## Etsuo Yonemochi

List of Publications by Year in descending order

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215 papers

3,881 citations

35 h-index 197818 49 g-index

222 all docs 222 docs citations

times ranked

222

3336 citing authors

#	Article	IF	CITATIONS
1	Computational approach to elucidate the formation and stabilization mechanism of amorphous formulation using molecular dynamics simulation and fragment molecular orbital calculation. International Journal of Pharmaceutics, 2022, 615, 121477.	5.2	2
2	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
3	Development of a retention prediction model in ion-pair reversed-phase HPLC for nucleoside triphosphates used as mRNA vaccine raw materials. Journal of Chromatography B: Analytical Technologies in the Biomedical and Life Sciences, 2022, 1193, 123168.	2.3	2
4	Formulation of Biopharmaceutical Dry Powder Inhaler Using the Void Forming Index (VFI) to Detect and Avoid Powder Caking in Dry Powder Inhaler Formulations. Chemical and Pharmaceutical Bulletin, 2022, 70, 245-253.	1.3	1
5	Non-Effective Improvement of Absorption for Some Nanoparticle Formulations Explained by Permeability under Non-Sink Conditions. Pharmaceutics, 2022, 14, 816.	4.5	1
6	Tablet Quality-Prediction Model Using Quality Material Attributes: Toward Flexible Switching Between Batch and Continuous Granulation. Journal of Pharmaceutical Innovation, 2021, 16, 588-602.	2.4	2
7	Understanding Crystal Cleavability and Physical Properties of Crystal Surfaces Using <i>in Silico</i> Simulation. Chemical and Pharmaceutical Bulletin, 2021, 69, 185-198.	1.3	1
8	Crystal Structure of Novel Terephthalate Salt of Antiarrhythmic Drug Disopyramide. Crystals, 2021, 11, 368.	2.2	2
9	Dose-Dependent Solubility–Permeability Interplay for Poorly Soluble Drugs under Non-Sink Conditions. Pharmaceutics, 2021, 13, 323.	4.5	6
10	Selection of Small Amounts of Glidant Capable of Improving the Tensile Strength of Ibuprofen Tablets. Chemical and Pharmaceutical Bulletin, 2021, 69, 374-382.	1.3	2
11	Crystal Structures of Antiarrhythmic Drug Disopyramide and Its Salt with Phthalic Acid. Crystals, 2021, 11, 379.	2.2	1
12	Manufacturability and Properties of Granules and Tablets Using the Eco-Friendly Granulation Method Green Fluidized Bed Granulation Compared to Direct Compression. Chemical and Pharmaceutical Bulletin, 2021, 69, 447-455.	1.3	3
13	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
14	The development of retention time prediction model using multilinear gradient profiles of seven pharmaceuticals. Journal of Pharmaceutical and Biomedical Analysis, 2021, 198, 114024.	2.8	2
15	Miscibility characterization of zein/methacrylic acid copolymer composite films and plasticization effects. International Journal of Pharmaceutics, 2021, 601, 120498.	5.2	9
16	Altered Media Flow and Tablet Position as Factors of How Air Bubbles Affect Dissolution of Disintegrating and Non-disintegrating Tablets Using a USP 4 Flow-Through Cell Apparatus. AAPS PharmSciTech, 2021, 22, 227.	3.3	2
17	Cholesteryl-Conjugated Ribonuclease A Exhibits Enzyme Activity in Aqueous Solution and Resistance to Dimethyl Sulfoxide. ACS Omega, 2021, 6, 533-543.	3.5	1
18	Improving the Accuracy of Crystal Structure Prediction Using FMO Crystal Energy: An Example of Target XXIII. Journal of Computer Chemistry Japan, 2021, 20, 92-93.	0.1	0

