Sung-joo Hwang

List of Publications by Year in descending order

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109321 110387 4,873 135 35 64 citations g-index h-index papers 136 136 136 5528 docs citations times ranked citing authors all docs

#	Article	IF	Citations
1	Preparation and characterization of chitosan microparticles intended for controlled drug delivery. International Journal of Pharmaceutics, 2002, 249, 165-174.	5.2	388
2	Preparation of alginate beads for floating drug delivery system: effects of CO2 gas-forming agents. International Journal of Pharmaceutics, 2002, 239, 81-91.	5.2	237
3	Preparation, characterization and in vivo evaluation of amorphous atorvastatin calcium nanoparticles using supercritical antisolvent (SAS) process. European Journal of Pharmaceutics and Biopharmaceutics, 2008, 69, 454-465.	4.3	224
4	Preparation and characterization of simvastatin/hydroxypropyl- \hat{l}^2 -cyclodextrin inclusion complex using supercritical antisolvent (SAS) process. European Journal of Pharmaceutics and Biopharmaceutics, 2007, 66, 413-421.	4.3	195
5	Improved physicochemical characteristics of felodipine solid dispersion particles by supercritical anti-solvent precipitation process. International Journal of Pharmaceutics, 2005, 301, 199-208.	5.2	170
6	Bioavailability of indomethacin-saccharin cocrystals. Journal of Pharmacy and Pharmacology, 2010, 62, 1560-1568.	2.4	148
7	Physicochemical properties and oral bioavailability of amorphous atorvastatin hemi-calcium using spray-drying and SAS process. International Journal of Pharmaceutics, 2008, 359, 211-219.	5.2	146
8	Enhancing the solubility and bioavailability of poorly water-soluble drugs using supercritical antisolvent (SAS) process. International Journal of Pharmaceutics, 2018, 538, 1-13.	5.2	144
9	Enhanced oral bioavailability of paclitaxel by coadministration of the P-glycoprotein inhibitor KR30031. Pharmaceutical Research, 2003, 20, 24-30.	3.5	120
10	Preparation and characterization of drug-loaded polymethacrylate microspheres by an emulsion solvent evaporation method. Journal of Microencapsulation, 2002, 19, 811-822.	2.8	110
11	Influence of the delivery systems using a microneedle array on the permeation of a hydrophilic molecule, calcein. European Journal of Pharmaceutics and Biopharmaceutics, 2008, 69, 1040-1045.	4.3	109
12	Micronization of cilostazol using supercritical antisolvent (SAS) process: Effect of process parameters. Powder Technology, 2007, 177, 64-70.	4.2	94
13	Recrystallization of fluconazole using the supercritical antisolvent (SAS) process. International Journal of Pharmaceutics, 2007, 328, 152-160.	5.2	75
14	Enhanced dissolution of megestrol acetate microcrystals prepared by antisolvent precipitation process using hydrophilic additives. International Journal of Pharmaceutics, 2010, 396, 91-98.	5.2	75
15	Enhanced bioavailability of sirolimus via preparation of solid dispersion nanoparticles using a supercritical antisolvent process. International Journal of Nanomedicine, 2011, 6, 2997.	6.7	73
16	Preparation and characterization of solid dispersions of itraconazole by using aerosol solvent extraction system for improvement in drug solubility and bioavailability. Archives of Pharmacal Research, 2005, 28, 866-874.	6.3	72
17	A mixed polymeric micellar formulation of itraconazole: Characteristics, toxicity and pharmacokinetics. Journal of Controlled Release, 2007, 117, 59-67.	9.9	70
18	In vivo efficacy of paclitaxel-loaded injectable in situ-forming gel against subcutaneous tumor growth. International Journal of Pharmaceutics, 2010, 392, 51-56.	5.2	68

