Young Wook Choi

List of Publications by Year in descending order

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Version: 2024-02-01

		218677	3	302126
89	1,989	26		39
papers	citations	h-index		g-index
				0.6.5
92	92	92		2665
all docs	docs citations	times ranked		citing authors

#	Article	IF	CITATIONS
1	Immunotherapeutic effects of recombinant Bacillus Calmette–Guérin containing ⟨i⟩sic⟨ i⟩ gene in ⟨i⟩ex vivo⟨ i⟩ and ⟨i⟩in vivo⟨ i⟩ bladder cancer models. Investigative and Clinical Urology, 2022, 63, 228.	2.0	2
2	Co-administration of tariquidar using functionalized nanostructured lipid carriers overcomes resistance to docetaxel in multidrug resistant MCF7/ADR cells. Journal of Drug Delivery Science and Technology, 2022, , 103323.	3.0	3
3	Enhanced dissolution and bioavailability of revaprazan using self-nanoemulsifying drug delivery system. Pharmaceutical Development and Technology, 2022, 27, 414-424.	2.4	6
4	Synergistic co-administration of docetaxel and curcumin to chemoresistant cancer cells using PEGylated and RIPL peptide-conjugated nanostructured lipid carriers. Cancer Nanotechnology, 2022, 13, .	3.7	5
5	Supersaturable self-microemulsifying drug delivery system enhances dissolution and bioavailability of telmisartan. Pharmaceutical Development and Technology, 2021, 26, 60-68.	2.4	7
6	Development of a Solid Supersaturable Micelle of Revaprazan for Improved Dissolution and Oral Bioavailability Using Box-Behnken Design. International Journal of Nanomedicine, 2021, Volume 16, 1245-1259.	6.7	9
7	Facilitated Buccal Insulin Delivery via Hydrophobic Ion-Pairing Approach: In vitro and ex vivo Evaluation. International Journal of Nanomedicine, 2021, Volume 16, 4677-4691.	6.7	9
8	Cochleate Formulation Enhances the Stability of Lansoprazole in Acidic Condition. Bulletin of the Korean Chemical Society, 2021, 42, 1281-1284.	1.9	3
9	Establishment of Three-Dimensional Bioprinted Bladder Cancer-on-a-Chip with a Microfluidic System Using Bacillus Calmette–Guérin. International Journal of Molecular Sciences, 2021, 22, 8887.	4.1	12
10	Optimization of a floating poloxamer 407-based hydrogel using the Box-Behnken design: in vitro characterization and in vivo buoyancy evaluation for intravesical instillation. European Journal of Pharmaceutical Sciences, 2021, 163, 105885.	4.0	12
11	European Regulatory Science and Regulatory Science Expert Training Project. Korean Journal of Clinical Pharmacy, 2021, 31, 171-179.	0.3	1
12	Bile acid transporter-mediated oral absorption of insulin via hydrophobic ion-pairing approach. Journal of Controlled Release, 2021, 338, 644-661.	9.9	22
13	Poloxamer 407â€based Floating Hydrogels for Intravesical Instillation: Statistical Optimization Using Central Composite Design, Gel Erosion, and Drug Release. Bulletin of the Korean Chemical Society, 2021, 42, 72-79.	1.9	9
14	Effects of periostin deficiency on kidney aging and lipid metabolism. Aging, 2021, 13, 22649-22665.	3.1	8
15	Analysis of Trends in Regulatory Science and Regulatory Science Experts Training Projects: US, Japan, Singapore, and Korea. Korean Journal of Clinical Pharmacy, 2021, 31, 257-267.	0.3	3
16	Enhanced oral bioavailability of valsartan in rats using a supersaturable self-microemulsifying drug delivery system with P-glycoprotein inhibitors. Pharmaceutical Development and Technology, 2020, 25, 178-186.	2.4	15