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19	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. Pharmaceuticals, 2020, 13, 147.	3.8	6
20	Synthesis and Characterization of Cholesteryl Conjugated Lysozyme (CHLysozyme). Molecules, 2020, 25, 3704.	3.8	2
21	Application of void forming index (VFI): Detection of the effect of physical properties of dry powder inhaler formulations on powder cohesion. International Journal of Pharmaceutics, 2020, 588, 119766.	5.2	4
22	Selfâ€Degradable Lipidâ€Like Materials Based on "Hydrolysis accelerated by the intraâ€Particle Enrichment of Reactant (HyPER)―for Messenger RNA Delivery. Advanced Functional Materials, 2020, 30, 1910575.	14.9	65
23	Structural origin of physicochemical properties differences upon dehydration and polymorphic transformation of ciprofloxacin hydrochloride revealed by structure determination from powder X-ray diffraction data. CrystEngComm, 2020, 22, 7272-7279.	2.6	14
24	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
25	Preface of the Special Issue "Pharmaceutical Crystals― Crystals, 2020, 10, 89.	2.2	1
26	Crystal Structural Analysis of DL-Mandelate Salt of Carvedilol and Its Correlation with Physicochemical Properties. Crystals, 2020, 10, 53.	2.2	2
27	New approach to optimizing risk management of the sticking problem using scale-independent critical material attributes and the quantitative process parameter. International Journal of Pharmaceutics, 2020, 577, 119032.	5 <b>.</b> 2	3
28	Degradation Pathway of a Taxane Derivative DS80100717 Drug Substance and Drug Product. Chemical and Pharmaceutical Bulletin, 2020, 68, 392-397.	1.3	1
29	Novel approach to evaluating granulation and segregation level considering the contribution of hydroxypropyl cellulose to the surface property change of granules. International Journal of Pharmaceutics, 2020, 581, 119254.	<b>5.</b> 2	2
30	Impact of Magnesium Stearate Content: Modeling of Drug Degradation Using a Modified Arrhenius Equation. Chemical and Pharmaceutical Bulletin, 2020, 68, 1049-1054.	1.3	4
31	Effect of sulfobutyl ether- $\hat{l}^2$ -cyclodextrin and propylene glycol alginate on the solubility of clozapine. Pharmaceutical Development and Technology, 2019, 24, 479-486.	2.4	7
32	A New Method for Classification of Salts and Cocrystals Using Solid-State UV Spectrophotometry. Chemical and Pharmaceutical Bulletin, 2019, 67, 945-952.	1.3	7
33	Foreword. Chemical and Pharmaceutical Bulletin, 2019, 67, 904-905.	1.3	0
34	Formulation design and evaluation of a transdermal drug delivery system containing a novel eptazocine salt with the Eudragit® E adhesive. Journal of Drug Delivery Science and Technology, 2019, 54, 101289.	3.0	7
35	Capturing a new hydrate polymorph of amodiaquine dihydrochloride dihydrate via heterogeneous crystallisation. CrystEngComm, 2019, 21, 2053-2057.	2.6	6
36	Development of microparticles coated with poly-Î <sup>3</sup> -glutamic acid to improve oral absorption of a poorly water-soluble drug. Pharmaceutical Development and Technology, 2019, 24, 992-1001.	2.4	4