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19	Application of the Discrete Element Method for Manufacturing Process Simulation in the Pharmaceutical Industry. Pharmaceutics, 2019, 11, 414.	4.5	67
20	Hydrogel patches containing Triclosan for acne treatment. European Journal of Pharmaceutics and Biopharmaceutics, 2003, 56, 407-412.	4.3	62
21	Preparation and Evaluation of Solid Dispersion of Atorvastatin Calcium with Soluplus® by Spray Drying Technique. Chemical and Pharmaceutical Bulletin, 2014, 62, 545-551.	1.3	62
22	Solubilization of the poorly water soluble drug, telmisartan, using supercritical anti-solvent (SAS) process. International Journal of Pharmaceutics, 2013, 441, 50-55.	5.2	58
23	Development and optimization of a novel oral controlled delivery system for tamsulosin hydrochloride using response surface methodology. International Journal of Pharmaceutics, 2007, 341, 97-104.	5.2	57
24	Mechanism of eutectic formation upon compaction and its effects on tablet properties. Thermochimica Acta, 2003, 404, 213-226.	2.7	53
25	Release characteristics of ibuprofen from excipient-loaded alginate gel beads. International Journal of Pharmaceutics, 1995, 116, 125-128.	5.2	52
26	Characteristics of felodipine-located poly($\hat{l}\mu$ -caprolactone) microspheres. Journal of Microencapsulation, 2005, 22, 193-203.	2.8	52
27	Advances in pharmaceutical materials and processing. Pharmaceutical Science & Technology Today, 1998, 1, 235-245.	0.7	49
28	The Delivery Strategy of Paclitaxel Nanostructured Lipid Carrier Coated with Platelet Membrane. Cancers, 2019, 11, 807.	3.7	46
29	Preparation and physicochemical characterization of phase inverted water/oil microemulsion containing cyclosporin A. International Journal of Pharmaceutics, 1997, 147, 131-134.	5.2	45
30	Characterization and stability studies of a novel liposomal cyclosporin A prepared using the supercritical fluid method: comparison with the modified conventional Bangham method. International Journal of Nanomedicine, 2013, 8, 365.	6.7	44
31	Controlled delivery of a hydrophilic drug from a biodegradable microsphere system by supercritical anti-solvent precipitation technique. Journal of Microencapsulation, 2006, 23, 741-749.	2.8	43
32	Supersaturatable formulations for the enhanced oral absorption of sirolimus. International Journal of Pharmaceutics, 2013, 445, 108-116.	5.2	41
33	Oral absorption of atorvastatin solid dispersion based on cellulose or pyrrolidone derivative polymers. International Journal of Biological Macromolecules, 2013, 59, 138-142.	7.5	40
34	Quality by design: screening of critical variables and formulation optimization of Eudragit E nanoparticles containing dutasteride. Archives of Pharmacal Research, 2013, 36, 593-601.	6.3	38
35	Cefuroxime axetil solid dispersions prepared using solution enhanced dispersion by supercritical fluids. Journal of Pharmacy and Pharmacology, 2010, 57, 1529-1537.	2.4	37
36	Preparation and Evaluation of Resveratrol-Loaded Composite Nanoparticles Using a Supercritical Fluid Technology for Enhanced Oral and Skin Delivery. Antioxidants, 2019, 8, 554.	5.1	37

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37	Preparation and pharmaceutical characterization of amorphous cefdinir using spray-drying and SAS-process. International Journal of Pharmaceutics, 2010, 396, 239-245.	5.2	35
38	Optimized formulation of solid self-microemulsifying sirolimus delivery systems. International Journal of Nanomedicine, 2013, 8, 1673.	6.7	35
39	Preparation, characterization, and evaluation of celecoxib eutectic mixtures with adipic acid/saccharin for improvement of wettability and dissolution rate. International Journal of Pharmaceutics, 2019, 554, 61-71.	5.2	35
