61	Fluorometric Studies of Pyrene Adsorption on Porous Crystalline Cellulose. <i>Journal of Colloid and Interface Science</i> , 1998, 205, 510-515.	9.4	23
62	Effect of Pore Size on the Gaseous Adsorption of Ethenzamide on Porous Crystalline Cellulose and the Physicochemical Stability of Ethenzamide after Storage.. <i>Chemical and Pharmaceutical Bulletin</i> , 1998, 46, 314-318.	1.3	22
63	Importance of excipient wettability on tablet characteristics prepared by moisture activated dry granulation (MADG). <i>International Journal of Pharmaceutics</i> , 2013, 456, 58-64.	5.2	22
64	Molecular Properties of Propranolol Hydrochloride Prepared as Drug-Resin Complexes. <i>Drug Development and Industrial Pharmacy</i> , 2001, 27, 359-364.	2.0	21
65	Glass-State Amorphous Salt Solids Formed by Freeze-Drying of Amines and Hydroxy Carboxylic Acids: Effect of Hydrogen-Bonding and Electrostatic Interactions. <i>Chemical and Pharmaceutical Bulletin</i> , 2008, 56, 821-826.	1.3	21
66	Competition of Thermodynamic and Dynamic Factors During Formation of Multicomponent Particles via Spray Drying. <i>Journal of Pharmaceutical Sciences</i> , 2013, 102, 518-529.	3.3	21
67	Evaluation of the crystalline and amorphous states of drug products by nanothermal analysis and Raman imaging. <i>Journal of Pharmaceutical and Biomedical Analysis</i> , 2013, 75, 105-111.	2.8	21
68	Studying the Morphology of Lyophilized Protein Solids Using X-ray Micro-CT: Effect of Post-freeze Annealing and Controlled Nucleation. <i>AAPS PharmSciTech</i> , 2014, 15, 1181-1188.	3.3	21
69	Freeze-drying of drug-additive binary systems III. Crystallization of β -cyclodextrin inclusion complex in freezing process. <i>International Journal of Pharmaceutics</i> , 1990, 61, 27-34.	5.2	20
70	Factors affecting the apparent solubility of ursodeoxycholic acid in the grinding process. <i>International Journal of Pharmaceutics</i> , 2003, 255, 49-56.	5.2	20
71	Applying terahertz technology for nondestructive detection of crack initiation in a film-coated layer on a swelling tablet. <i>Results in Pharma Sciences</i> , 2012, 2, 29-37.	4.2	20
72	Physicochemical and crystal structure analysis of pranlukast pseudo-polymorphs II: Solvate and cocrystal. <i>Journal of Pharmaceutical and Biomedical Analysis</i> , 2015, 111, 44-50.	2.8	20

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73	Application of Eudragit RS to Thermo-Sensitive Drug Delivery Systems. I. Thermo-Sensitive Drug Release from Acetaminophen Matrix Tablets Consisting of Eudragit RS/PEG 400 Blend Polymers.. Chemical and Pharmaceutical Bulletin, 2002, 50, 408-412.	1.3	19
74	Polymorphic and pseudomorphic transformation behavior of acyclovir based on thermodynamics and crystallography. Journal of Thermal Analysis and Calorimetry, 2013, 113, 1261-1267.	3.6	19
75	Stability of aspirin in controlled pore glass solid dispersions.. Chemical and Pharmaceutical Bulletin, 1991, 39, 1027-1031.	1.3	18
76	Characterization of Polymorphs of a Novel Quinolinone Derivative, TA-270 (4-Hydroxy-1-methyl-3-octyloxy-7-sinapinoylamino-2(1H)-quinolinone).. Chemical and Pharmaceutical Bulletin, 2001, 49, 1321-1325.	1.3	18
77	Estimation of Initial Dissolution Rate of Drug Substance by Thermal Analysis: Application for Carbamazepine Hydrate. Pharmaceutical Development and Technology, 2002, 7, 89-95.	2.4	18
78	Changes in surface properties by granulation and physicochemical stability of granulated amorphous cefditoren pivoxil with additives. International Journal of Pharmaceutics, 2004, 280, 67-75.	5.2	17
79	Stabilization of Protein Structure in Freeze-Dried Amorphous Organic Acid Buffer Salts. Chemical and Pharmaceutical Bulletin, 2009, 57, 1231-1236.	1.3	17
80	Determination for dry layer resistance of sucrose under various primary drying conditions using a novel simulation program for designing pharmaceutical lyophilization cycle. International Journal of Pharmaceutics, 2013, 452, 180-187.	5.2	17
81	Sequential intravenous injection of anionic polymer and cationic lipoplex of siRNA could effectively deliver siRNA to the liver. International Journal of Pharmaceutics, 2014, 476, 289-298.	5.2	17