17	Current status of the development of intravesical drug delivery systems for the treatment of bladder cancer. Expert Opinion on Drug Delivery, 2020, 17, 1555-1572.	5.0	33
18	Liposome-Encapsulated Bacillus Calmette–Guérin Cell Wall Skeleton Enhances Antitumor Efficiency for Bladder Cancer In Vitro and In Vivo via Induction of AMP-Activated Protein Kinase. Cancers, 2020, 12, 3679.	3.7	17

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19	Optimization of solid self-dispersing micelle for enhancing dissolution and oral bioavailability of valsartan using Box-Behnken design. International Journal of Pharmaceutics, 2020, 585, 119483.	5.2	12
20	Enhanced docetaxel delivery using sterically stabilized RIPL peptide-conjugated nanostructured lipid carriers: In vitro and in vivo antitumor efficacy against SKOV3 ovarian cancer cells. International Journal of Pharmaceutics, 2020, 583, 119393.	5.2	17
21	Alnus Sibirica Extracts Suppress the Expression of Inflammatory Cytokines Induced by Lipopolysaccharides, Tumor Necrosis Factor- \hat{l}_{\pm} , and Interferon- \hat{l}_{3} in Human Dermal Fibroblasts. Molecules, 2019, 24, 2883.	3.8	11
22	Design and In Vivo Pharmacokinetic Evaluation of Triamcinolone Acetonide Microcrystals-Loaded PLGA Microsphere for Increased Drug Retention in Knees after Intra-Articular Injection. Pharmaceutics, 2019, 11, 419.	4.5	20
23	<p>Intravesical delivery of rapamycin via folate-modified liposomes dispersed in thermo-reversible hydrogel</p> . International Journal of Nanomedicine, 2019, Volume 14, 6249-6268.	6.7	42
24	Improved Dissolution and Oral Bioavailability of Valsartan Using a Solidified Supersaturable Self-Microemulsifying Drug Delivery System Containing Gelucire® 44/14. Pharmaceutics, 2019, 11, 58.	4.5	23
25	Improved Drug Loading and Sustained Release of Entecavirâ€loaded PLGA Microsphere Prepared by Spray Drying Technique. Bulletin of the Korean Chemical Society, 2019, 40, 306-312.	1.9	10
26	Enhanced Intracellular Delivery of BCG Cell Wall Skeleton into Bladder Cancer Cells Using Liposomes Functionalized with Folic Acid and Pep-1 Peptide. Pharmaceutics, 2019, 11, 652.	4.5	14
27	The immunotherapeutic effects of recombinant Bacillus Calmette-Guérin resistant to antimicrobial peptides on bladderÂcancerÂcells. Biochemical and Biophysical Research Communications, 2019, 509, 167-174.	2.1	13
28	Rapamycin enhances growth inhibition on urothelial carcinoma cells through LKB1 deficiencyâ€mediated mitochondrial dysregulation. Journal of Cellular Physiology, 2019, 234, 13083-13096.	4.1	11
29	Steric stabilization of RIPL peptide-conjugated liposomes and in vitro assessment. Journal of Pharmaceutical Investigation, 2019, 49, 115-125.	5.3	12
30	Recent advances in intra-articular drug delivery systems to extend drug retention in joint. Journal of Pharmaceutical Investigation, 2019, 49, 9-15.	5.3	12
31	Formulation and in vivo pharmacokinetic evaluation of ethyl cellulose-coated sustained release multiple-unit system of tacrolimus. International Journal of Biological Macromolecules, 2018, 109, 544-550.	7.5	17
32	Combined Poly(Lactide-Co-Glycolide) Microspheres Containing Diphtheria Toxoid for a Single-shot Immunization. AAPS PharmSciTech, 2018, 19, 1160-1167.	3.3	4
33	pH-sensitive PEGylation of RIPL peptide-conjugated nanostructured lipid carriers: design and in vitro evaluation. International Journal of Nanomedicine, 2018, Volume 13, 6661-6675.	6.7	15