40	Supercritical fluid-mediated liposomes containing cyclosporin A for the treatment of dry eye syndrome in a rabbit model: comparative study with the conventional cyclosporin A emulsion. International Journal of Nanomedicine, 2014, 9, 3791.	6.7	33
41	Design of salmon calcitonin particles for nasal delivery using spray-drying and novel supercritical fluid-assisted spray-drying processes. International Journal of Pharmaceutics, 2015, 478, 288-296.	5.2	33
42	Oxaliplatin-loaded chemically cross-linked hydrogels for prevention of postoperative abdominal adhesion and colorectal cancer therapy. International Journal of Pharmaceutics, 2019, 565, 50-58.	5.2	33
43	Topical Oleo-Hydrogel Preparation of Ketoprofen with Enhanced Skin Permeability. Drug Development and Industrial Pharmacy, 1999, 25, 717-726.	2.0	32
44	Liposomal drug products and recent advances in the synthesis of supercritical fluid-mediated liposomes. Nanomedicine, 2013, 8, 1529-1548.	3.3	32
45	Effect of operating parameters on PVP/tadalafil solid dispersions prepared using supercritical anti-solvent process. Journal of Supercritical Fluids, 2014, 90, 126-133.	3.2	32
46	Effect of Stabilizers on Encapsulation Efficiency and Release Behavior of Exenatide-Loaded PLGA Microsphere Prepared by the W/O/W Solvent Evaporation Method. Pharmaceutics, 2019, 11, 627.	4.5	31
47	Development of a Resveratrol Nanosuspension Using the Antisolvent Precipitation Method without Solvent Removal, Based on a Quality by Design (QbD) Approach. Pharmaceutics, 2019, 11, 688.	4.5	31
48	Preparation and evaluation of cyclosporin A-containing proliposomes: a comparison of the supercritical antisolvent process with the conventional film method. International Journal of Nanomedicine, 2014, 9, 5079.	6.7	30
49	Preparation and Evaluation of Intraperitoneal Long-Acting Oxaliplatin-Loaded Multi-Vesicular Liposomal Depot for Colorectal Cancer Treatment. Pharmaceutics, 2020, 12, 736.	4.5	30
50	Enhancement of dissolution and bioavailability of ezetimibe by amorphous solid dispersion nanoparticles fabricated using supercritical antisolvent process. Journal of Pharmaceutical Investigation, 2015, 45, 641-649.	5.3	29
51	Enhancement of Wettability and Dissolution Properties of Cilostazol Using the Supercritical Antisolvent Process: Effect of Various Additives. Chemical and Pharmaceutical Bulletin, 2010, 58, 230-233.	1.3	28
52	The advent of a novel manufacturing technology in pharmaceutics: superiority of fused deposition modeling 3D printer. Journal of Pharmaceutical Investigation, 2020, 50, 131-145.	5.3	28
53	Characterization and therapeutic efficacy evaluation of glimepiride and L-arginine co-amorphous formulation prepared by supercritical antisolvent process: Influence of molar ratio and preparation methods. International Journal of Pharmaceutics, 2020, 581, 119232.	5.2	27
54	Phase Behavior of Poly(l-lactide) in Supercritical Mixtures of Carbon Dioxide and Chlorodifluoromethane. Journal of Chemical & Engineering Data, 2000, 45, 1162-1166.	1.9	26

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55	Surface modification of paclitaxel-loaded liposomes using $d\cdot \hat{l}\pm$ -tocopheryl polyethylene glycol 1000 succinate: Enhanced cellular uptake and cytotoxicity in multidrug resistant breast cancer cells. Chemistry and Physics of Lipids, 2018, 213, 39-47.	3.2	26
56	An update on niche composition, signaling and functional regulation of the adipose-derived stem cells. Expert Opinion on Biological Therapy, 2014, 14, 1091-1102.	3.1	25