82	Thermal behavior of ground mixtures of heptakis (2,6-di-O-methyl)- β -cyclodextrin and benzoic acid. Journal of Inclusion Phenomena and Macrocyclic Chemistry, 1993, 15, 91-101.	1.6	16
83	Physicochemical Properties of Amorphous Ursodeoxycholic Acid Obtained by Grinding.. Chemical and Pharmaceutical Bulletin, 1994, 42, 1948-1950.	1.3	16
84	Clarifying the mechanism of aggregation of particles in high-shear granulation based on their surface properties by using micro-spectroscopy. International Journal of Pharmaceutics, 2014, 461, 495-504.	5.2	16
85	Optimization of Primary Drying Condition for Pharmaceutical Lyophilization Using a Novel Simulation Program with a Predictive Model for Dry Layer Resistance. Chemical and Pharmaceutical Bulletin, 2014, 62, 153-159.	1.3	15
86	Therapeutic effect for liver-metastasized tumor by sequential intravenous injection of anionic polymer and cationic lipoplex of siRNA. Journal of Drug Targeting, 2016, 24, 309-317.	4.4	15
87	Freeze-drying of drug-additive binary systems. II. Relationship between decarboxylation behavior and molecular states of p-aminosalicylic acid.. Chemical and Pharmaceutical Bulletin, 1989, 37, 3088-3091.	1.3	14
88	Physicochemical Properties of Ursodeoxycholic Acid Dispersed in Controlled Pore Glass. Journal of Colloid and Interface Science, 1999, 216, 276-284.	9.4	14
89	Specific complexation of ursodeoxycholic acid with guest compounds induced by co-grinding. Physical Chemistry Chemical Physics, 2000, 2, 2815-2820.	2.8	14
90	Solid-State Fluorescence Study of Naphthalene Adsorption on Porous Material. Journal of Colloid and Interface Science, 2002, 248, 239-243.	9.4	14

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91	Practical Approach for Measuring Heat Capacity of Pharmaceutical Crystals/Glasses by Modulated-Temperature Differential Scanning Calorimetry. <i>Chemical and Pharmaceutical Bulletin</i> , 2013, 61, 315-319.	1.3	14
92	Zoledronic acid enhances antitumor efficacy of liposomal doxorubicin. <i>International Journal of Oncology</i> , 2015, 47, 211-219.	3.3	14
93	Enhanced dissolution and skin permeation profiles of epalrestat with β -cyclodextrin derivatives using a cogrinding method. <i>European Journal of Pharmaceutical Sciences</i> , 2017, 106, 79-86.	4.0	14
94	Structural origin of physicochemical properties differences upon dehydration and polymorphic transformation of ciprofloxacin hydrochloride revealed by structure determination from powder X-ray diffraction data. <i>CrystEngComm</i> , 2020, 22, 7272-7279.	2.6	14
95	Molecular State of Chlorpheniramine in Resinates.. <i>Chemical and Pharmaceutical Bulletin</i> , 2000, 48, 231-234.	1.3	13
96	Prediction of the induction period of crystallization of naproxen in solid dispersion using differential scanning calorimetry. <i>Journal of Thermal Analysis and Calorimetry</i> , 2010, 99, 15-19.	3.6	13
97	The importance of binder moisture content in Metformin HCL high-dose formulations prepared by moist aqueous granulation (MAG). <i>Results in Pharma Sciences</i> , 2015, 5, 1-7.	4.2	13
98	Mechanisms for Improved Hygroscopicity of L-Arginine Valproate Revealed by X-Ray Single Crystal Structure Analysis. <i>Journal of Pharmaceutical Sciences</i> , 2017, 106, 859-865.	3.3	13
99	Fluorometric study of the molecular states of 2,5-diphenyloxazole in ground mixtures with gamma-cyclodextrin. <i>Pharmaceutical Research</i> , 1994, 11, 331-336.	3.5	12
100	Uniformity and physical states of troglitazone in solid dispersions determined by electron probe microanalysis and microthermal analysis. <i>International Journal of Pharmaceutics</i> , 2004, 280, 39-46.	5.2	12
101	Effects of Grinding and Humidification on the Transformation of Conglomerate to Racemic Compound in Optically Active Drugs. <i>Journal of Pharmacy and Pharmacology</i> , 2011, 49, 384-389.	2.4	12
102	Crystallographic Analysis of Phase Dissociation Related to Anomalous Solubility of Irsogladine Maleate. <i>Crystal Growth and Design</i> , 2016, 16, 6714-6718.	3.0	12