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37	Importance of free water in controlling granule and tablet properties in a novel granulation method, green fluidized bed granulation (GFBG). International Journal of Pharmaceutics, 2019, 570, 118647.	5.2	6
38	Novel, lean and environment-friendly granulation method: Green fluidized bed granulation (GFBG). International Journal of Pharmaceutics, 2019, 557, 18-25.	5.2	11
39	Simultaneous Improvement of Epalrestat Photostability and Solubility via Cocrystallization: A Case Study. Crystal Growth and Design, 2018, 18, 373-379.	3.0	28
40	Improving mechanical properties of desloratadine via multicomponent crystal formation. European Journal of Pharmaceutical Sciences, 2018, 111, 65-72.	4.0	26
41	Ethyl Haematommate fromStereocaulon graminosumSchaer.: Isolation and Crystal Structure. Natural Product Sciences, 2018, 24, 115.	0.9	1
42	Formation Mechanism of Lipid Membrane and Vesicle Using Small Angle X-ray Scattering and Dissipative Particle Dynamics (DPD) Method. Journal of Computer Chemistry Japan, 2018, 17, 172-179.	0.1	6
43	Solubility Improvement of Benexate through Salt Formation Using Artificial Sweetener. Pharmaceutics, 2018, 10, 64.	4.5	27
44	Tumor delivery of liposomal doxorubicin prepared with poly- <scp>L</scp> -glutamic acid as a drug-trapping agent. Journal of Liposome Research, 2017, 27, 99-107.	3.3	10
45	Characterization of complexes between phenethylamine enantiomers and βâ€eyclodextrin derivatives by capillary electrophoresis—Determination of binding constants and complex mobilities. Electrophoresis, 2017, 38, 1188-1200.	2.4	12
46	Enhanced dissolution and skin permeation profiles of epalrestat with $\hat{l}^2$ -cyclodextrin derivatives using a cogrinding method. European Journal of Pharmaceutical Sciences, 2017, 106, 79-86.	4.0	14
47	Solubility improvement of epalrestat by layered structure formation via cocrystallization. CrystEngComm, 2017, 19, 2614-2622.	2.6	45
48	Molecular Dynamics of Amorphous Sulfamethazine With Structurally Related Sulfonamide Impurities Evaluated Using Thermal Analysis. Journal of Pharmaceutical Sciences, 2017, 106, 1062-1068.	3.3	4
49	Mechanisms for Improved Hygroscopicity of L-Arginine Valproate Revealed by X-Ray Single Crystal Structure Analysis. Journal of Pharmaceutical Sciences, 2017, 106, 859-865.	3.3	13
50	Void forming index: A new parameter for detecting microstructural transformation caused by powder agglomeration. International Journal of Pharmaceutics, 2017, 532, 118-123.	5.2	3
51	Investigation of Discoloration of Furosemide Tablets in a Light-Shielded Environment. Chemical and Pharmaceutical Bulletin, 2017, 65, 373-380.	1.3	2
52	Effect of Magnesium Stearate Mono- and Dihydrate Dispersibilities on Physical Properties of Tablets. Chemical and Pharmaceutical Bulletin, 2017, 65, 1028-1034.	1.3	0
53	Epalrestat tetrahydrofuran monosolvate: crystal structure and phase transition. Acta Crystallographica Section E: Crystallographic Communications, 2017, 73, 941-944.	0.5	3
54	A new solvate of epalerstat, a drug for diabetic neuropathy. Acta Crystallographica Section E: Crystallographic Communications, 2017, 73, 1264-1267.	0.5	2

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55	Evaluation of the Manufacturing Process for Pharmaceuticals by Using Micro-imaging Analysis. Oleoscience, 2017, 17, 379-385.	0.0	O
56	General understanding on physical stability of pharmaceutical glasses. Asian Journal of Pharmaceutical Sciences, 2016, 11, 54-55.	9.1	0
57	The effect of structurally related impurities on crystallinity reduction of sulfamethazine by grinding. International Journal of Pharmaceutics, 2016, 515, 416-421.	5.2	5
58	Isostructural Multicomponent Gliclazide Crystals with Improved Solubility. Crystal Growth and Design, 2016, 16, 6568-6573.	3.0	36
59	Crystal Structure Determination of Dimenhydrinate after More than 60 Years: Solving Salt–Cocrystal Ambiguity via Solid-State Characterizations and Solubility Study. Crystal Growth and Design, 2016, 16, 5223-5229.	3.0	33
60	Drug-drug Multi-component Crystal of Acedoben–Dimepranol 2:1. X-ray Structure Analysis Online, 2016, 32, 39-40.	0.2	1
61	Crystal Structure of an Epalrestat Dimethanol Solvate. X-ray Structure Analysis Online, 2016, 32, 7-9.	0.2	6
62	Characterization and Quality Control of Pharmaceutical Cocrystals. Chemical and Pharmaceutical Bulletin, 2016, 64, 1421-1430.	1.3	46
63	Crystallographic Analysis of Phase Dissociation Related to Anomalous Solubility of Irsogladine Maleate. Crystal Growth and Design, 2016, 16, 6714-6718.	3.0	12
64	Drug–Drug Multicomponent Crystals as an Effective Technique to Overcome Weaknesses in Parent Drugs. Crystal Growth and Design, 2016, 16, 3577-3581.	3.0	52
65	The effect of water activity on granule characteristics and tablet properties produced by moisture activated dry granulation (MADG). Powder Technology, 2016, 294, 113-118.	4.2	11
66	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
67	The importance of binder moisture content in Metformin HCL high-dose formulations prepared by moist aqueous granulation (MAG). Results in Pharma Sciences, 2015, 5, 1-7.	4.2	13
68	siRNA Delivery into Tumor Cells by Cationic Cholesterol Derivative-Based Nanoparticles and Liposomes. Biological and Pharmaceutical Bulletin, 2015, 38, 30-38.	1.4	24
69	Zoledronic acid enhances antitumor efficacy of liposomal doxorubicin. International Journal of Oncology, 2015, 47, 211-219.	3.3	14
70	Physicochemical and crystal structure analysis of pranlukast pseudo-polymorphs I: Anhydrates and hydrate. Journal of Pharmaceutical and Biomedical Analysis, 2015, 107, 11-16.	2.8	5
71	siRNA delivery to lung-metastasized tumor by systemic injection with cationic liposomes. Journal of Liposome Research, 2015, 25, 279-286.	3.3	35
72	Physicochemical and crystal structure analysis of pranlukast pseudo-polymorphs II: Solvate and cocrystal. Journal of Pharmaceutical and Biomedical Analysis, 2015, 111, 44-50.	2.8	20