57	Dissolution and oral absorption of pranlukast nanosuspensions stabilized by hydroxypropylmethyl cellulose. International Journal of Biological Macromolecules, 2014, 67, 53-57.	7.5	25
58	LC–MS determination and bioavailability study of loperamide hydrochloride after oral administration of loperamide capsule in human volunteers. Journal of Pharmaceutical and Biomedical Analysis, 2004, 36, 421-427.	2.8	24
59	Enhancement of the dissolution rate and bioavailability of fenofibrate by a melt-adsorption method using supercritical carbon dioxide. International Journal of Nanomedicine, 2012, 7, 5565.	6.7	24
60	Enhanced solubility and oral absorption of sirolimus using D-α-tocopheryl polyethylene glycol succinate micelles. Artificial Cells, Nanomedicine and Biotechnology, 2013, 41, 85-91.	2.8	24
61	Functional characterization of mesenchymal stem cells labeled with a novel PVPâ€coated superparamagnetic iron oxide. Contrast Media and Molecular Imaging, 2009, 4, 118-126.	0.8	23
62	Application of diethylene glycol monoethyl ether in solubilization of poorly water-soluble drugs. Journal of Pharmaceutical Investigation, 2020, 50, 231-250.	5.3	23
63	Effect of Solvent Type on the Nanoparticle Formation of Atorvastatin Calcium by the Supercritical Antisolvent Process. Chemical and Pharmaceutical Bulletin, 2012, 60, 543-547.	1.3	22
64	Oral absorption of a valsartan-loaded spray-dried emulsion based on hydroxypropylmethyl cellulose. International Journal of Biological Macromolecules, 2014, 69, 222-228.	7.5	22
65	Lysophosphatidic acid increases the proliferation and migration of adipose-derived stem cells via the generation of reactive oxygen species. Molecular Medicine Reports, 2015, 12, 5203-5210.	2.4	22
66	Dissolution and bioavailability of lercanidipine–hydroxypropylmethyl cellulose nanoparticles with surfactant. International Journal of Biological Macromolecules, 2015, 72, 218-222.	7.5	22
67	Enhanced Supersaturation and Oral Absorption of Sirolimus Using an Amorphous Solid Dispersion Based on Eudragit® E. Molecules, 2015, 20, 9496-9509.	3.8	21
68	Cyclosporine A micellar delivery system for dry eyes. International Journal of Nanomedicine, 2016, 11, 2921.	6.7	21
69	Solid-State Carbon NMR Characterization and Investigation of Intrinsic Dissolution Behavior of Fluconazole Polymorphs, Anhydrate Forms I and II. Chemical and Pharmaceutical Bulletin, 2010, 58, 1243-1247.	1.3	20
70	Fluoxetine Decreases the Proliferation and Adipogenic Differentiation of Human Adipose-Derived Stem Cells. International Journal of Molecular Sciences, 2015, 16, 16655-16668.	4.1	20
71	Development of Houttuynia cordata Extract-Loaded Solid Lipid Nanoparticles for Oral Delivery: High Drug Loading Efficiency and Controlled Release. Molecules, 2017, 22, 2215.	3.8	20
72	Pharmacokinetic Profile and Anti-Adhesive Effect of Oxaliplatin-PLGA Microparticle-Loaded Hydrogels in Rats for Colorectal Cancer Treatment. Pharmaceutics, 2019, 11, 392.	4.5	19

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73	Controlled release tamsulosin hydrochloride from alginate beads with waxy materialsâ€. Journal of Pharmacy and Pharmacology, 2010, 57, 1521-1528.	2.4	18
74	Solubility of oxcarbazepine in eight solvents within the temperature range T = (288.15–308.15) K. Journal of Chemical Thermodynamics, 2017, 104, 45-49.	2.0	18
75	Preparation, Characterization, and In Vivo Pharmacokinetic Study of the Supercritical Fluid-Processed Liposomal Amphotericin B. Pharmaceutics, 2019, 11, 589.	4.5	18
76	The organ targetability of small and large albumin microspheres containing free and HSA conjugated methotrexate. International Journal of Pharmaceutics, 1993, 89, 91-102.	5.2	17