103	Characterization of complexes between phenethylamine enantiomers and β -cyclodextrin derivatives by capillary electrophoresis—Determination of binding constants and complex mobilities. <i>Electrophoresis</i> , 2017, 38, 1188-1200.	2.4	12
104	Peculiar Peak Shifts in the IR Spectrum of Benzoic Acid Crystals by Compression with Methylated Additives.. <i>Chemical and Pharmaceutical Bulletin</i> , 1995, 43, 666-670.	1.3	11
105	Complex Formation between Deoxycholic Acid and Menadione by Grinding and Sealed Heating Methods.. <i>Chemical and Pharmaceutical Bulletin</i> , 1997, 45, 1358-1362.	1.3	11
106	Quantitative relationship between solubility, initial dissolution rate and heat of solution of chiral drugs. <i>Pharmaceutical Research</i> , 2000, 17, 90-93.	3.5	11
107	The effect of water activity on granule characteristics and tablet properties produced by moisture activated dry granulation (MADG). <i>Powder Technology</i> , 2016, 294, 113-118.	4.2	11
108	Novel, lean and environment-friendly granulation method: Green fluidized bed granulation (GFBG). <i>International Journal of Pharmaceutics</i> , 2019, 557, 18-25.	5.2	11

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109	Acceleration of the addition reaction of succinic anhydride and p-nitroaniline in controlled-pore glass solid dispersions.. Chemical and Pharmaceutical Bulletin, 1989, 37, 3083-3087.	1.3	10
110	Molecular States of 2-Naphthoic Acid in Solid Dispersions with Porous Crystalline Cellulose, as Investigated by Fluorescence Spectroscopy. Bulletin of the Chemical Society of Japan, 2000, 73, 1567-1572.	3.2	10
111	Potential of synchrotron X-ray powder diffractometry for detection and quantification of small amounts of crystalline drug substances in pharmaceutical tablets. Journal of Pharmaceutical and Biomedical Analysis, 2011, 56, 448-453.	2.8	10
112	Tumor delivery of liposomal doxorubicin prepared with poly-L-glutamic acid as a drug-trapping agent. Journal of Liposome Research, 2017, 27, 99-107.	3.3	10
113	Stabilization mechanism of amorphous carbamazepine by transglycosylated rutin, a non-polymeric amorphous additive with a high glass transition temperature. International Journal of Pharmaceutics, 2021, 600, 120491.	5.2	10
114	Thermal Behavior of Methyl p-hydroxybenzoate in Controlled-Pore Glass Solid Dispersion. Journal of Colloid and Interface Science, 1995, 173, 186-191.	9.4	9
115	Dissolution studies in organic solvents for evaluating hydrogen-bond matrix of cellulose in the ground mixture. International Journal of Pharmaceutics, 1995, 113, 97-102.	5.2	9
116	Evaluation of physicochemical properties on the blending process of pharmaceutical granules with magnesium stearate by thermal effusivity sensor. Journal of Thermal Analysis and Calorimetry, 2013, 113, 1281-1285.	3.6	9
117	Evaluation of the physical properties of dry surface-modified ibuprofen using a powder rheometer (FT4) and analysis of the influence of pharmaceutical additives on improvement of the powder flowability. International Journal of Pharmaceutics, 2020, 579, 119165.	5.2	9
118	Miscibility characterization of zein/methacrylic acid copolymer composite films and plasticization effects. International Journal of Pharmaceutics, 2021, 601, 120498.	5.2	9
119	Chitosan film containing antifungal agent-loaded SLNs for the treatment of candidiasis using a Box-Behnken design. Carbohydrate Polymers, 2022, 283, 119178.	10.2	9
120	Complex formation between naphthalene and dimethyl- β -cyclodextrin by heating in a sealed ampoule. Journal of the Chemical Society, Faraday Transactions, 1994, 90, 3117-3119.	1.7	8
121	Interaction of Microcrystalline Cellulose and Water in Granules Prepared by a High-Shear Mixer.. Chemical and Pharmaceutical Bulletin, 2001, 49, 373-378.	1.3	8
122	Specific Complexation of Ursodeoxycholic Acid with Guest Compounds Induced by Co-grinding. II. Effect of Grinding Temperature on the Mechanochemical Complexation. Bulletin of the Chemical Society of Japan, 2003, 76, 515-521.	3.2	8
123	Microscopic molecular mobility of amorphous AG-041R measured by solid-state ^{13}C NMR. International Journal of Pharmaceutics, 2004, 275, 73-83.	5.2	8