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73	Correlation between Glass-Forming Ability and Fragility of Pharmaceutical Compounds. Journal of Physical Chemistry B, 2015, 119, 4873-4880.	2.6	51
74	Triboelectrification of active pharmaceutical ingredients: week acids and their salts. International Journal of Pharmaceutics, 2015, 493, 434-438.	<b>5.</b> 2	1
75	Evaluation of antitumor effect of zoledronic acid entrapped in folate-linked liposome for targeting to tumor-associated macrophages. Journal of Liposome Research, 2015, 25, 131-140.	3.3	44
76	In vivo siRNA delivery system for targeting to the liver by poly-l-glutamic acid-coated lipoplex. Results in Pharma Sciences, 2014, 4, 1-7.	4.2	30
77	Low-Density Microparticles with Petaloid Surface Structure for Pulmonary Drug Delivery. Journal of Pharmaceutical Sciences, 2014, 103, 1309-1313.	3.3	8
78	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.	<b>5.2</b>	17
79	Relationship between Crystallization Tendencies during Cooling from Melt and Isothermal Storage: Toward a General Understanding of Physical Stability of Pharmaceutical Glasses. Molecular Pharmaceutics, 2014, 11, 1835-1843.	4.6	48
80	Studying the Morphology of Lyophilized Protein Solids Using X-ray Micro-CT: Effect of Post-freeze Annealing and Controlled Nucleation. AAPS PharmSciTech, 2014, 15, 1181-1188.	3.3	21
81	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.	<b>5.2</b>	16
82	Effects of Formulation and Process Factors on the Crystal Structure of Freeze-Dried Myo-Inositol. Journal of Pharmaceutical Sciences, 2014, 103, 2347-2355.	3.3	6
83	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
84	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
85	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
86	Competition of Thermodynamic and Dynamic Factors During Formation of Multicomponent Particles via Spray Drying. Journal of Pharmaceutical Sciences, 2013, 102, 518-529.	3.3	21
87	Evaluation of the crystalline and amorphous states of drug products by nanothermal analysis and Raman imaging. Journal of Pharmaceutical and Biomedical Analysis, 2013, 75, 105-111.	2.8	21
88	Importance of excipient wettability on tablet characteristics prepared by moisture activated dry granulation (MADG). International Journal of Pharmaceutics, 2013, 456, 58-64.	<b>5.</b> 2	22
89	Diffusivity of amorphous drug in solid dispersion. Journal of Thermal Analysis and Calorimetry, 2013, 113, 1505-1510.	3.6	6
90	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.	<b>5.</b> 2	17