34	Optimization of self-microemulsifying drug delivery system for phospholipid complex of telmisartan using D-optimal mixture design. PLoS ONE, 2018, 13, e0208339.	2.5	17
35	Sterically Stabilized RIPL Peptide-Conjugated Nanostructured Lipid Carriers: Characterization, Cellular Uptake, Cytotoxicity, and Biodistribution. Pharmaceutics, 2018, 10, 199.	4.5	14
36	Development and Evaluation of a Water Soluble Fluorometholone Eye Drop Formulation Employing Polymeric Micelle. Pharmaceutics, 2018, 10, 208.	4.5	22

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37	Enhanced Chemical Stability of Hirsutenone Incorporated into a Nanostructured Lipid Carrier Formulation Containing Antioxidants. Bulletin of the Korean Chemical Society, 2018, 39, 1287-1293.	1.9	2
38	Immediate release tablet formulation of varenicline salicylate and comparative pharmacokinetic study in human volunteers. Drug Design, Development and Therapy, 2018, Volume 12, 3377-3392.	4.3	4
39	Facilitated permeation of insulin across TR146 cells by cholic acid derivatives-modified elastic bilosomes. International Journal of Nanomedicine, 2018, Volume 13, 5173-5186.	6.7	28
40	Effect of Poly(Lactide-Co-Glycolide) Nanoparticles on Local Retention of Fluorescent Material: An Experimental Study in Mice. Korean Journal of Radiology, 2018, 19, 950.	3.4	3
41	Novel Extended-Release Multiple-Unit System of Imidafenacin Prepared by Fluid-Bed Coating Technique. AAPS PharmSciTech, 2018, 19, 2639-2645.	3.3	3
42	RIPL peptide-conjugated nanostructured lipid carriers for enhanced intracellular drug delivery to hepsin-expressing cancer cells. International Journal of Nanomedicine, 2018, Volume 13, 3263-3278.	6.7	24
43	Tablet Formulation of a Polymeric Solid Dispersion Containing Amorphous Alkalinized Telmisartan. AAPS PharmSciTech, 2018, 19, 2990-2999.	3.3	16
44	Development of a chitosan based double layer-coated tablet as a platform for colon-specific drug delivery. Drug Design, Development and Therapy, 2017, Volume11, 45-57.	4.3	26
45	Surface modification of lipid-based nanocarriers for cancer cell-specific drug targeting. Journal of Pharmaceutical Investigation, 2017, 47, 203-227.	5.3	96
46	Enhanced Transdermal Delivery by Combined Application of Dissolving Microneedle Patch on Serum-Treated Skin. Molecular Pharmaceutics, 2017, 14, 2024-2031.	4.6	34
47	Docetaxel-loaded RIPL peptide (IPLVVPLRRRRRRRRC)-conjugated liposomes: Drug release, cytotoxicity, and antitumor efficacy. International Journal of Pharmaceutics, 2017, 523, 229-237.	5.2	38
48	A Polyvinylpyrrolidone-Based Supersaturable Self-Emulsifying Drug Delivery System for Enhanced Dissolution of Cyclosporine A. Polymers, 2017, 9, 124.	4.5	19
49	Surface-Modification of RIPL Peptide-Conjugated Liposomes to Achieve Steric Stabilization and pH Sensitivity. Journal of Nanoscience and Nanotechnology, 2017, 17, 1008-1017.	0.9	5
50	Enhanced oral bioavailability of valsartan using a polymer-based supersaturable self-microemulsifying drug delivery system. International Journal of Nanomedicine, 2017, Volume 12, 3533-3545.	6.7	53
51	Solid formulation of a supersaturable self-microemulsifying drug delivery system for valsartan with improved dissolution and bioavailability. Oncotarget, 2017, 8, 94297-94316.	1.8	21
52	Poloxamer 407 Hydrogels for Intravesical Instillation to Mouse Bladder: Gel-Forming Capacity and Retention Performance. The Korean Journal of Urological Oncology, 2017, 15, 178-186.	0.1	10