77	Isolation of adipose-derived stem cells by using a subfractionation culturing method. Expert Opinion on Biological Therapy, 2014, 14, 1551-1560.	3.1	17
78	Preparation and characterization of mucoadhesive enteric-coating ginsenoside-loaded microparticles. Archives of Pharmacal Research, 2015, 38, 761-768.	6.3	17
79	Diazepam sorption to PVC- and non-PVC-based tubes in administration sets with quantitative determination using a high-performance liquid chromatographic method. International Journal of Pharmaceutics, 2016, 506, 414-419.	5.2	17
80	Solubility of Simvastatin and Lovastatin in Mixtures of Dichloromethane and Supercritical Carbon Dioxide. Journal of Chemical & Engineering Data, 2007, 52, 1273-1279.	1.9	16
81	A Mixed Micellar Formulation for the Transdermal Delivery of an Indirubin Analog. Pharmaceutics, 2020, 12, 175.	4.5	16
82	Chitosan microparticle preparation for controlled drug release by response surface methodology. Journal of Microencapsulation, 2003, 20, 791-797.	2.8	15
83	Influence of water soluble additives and HPMCP on drug release from sureleaseê-coated pellets containing tamsulosin hydrochloride. Archives of Pharmacal Research, 2007, 30, 1008-1013.	6.3	14
84	Intracorneal melatonin delivery using 2-hydroxypropyl- \hat{l}^2 -cyclodextrin ophthalmic solution for granular corneal dystrophy type 2. International Journal of Pharmaceutics, 2017, 529, 608-616.	5.2	14
85	The influence of Surelease and sodium alginate on the in-vitro release of tamsulosin hydrochloride in pellet dosage form. Journal of Pharmacy and Pharmacology, 2010, 57, 735-742.	2.4	13
86	Fabrication and evaluation of celecoxib microparticle surface modified by hydrophilic cellulose and surfactant. International Journal of Biological Macromolecules, 2015, 72, 1473-1478.	7. 5	13
87	Sustained release of risperidone from biodegradable microspheres prepared by in-situ suspension-evaporation process. International Journal of Pharmaceutics, 2016, 503, 8-15.	5.2	13
88	Evaluation of tacrolimus sorption to PVC- and non-PVC-based tubes in administration sets: Pump method vs. drip method. International Journal of Pharmaceutics, 2017, 528, 172-179.	5.2	13
89	Preparation and characterization of glimepiride eutectic mixture with l-arginine for improvement of dissolution rate. International Journal of Pharmaceutics, 2020, 581, 119288.	5.2	13
90	Micronization of a poorly water-soluble drug, fenofibrate, via supercritical-fluid-assisted spray-drying. Journal of Pharmaceutical Investigation, 2022, 52, 353-366.	5.3	13

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91	Pharmacokinetics of methotrexate after intravenous and intramuscular injection of methotrexate-bearing positively charged liposomes to rats. Biopharmaceutics and Drug Disposition, 1995, 16, 279-293.	1.9	11
92	Paroxetine hydrochloride controlled release POLYOX® matrix tablets: Screening of formulation variables using Plackett-Burman screening design. Archives of Pharmacal Research, 2008, 31, 399-405.	6.3	11
93	Delivery of interleukin-18 gene to lung cancer cells using cationic emulsion. Journal of Drug Targeting, 2009, 17, 19-28.	4.4	11
94	Biocompatible Polyhydroxyethylaspartamide-based Micelles with Gadolinium for MRI Contrast Agents. Nanoscale Research Letters, 2010, 5, 1970-1976.	5.7	11
95	The Effect of Sodium Alginate on Physical and Dissolution Properties of Surelease-Matrix Pellets Prepared by a Novel Pelletizer. Chemical and Pharmaceutical Bulletin, 2007, 55, 1631-1634.	1.3	10
96	Comparative study of telmisartan tablets prepared via the wet granulation method and pritorâ, prepared using the spray-drying method. Archives of Pharmacal Research, 2011, 34, 463-468.	6.3	10