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91	Practical Approach for Measuring Heat Capacity of Pharmaceutical Crystals/Glasses by Modulated-Temperature Differential Scanning Calorimetry. Chemical and Pharmaceutical Bulletin, 2013, 61, 315-319.	1.3	14
92	Study of Cohesive Properties of Pharmaceutical Powders for Punch Characterized by Surface Free Energy and Cohesive Property Analysis. Journal of the Society of Powder Technology, Japan, 2013, 50, 656-661.	0.1	4
93	Transdermal Delivery of Small Interfering RNA with Elastic Cationic Liposomes in Mice. Journal of Pharmaceutics, 2013, 2013, 1-6.	4.7	7
94	Determination of Surface Free Energy and Contact Angle for Hydrolyzed Shellac. Advanced Materials Research, 2012, 506, 270-273.	0.3	3
95	Component Crystallization and Physical Collapse during Freeze-Drying of <small>L</small> -Arginine–Citric Acid Mixtures. Chemical and Pharmaceutical Bulletin, 2012, 60, 1176-1181.	1.3	7
96	Applying terahertz technology for nondestructive detection of crack initiation in a film-coated layer on a swelling tablet. Results in Pharma Sciences, 2012, 2, 29-37.	4.2	20
97	Mechanism of Dehydration–Hydration Processes of Lisinopril Dihydrate Investigated by ab Initio Powder X-ray Diffraction Analysis. Crystal Growth and Design, 2012, 12, 6165-6172.	3.0	32
98	Cocrystallization and amorphization induced by drug–excipient interaction improves the physical properties of acyclovir. International Journal of Pharmaceutics, 2012, 422, 160-169.	5.2	108
99	Swelling kinetics of spray-dried chitosan acetate assessed by magnetic resonance imaging and their relation to drug release kinetics of chitosan matrix tablets. European Journal of Pharmaceutics and Biopharmaceutics, 2011, 77, 320-326.	4.3	41
100	Effects of Grinding and Humidification on the Transformation of Conglomerate to Racemic Compound in Optically Active Drugs. Journal of Pharmacy and Pharmacology, 2011, 49, 384-389.	2.4	12
101	Characterization of Amorphous Ursodeoxycholic Acid Prepared by Spray-drying. Journal of Pharmacy and Pharmacology, 2011, 50, 1213-1219.	2.4	28
102	Do Amorphous Troglitazones Prepared from Two Diastereomer-Pairs Have the Same Molecular Mobility and Crystallization Rate at the Surface?. Chemical and Pharmaceutical Bulletin, 2011, 59, 1452-1457.	1.3	3
103	Study of the Pseudo-Crystalline Transformation from Form I to Form II of Thiamine Hydrochloride (Vitamin B1). Chemical and Pharmaceutical Bulletin, 2011, 59, 57-62.	1.3	7
104	Development of a Rapid Process Monitoring Method for Dry-Coated Tableting Process by Using Near-Infrared Spectroscopy. Chemical and Pharmaceutical Bulletin, 2011, 59, 868-873.	1.3	4
105	Evaluation of the Change in Surface Properties of Particles Induced by Mechanofusion Process. Journal of the Society of Powder Technology, Japan, 2011, 48, 618-624.	0.1	1
106	Molecular States of p-Dimethylaminobenzonitrile Coground with .BETACyclodextrin Investigated Using Solid-State Fluorescence Spectroscopy. Chemical and Pharmaceutical Bulletin, 2011, 59, 1299-1302.	1.3	3
107	Change of Molecular States of Drug by Ground with Cyclodextrin. Journal of the Society of Powder Technology, Japan, 2011, 48, 612-617.	0.1	0
108	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