53	Increased localized delivery of piroxicam by cationic nanoparticles after intra-articular injection. Drug Design, Development and Therapy, 2016, Volume 10, 3779-3787.	4.3	24
54	Enhanced dissolution and oral absorption of tacrolimus by supersaturable self-emulsifying drug delivery system. International Journal of Nanomedicine, 2016, 11, 1109.	6.7	20

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55	Cell penetrating peptides as an innovative approach for drug delivery; then, present and the future. Journal of Pharmaceutical Investigation, 2016, 46, 205-220.	5.3	29
56	Development of a solidified self-microemulsifying drug delivery system (S-SMEDDS) for atorvastatin calcium with improved dissolution and bioavailability. International Journal of Pharmaceutics, 2016, 506, 302-311.	5.2	60
57	Formulation of controlled-release pelubiprofen tablet using Kollidon \hat{A}^{\otimes} SR. International Journal of Pharmaceutics, 2016, 511, 864-875.	5.2	10
58	A Novel Stable Crystalline Triamcinolone Acetonideâ€loaded PLGA Microsphere for Prolonged Release After Intraâ€Articular Injection. Bulletin of the Korean Chemical Society, 2016, 37, 1496-1500.	1.9	6
59	Enhanced topical delivery of tacrolimus by a carbomer hydrogel formulation with transcutol P. Drug Development and Industrial Pharmacy, 2016, 42, 1636-1642.	2.0	28
60	Topical Semisolid Formulations of Hirsutenone and Accelerated Stability Assessment. Bulletin of the Korean Chemical Society, 2015, 36, 1688-1693.	1.9	1
61	Cationic PLGA/Eudragit RL nanoparticles for increasing retention time in synovial cavity after intra-articular injection in knee joint. International Journal of Nanomedicine, 2015, 10, 5263.	6.7	29
62	Design of Multifunctional Liposomal Nanocarriers for Folate Receptor-Specific Intracellular Drug Delivery. Molecular Pharmaceutics, 2015, 12, 4200-4213.	4.6	40
63	Development and optimization of a self-microemulsifying drug delivery system for atorvastatin calcium by using D-optimal mixture design. International Journal of Nanomedicine, 2015, 10, 3865.	6.7	48
64	Fujicalin $\hat{A}^{@}$ -based solid supersaturable self-emulsifying drug delivery system (S-SEDDS) of tacrolimus for enhanced dissolution rate and oral absorption. Journal of Pharmaceutical Investigation, 2015, 45, 651-658.	5.3	10
65	Formulation of a modified-release pregabalin tablet using hot-melt coating with glyceryl behenate. International Journal of Pharmaceutics, 2015, 495, 1-8.	5.2	19
66	Improved oral absorption of dutasteride via Soluplus $\hat{A}^{@}$ -based supersaturable self-emulsifying drug delivery system (S-SEDDS). International Journal of Pharmaceutics, 2015, 478, 341-347.	5.2	56
67	Nanostructured lipid carrier-loaded hyaluronic acid microneedles for controlled dermal delivery of a lipophilic molecule. International Journal of Nanomedicine, 2014, 9, 289.	6.7	42
68	In situ intestinal permeability and in vivo oral bioavailability of celecoxib in supersaturating self-emulsifying drug delivery system. Archives of Pharmacal Research, 2014, 37, 626-635.	6.3	40
69	RIPL peptide (IPLVVPLRRRRRRRRC)-conjugated liposomes for enhanced intracellular drug delivery to hepsin-expressing cancer cells. European Journal of Pharmaceutics and Biopharmaceutics, 2014, 87, 489-499.	4.3	34
70	Inclusion compound formulation of hirsutenone with beta-cyclodextrin. Journal of Pharmaceutical Investigation, 2013, 43, 453-459.	5.3	9