97	Pharmaceutical Applications of Supercritical Fluid Extraction of Emulsions for Micro-/Nanoparticle Formation. Pharmaceutics, 2021, 13, 1928.	4.5	10
98	Gelatin Coating for the Improvement of Stability and Cell Uptake of Hydrophobic Drug-Containing Liposomes. Molecules, 2022, 27, 1041.	3.8	10
99	Preparation and evaluation of a titrated extract of Centella asiatica injection in the form of an extemporaneous micellar solution. International Journal of Pharmaceutics, 1997, 146, 63-70.	5.2	9
100	pH-independent sustained release matrix tablet containing doxazosin mesylate: Effect of citric acid. Archives of Pharmacal Research, 2010, 33, 2003-2009.	6.3	9
101	LC-MS determination and bioavailability study of imidapril hydrochloride after the oral administration of imidapril tablets in human volunteers. Archives of Pharmacal Research, 2005, 28, 463-468.	6.3	8
102	Quantitative determination of sirolimus in dog blood using liquid chromatography–tandem mass spectrometry, and its applications to pharmacokinetic studies. Journal of Pharmaceutical and Biomedical Analysis, 2010, 53, 1042-1047.	2.8	8
103	A novel thermosensitive poloxamer-hyaluronic acid- kappa-carrageenan-based hydrogel anti-adhesive agent loaded with 5-fluorouracil: A preclinical study in Sprague-Dawley rats. International Journal of Pharmaceutics, 2022, 621, 121771.	5.2	8
104	Statistical Optimization of Tamsulosin Hydrochloride Controlled Release Pellets Coated with the Blend of HPMCP and HPMC. Chemical and Pharmaceutical Bulletin, 2007, 55, 936-939.	1.3	7
105	Pharmacologic Properties of the Carrier Solutions for Hyperthermic Intraperitoneal Chemotherapy: Comparative Analyses Between Water and Lipid Carrier Solutions in the Rat Model. Annals of Surgical Oncology, 2018, 25, 3185-3192.	1.5	7
106	Optimization of tamsulosin hydrochloride controlled release pellets coated with Surelease and neutralized HPMCPâ€. Journal of Pharmacy and Pharmacology, 2010, 58, 1611-1616.	2.4	6
107	The feasibility study of transdermal drug delivery systems for antidepressants possessing hydrophilicity or hydrophobicity. Journal of Pharmaceutical Investigation, 2012, 42, 109-114.	5.3	6
108	Biodistribution of newly synthesized PHEA-based polymer-coated SPION in Sprague Dawley rats as magnetic resonance contrast agent. International Journal of Nanomedicine, 2013, 8, 4077.	6.7	6

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109	Boiling Method-Based Zinc Oxide Nanorods for Enhancement of Adipose-Derived Stem Cell Proliferation. Tissue Engineering - Part C: Methods, 2016, 22, 847-855.	2.1	6
110	Formulation, Preparation, Characterization, and Evaluation of Dicarboxylic Ionic Liquid Donepezil Transdermal Patches. Pharmaceutics, 2022, 14, 205.	4.5	6
111	Preparation of controlled release spheronized beads by a simple extrusion and modified spheronization process. Archives of Pharmacal Research, 2005, 28, 619-625.	6.3	5
112	Ocular delivery systems for the administration of antibody therapeutics. Journal of Pharmaceutical Investigation, 2017, 47, 373-382.	5.3	5
113	Pharmaceutical Characterization and In Vivo Evaluation of Orlistat Formulations Prepared by the Supercritical Melt-Adsorption Method Using Carbon Dioxide: Effects of Mesoporous Silica Type. Pharmaceutics, 2020, 12, 333.	4.5	5
114	Melt Amorphisation of Orlistat with Mesoporous Silica Using a Supercritical Carbon Dioxide: Effects of Pressure, Temperature, and Drug Loading Ratio and Comparison with Other Conventional Amorphisation Methods. Pharmaceutics, 2020, 12, 377.	4.5	5