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109	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. Powder Technology, 2011, 210, 122-131.	4.2	41
110	Investigation of the dynamic process during spray-drying to improve aerodynamic performance of inhalation particles. International Journal of Pharmaceutics, 2010, 390, 250-259.	5.2	40
111	Prediction of the induction period of crystallization of naproxen in solid dispersion using differential scanning calorimetry. Journal of Thermal Analysis and Calorimetry, 2010, 99, 15-19.	3.6	13
112	Effects of Solute Miscibility on the Micro- and Macroscopic Structural Integrity of Freeze-Dried Solids. Journal of Pharmaceutical Sciences, 2010, 99, 4710-4719.	3.3	7
113	Development of a method for nondestructive NIR transmittance spectroscopic analysis of acetaminophen and caffeine anhydrate in intact bilayer tablets. Journal of Pharmaceutical and Biomedical Analysis, 2010, 53, 396-402.	2.8	34
114	Freeze-drying of proteins with glass-forming oligosaccharide-derived sugar alcohols. International Journal of Pharmaceutics, 2010, 389, 107-113.	5.2	61
115	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. Journal of Physical Chemistry C, 2010, 114, 580-586.	3.1	42
116	Reevaluation of solubility of tolbutamide and polymorphic transformation from Form I to unknown crystal form. International Journal of Pharmaceutics, 2009, 369, 12-18.	5.2	32
117	Design of Highly Dispersive Particles for Pulmonary Drug Delivery. Journal of the Society of Powder Technology, Japan, 2009, 46, 698-703.	0.1	0
118	Stabilization of Protein Structure in Freeze-Dried Amorphous Organic Acid Buffer Salts. Chemical and Pharmaceutical Bulletin, 2009, 57, 1231-1236.	1.3	17
119	Freeze-Drying of Proteins in Glass Solids Formed by Basic Amino Acids and Dicarboxylic Acids. Chemical and Pharmaceutical Bulletin, 2009, 57, 43-48.	1.3	46
120	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. Journal of Pharmaceutical and Biomedical Analysis, 2008, 47, 819-827.	2.8	33
121	Effect of grinding on the dehydration behavior of nedocromil sodium hydrates. Journal of Thermal Analysis and Calorimetry, 2008, 92, 471-476.	3.6	1
122	Evaluation of dispersion state of the two racemic compounds of troglitazone in pharmaceutical granules using IR-to-THz imaging. Infrared Physics and Technology, 2008, 51, 450-453.	2.9	5
123	Applicability of DPI formulations for novel neurokinin receptor antagonist. International Journal of Pharmaceutics, 2008, 356, 102-109.	5.2	8
124	Evaluation of solid dispersions on a molecular level by the Raman mapping technique. International Journal of Pharmaceutics, 2008, 361, 12-18.	5.2	61
125	Effect of the type of lubricant on the characteristics of orally disintegrating tablets manufactured using the phase transition of sugar alcohol. European Journal of Pharmaceutics and Biopharmaceutics, 2008, 69, 986-992.	4.3	27
126	Application and Mechanism of Inhalation Profile Improvement of DPI Formulations by Mechanofusion with Magnesium Stearate. Chemical and Pharmaceutical Bulletin, 2008, 56, 617-625.	1.3	38