71	A retinyl palmitate-loaded solid lipid nanoparticle system: Effect of surface modification with dicetyl phosphate on skin permeation in vitro and anti-wrinkle effect in vivo. International Journal of Pharmaceutics, 2013, 452, 311-320.	5.2	70
72	Enhanced dissolution of celecoxib by supersaturating self-emulsifying drug delivery system (S-SEDDS) formulation. Archives of Pharmacal Research, 2013, 36, 69-78.	6. 3	44

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73	Topical formulation of retinyl retinoate employing nanostructured lipid carriers. Journal of Pharmaceutical Investigation, 2012, 42, 243-250.	5. 3	21
74	Immediate release of ibuprofen from Fujicalin (sup) \hat{A}^{\otimes} (sup)-based fast-dissolving self-emulsifying tablets. Drug Development and Industrial Pharmacy, 2011, 37, 1298-1305.	2.0	45
75	Tat peptide-admixed elastic liposomal formulation of hirsutenone for the treatment of atopic dermatitis in Nc/Nga mice. International Journal of Nanomedicine, 2011, 6, 2459.	6.7	14
76	Design, synthesis, bioconversion, and pharmacokinetics evaluation of new ester prodrugs of olmesartan. European Journal of Medicinal Chemistry, 2011, 46, 3564-3569.	5 . 5	5
77	Solid dispersion formulations of megestrol acetate with copovidone for enhanced dissolution and oral bioavailability. Archives of Pharmacal Research, 2011, 34, 127-135.	6.3	14
78	Facilitated Skin Permeation of Oregonin by Elastic Liposomal Formulations and Suppression of Atopic Dermatitis in NC/Nga Mice. Biological and Pharmaceutical Bulletin, 2010, 33, 100-106.	1.4	41
79	Pep-1 peptide-conjugated elastic liposomal formulation of taxifolin glycoside for the treatment of atopic dermatitis in NC/Nga mice. International Journal of Pharmaceutics, 2010, 402, 198-204.	5.2	39
80	Identification and assessment of permeability enhancing vehicles for transdermal delivery of glucosamine hydrochloride. Archives of Pharmacal Research, 2010, 33, 293-299.	6.3	11
81	Stability-enhanced solid dispersion formulation of amorphous raloxifene hydrochloride. Korean Journal of Chemical Engineering, 2010, 27, 1906-1909.	2.7	9
82	Influence of Liposome Type and Skin Model on Skin Permeation and Accumulation Properties of Genistein. Journal of Dispersion Science and Technology, 2010, 31, 1061-1066.	2.4	10
83	Solubilized formulation of olmesartan medoxomil for enhancing oral bioavailability. Archives of Pharmacal Research, 2009, 32, 1629-1635.	6.3	36
84	Pharmaceutical evaluation of genistein-loaded pluronic micelles for oral delivery. Archives of Pharmacal Research, 2007, 30, 1138-1143.	6.3	89
85	Anticancer Efficacy and Toxicity of Oral GMO-paclitaxel in a Hormone Refractory Prostate Cancer Model. Korean Journal of Urology, 2006, 47, 143.	0.2	3
86	High-performance liquid chromatographic determination of doxazosin in human plasma for bioequivalence study of controlled release doxazosin tablets. Biomedical Chromatography, 2006, 20, 1172-1177.	1.7	24
87	Formulation of microemulsion systems for transdermal delivery of aceclofenac. Archives of Pharmacal Research, 2005, 28, 1097-1102.	6.3	57
88	A cationic lipid emulsion/DNA complex as a physically stable and serum-resistant gene delivery system. Pharmaceutical Research, 2000, 17, 314-320.	3 . 5	78
89	Effects of solvent selection and fabrication method on the characteristics of biodegradable poly(lactide-co-glycolide) microspheres containing ovalbumin. Archives of Pharmacal Research, 2000, 23, 385-390.	6. 3	28