115	Preparation of substained-release microspheres of phenylpropanolamine HCl and their release characteristics. Archives of Pharmacal Research, 1990, 13, 293-297.	6.3	4
116	Design of pH-independent extended release matrix tablets of minocycline hydrochloride for the treatment of dementia. Archives of Pharmacal Research, 2009, 32, 1593-1598.	6.3	4
117	Pharmacokinetic and bioequivalence study of itopride HCl in healthy volunteers. Arzneimittelforschung, 2010, 60, 137-140.	0.4	4
118	Development and Optimization of a Novel Sustained-release Tablet Formulation for Bupropion Hydrochloride using Box-Behnken Design. Journal of Pharmaceutical Investigation, 2010, 40, 313-319.	5.3	4
119	Phase behavior of water-insoluble simvastatin drug in supercritical mixtures of chlorodifluoromethane and carbon dioxide. Korean Journal of Chemical Engineering, 2006, 23, 1009-1015.	2.7	3
120	Evaluation of Drug Sorption to PVC- and Non-PVC-based Tubes in Administration Sets Using a Pump. Journal of Visualized Experiments, 2017, , .	0.3	3
121	Evaluation of nitroglycerin and cyclosporin A sorption to polyvinylchloride- and non-polyvinylchloride-based tubes in administration sets. Journal of Pharmaceutical Investigation, 2018, 48, 665-672.	5.3	3
122	Development of Self-microemulsifying Drug Delivery System for Enhancing the Bioavailability of Atorvastatin. Journal of Pharmaceutical Investigation, 2011, 41, 103-109.	5.3	3
123	Molecular Mechanism of Cinnamomum cassia against Gastric Damage and Identification of Active Compounds. Biomolecules, 2022, 12, 525.	4.0	3
124	Comparison of adhesion and dissolution of fentanyl patches: Fentadur® and Durogesic DTrans®. Journal of Pharmaceutical Investigation, 2015, 45, 475-480.	5.3	2
125	Preformulation of FK506 Prodrugs for Improving Solubility. Bulletin of the Korean Chemical Society, 2016, 37, 1313-1319.	1.9	2
126	Preparation of Sprayâ€dried Emulsion of Sirolimus for Enhanced Oral Bioavailability. Bulletin of the Korean Chemical Society, 2018, 39, 1215-1218.	1.9	2

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127	Microencapsulation of isoprinosine with ethylcellulose. Archives of Pharmacal Research, 1991, 14, 298-304.	6.3	1
128	Preparation andIn vitro release characteristics of hydrophilic albumin microspheres containing methotrexate and methotrexate-human serum albumin conjugates. Archives of Pharmacal Research, 1992, 15, 162-168.	6.3	1
129	Improvement of Dissolution Rate of Oxcarbazepine Using Surfaceâ€modified Solid Dispersion with Vinylpyrrolidoneâ€√inyl Acetate Copolymer and Sucrose Laurate. Bulletin of the Korean Chemical Society, 2018, 39, 995-998.	1.9	1
130	Case Study of Pharmaceutical Ingredients Derived from Clay Minerals. Economic and Environmental Geology, 2015, 48, 221-229.	0.4	1
131	Preparation and Characterization of Fenofibrate Microparticles with Surface-Active Additives: Application of a Supercritical Fluid-Assisted Spray-Drying Process. Pharmaceutics, 2021, 13, 2061.	4.5	1
132	Dataset for hierarchical tetramodal-porous architecture of zinc oxide nanoparticles microfluidically synthesized via dual-step nanofabrication. Data in Brief, 2022, 42, 108137.	1.0	1
133	Comparison on physical property, dissolution and disintegration of four launched orally disintegration film (ODF) products for erectile dysfunction. Journal of Pharmaceutical Investigation, 2014, 44, 297-307.	5.3	0
134	Effect of Polymer Type on the Dissolution Profile of a Solid Dispersion of Cilostazol. Bulletin of the Korean Chemical Society, 2019, 40, 370-373.	1.9	0
135	Pharmacokinetic and Bioequivalence Study of Zolpidem Tartate in Healthy Volunteers. Journal of Pharmaceutical Investigation, 2011, 41, 191-196.	5. 3	0