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127	Glass-State Amorphous Salt Solids Formed by Freeze-Drying of Amines and Hydroxy Carboxylic Acids: Effect of Hydrogen-Bonding and Electrostatic Interactions. Chemical and Pharmaceutical Bulletin, 2008, 56, 821-826.	1.3	21
128	Importance of Physicochemical Characterization of Pharmaceuticals. Journal of the Japan Society of Colour Material, 2008, 81, 48-53.	0.1	0
129	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
130	Novel Approach to DPI Carrier Lactose with Mechanofusion Process with Additives and Evaluation by IGC. Chemical and Pharmaceutical Bulletin, 2006, 54, 1508-1514.	1.3	58
131	Evaluation of rapidly disintegrating tablets containing glycine and carboxymethylcellulose. International Journal of Pharmaceutics, 2006, 310, 101-109.	5.2	92
132	Solid-state 13C NMR study of indomethacin polymorphism. International Journal of Pharmaceutics, 2006, 318, 146-153.	5.2	50
133	Application of XRD-DSC system to the optimization of manufacturing process for the freeze-dried pharmaceuticals. Journal of Thermal Analysis and Calorimetry, 2006, 85, 693-697.	3.6	4
134	Estimation of physical stability of amorphous solid dispersion using differential scanning calorimetry. Journal of Thermal Analysis and Calorimetry, 2006, 85, 689-692.	3.6	42
135	Application of microcalorimetry to the formulation study. Journal of Thermal Analysis and Calorimetry, 2006, 85, 675-680.	3.6	4
136	Mechanism of glass ampoule breakage prevention during the freeze-drying process of sodium thiopental lyophilization products on addition of sodium chloride. Journal of Thermal Analysis and Calorimetry, 2006, 85, 731-739.	3.6	4
137	Application of NIR Spectroscopy for Evaluation of Crystalline State in Granulation and Tabletting Process. Journal of the Society of Powder Technology, Japan, 2005, 42, 632-637.	0.1	0
138	Development of Fast Disintegrating Compressed Tablets Using Amino Acid as Disintegratation Accelerator: Evaluation of Wetting and Disintegration of Tablet on the Basis of Surface Free Energy. Chemical and Pharmaceutical Bulletin, 2005, 53, 1536-1539.	1.3	43
139	Effects of sugar ester and hydroxypropyl methylcellulose on the physicochemical stability of amorphous cefditoren pivoxil in aqueous suspension. International Journal of Pharmaceutics, 2005, 290, 91-99.	5.2	44
140	Effects of water content in physical mixture and heating temperature on crystallinity of troglitazone-PVP K30 solid dispersions prepared by closed melting method. International Journal of Pharmaceutics, 2005, 302, 103-112.	5.2	35
141	Formulation design of a novel fast-disintegrating tablet. International Journal of Pharmaceutics, 2005, 306, 83-90.	5.2	105
142	Investigation of optimal manufacturing process for freeze-dried formulations: Observation of frozen solutions by low temperature X-ray diffraction measurements. Thermochimica Acta, 2005, 431, 127-132.	2.7	1
143	Evaluation of the physical stability and local crystallization of amorphous terfenadine using XRD–DSC and micro-TA. Thermochimica Acta, 2005, 432, 70-75.	2.7	24
144	Application of Eudragit RS to thermo-sensitive drug delivery systems. Journal of Controlled Release, 2005, 102, 49-57.	9.9	43

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145	Comparison of Molecular Mobility in the Glassy State Between Amorphous Indomethacin and Salicin Based on Spin-Lattice Relaxation Times. Pharmaceutical Research, 2005, 22, 797-805.	3.5	40
146	Measurement of surface glass transition temperature of amorphous cefditoren pivoxil granules by inverse gas chromatography. Journal of Drug Delivery Science and Technology, 2005, 15, 439-442.	3.0	3
147	Physicochemical properties and surface free energy of ground talc. Solid State Ionics, 2004, 172, 459-462.	2.7	34
148	Microscopic molecular mobility of amorphous AG-041R measured by solid-state 13C NMR. International Journal of Pharmaceutics, 2004, 275, 73-83.	<b>5.</b> 2	8
149	Uniformity and physical states of troglitazone in solid dispersions determined by electron probe microanalysis and microthermal analysis. International Journal of Pharmaceutics, 2004, 280, 39-46.	5 <b>.</b> 2	12
150	Changes in surface properties by granulation and physicochemical stability of granulated amorphous cefditoren pivoxil with additives. International Journal of Pharmaceutics, 2004, 280, 67-75.	<b>5.</b> 2	17
151	Amorphous ultrafine particle preparation for improvement of bioavailability of insoluble drugs: grinding characteristics of fine grinding mills. International Journal of Mineral Processing, 2004, 74, S165-S172.	2.6	40
152	Change in the physicochemical properties of ursodeoxycholic acid by grinding. Solid State Ionics, 2004, 172, 455-458.	2.7	4
153	Quantitative Determination of Amorphous Nicardipine Hydrochloride in Long Acting Formula (NIC-LA) Using Light Anhydrous Silicic Acid. Chemical and Pharmaceutical Bulletin, 2004, 52, 1451-1457.	1.3	4
154	Factors affecting the apparent solubility of ursodeoxycholic acid in the grinding process. International Journal of Pharmaceutics, 2003, 255, 49-56.	5.2	20
155	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
156	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
157	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
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