124	Effect of physical properties of troglitazone crystal on the molecular interaction with PVP during heating. International Journal of Pharmaceutics, 2007, 336, 82-89.	5.2	8
125	Applicability of DPI formulations for novel neurokinin receptor antagonist. International Journal of Pharmaceutics, 2008, 356, 102-109.	5.2	8
126	Low-Density Microparticles with Petaloid Surface Structure for Pulmonary Drug Delivery. Journal of Pharmaceutical Sciences, 2014, 103, 1309-1313.	3.3	8

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127	A 1:1 Deoxycholic Acid-Salicylic Acid Complex. <i>Acta Crystallographica Section C: Crystal Structure Communications</i> , 1997, 53, 803-805.	0.4	7
128	Effects of Solute Miscibility on the Micro- and Macroscopic Structural Integrity of Freeze-Dried Solids. <i>Journal of Pharmaceutical Sciences</i> , 2010, 99, 4710-4719.	3.3	7
129	Study of the Pseudo-Crystalline Transformation from Form I to Form II of Thiamine Hydrochloride (Vitamin B1). <i>Chemical and Pharmaceutical Bulletin</i> , 2011, 59, 57-62.	1.3	7
130	Component Crystallization and Physical Collapse during Freeze-Drying of Arginine-Citric Acid Mixtures. <i>Chemical and Pharmaceutical Bulletin</i> , 2012, 60, 1176-1181.	1.3	7
131	Transdermal Delivery of Small Interfering RNA with Elastic Cationic Liposomes in Mice. <i>Journal of Pharmaceutics</i> , 2013, 2013, 1-6.	4.7	7
132	Effect of sulfobutyl ether- β -cyclodextrin and propylene glycol alginate on the solubility of clozapine. <i>Pharmaceutical Development and Technology</i> , 2019, 24, 479-486.	2.4	7
133	A New Method for Classification of Salts and Cocrystals Using Solid-State UV Spectrophotometry. <i>Chemical and Pharmaceutical Bulletin</i> , 2019, 67, 945-952.	1.3	7
134	Formulation design and evaluation of a transdermal drug delivery system containing a novel eptazocine salt with the Eudragit® E adhesive. <i>Journal of Drug Delivery Science and Technology</i> , 2019, 54, 101289.	3.0	7
135	Mechanochemical Complexation between Deoxycholic Acid and Salicylic Acid. <i>Journal of Inclusion Phenomena and Macrocyclic Chemistry</i> , 1998, 31, 367-379.	1.6	6
136	Effect of Guest Species on Inclusion Compound Formation of Deoxycholic Acid by Co-Grinding. <i>Bulletin of the Chemical Society of Japan</i> , 1998, 71, 1573-1579.	3.2	6
137	Near IR spectroscopy to quantify the silica content and difference between silicified microcrystalline cellulose and physical mixtures of microcrystalline cellulose and silica. <i>European Journal of Pharmaceutical Sciences</i> , 2000, 10, 77-80.	4.0	6
138	Diffusivity of amorphous drug in solid dispersion. <i>Journal of Thermal Analysis and Calorimetry</i> , 2013, 113, 1505-1510.	3.6	6
139	Effects of Formulation and Process Factors on the Crystal Structure of Freeze-Dried Myo-Inositol. <i>Journal of Pharmaceutical Sciences</i> , 2014, 103, 2347-2355.	3.3	6
140	Crystal Structure of an Epalrestat Dimethanol Solvate. <i>X-ray Structure Analysis Online</i> , 2016, 32, 7-9.	0.2	6
141	Formation Mechanism of Lipid Membrane and Vesicle Using Small Angle X-ray Scattering and Dissipative Particle Dynamics (DPD) Method. <i>Journal of Computer Chemistry Japan</i> , 2018, 17, 172-179.	0.1	6
142	Capturing a new hydrate polymorph of amodiaquine dihydrochloride dihydrate via heterogeneous crystallisation. <i>CrystEngComm</i> , 2019, 21, 2053-2057.	2.6	6
143	Importance of free water in controlling granule and tablet properties in a novel granulation method, green fluidized bed granulation (GFBC). <i>International Journal of Pharmaceutics</i> , 2019, 570, 118647.	5.2	6
144	Effect of Polymers and Storage Relative Humidity on Amorphous Rebamipide and Its Solid Dispersion Transformation: Multiple Spectra Chemometrics of Powder X-Ray Diffraction and Near-Infrared Spectroscopy. <i>Pharmaceutics</i> , 2020, 13, 147.	3.8	6

#	ARTICLE	IF	CITATIONS
145	Dose-Dependent Solubility-Permeability Interplay for Poorly Soluble Drugs under Non-Sink Conditions. <i>Pharmaceutics</i> , 2021, 13, 323.	4.5	6
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