Xiangtao Wang

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Enhanced tumor accumulation and therapeutic efficacy of liposomal drugs through over-threshold dosing. Journal of Nanobiotechnology, 2022, 20, 137.	9.1	7
2	Photothermal combined with intratumoral injection of annonaceous acetogenin nanoparticles for breast cancer therapy. Colloids and Surfaces B: Biointerfaces, 2022, 213, 112426.	5.0	2
3	Enhancement of oral bioavailability of quercetin by metabolic inhibitory nanosuspensions compared to conventional nanosuspensions. Drug Delivery, 2021, 28, 1226-1236.	5.7	29
4	Pterostilbene nanoparticles with small particle size show excellent anti-breast cancer activity in vitro and in vivo. Nanotechnology, 2021, 32, 325102.	2.6	8
5	Annonaceous acetogenins nanosuspensions stabilized by poloxamer 188: Preparation, properties and in vivo evaluation. Journal of Drug Delivery Science and Technology, 2021, 66, 102676.	3.0	4
6	A comparative study on the <i>inÂvitro</i> and <i>inÂvivo</i> antitumor efficacy of icaritin and hydrous icaritin nanorods. Drug Delivery, 2020, 27, 1176-1187.	5.7	9
7	Preparation of high drug-loading celastrol nanosuspensions and their anti-breast cancer activities in vitro and in vivo. Scientific Reports, 2020, 10, 8851.	3.3	28
8	Preparation of hydroxy genkwanin nanosuspensions and their enhanced antitumor efficacy against breast cancer. Drug Delivery, 2020, 27, 816-824.	5.7	16
9	Nanoadsorbents preparing from oligoethylene glycol dendron and citric acid: Enhanced adsorption effect for the removal of heavy metal ions. Colloids and Surfaces B: Biointerfaces, 2020, 189, 110876.	5.0	10
10	Soybean lecithin stabilizes disulfiram nanosuspensions with a high drug-loading content: remarkably improved antitumor efficacy. Journal of Nanobiotechnology, 2020, 18, 4.	9.1	14
11	Hydrous icaritin nanorods with excellent stability improves the <i>inÂvitro</i> and <i>inÂvivo</i> activity against breast cancer. Drug Delivery, 2020, 27, 228-237.	5.7	10
12	Nanoadsorbents Based on NIPAM and Citric Acid: Removal Efficacy of Heavy Metal Ions in Different Media. ACS Omega, 2019, 4, 14162-14168.	3.5	12
13	Surface modification of pH-sensitive honokiol nanoparticles based on dopamine coating for targeted therapy of breast cancer. Colloids and Surfaces B: Biointerfaces, 2019, 177, 1-10.	5.0	16
14	A comparative study of polydopamine modified and conventional chemical synthesis method in doxorubicin liposomes form the aspect of tumor targeted therapy. International Journal of Pharmaceutics, 2019, 559, 76-85.	5.2	15
15	Hydroxycamptothecin nanoparticles based on poly/oligo (ethylene glycol): Architecture effects of nanocarriers on antitumor efficacy. European Journal of Pharmaceutics and Biopharmaceutics, 2019, 134, 178-184.	4.3	14
16	Surface modification of doxorubicin-loaded nanoparticles based on polydopamine with pH-sensitive property for tumor targeting therapy. Drug Delivery, 2018, 25, 564-575.	5.7	64
17	Folate-targeting annonaceous acetogenins nanosuspensions: significantly enhanced antitumor efficacy in HeLa tumor-bearing mice. Drug Delivery, 2018, 25, 880-887.	5.7	35
18	Shape of Nanoparticles as a Design Parameter to Improve Docetaxel Antitumor Efficacy. Bioconjugate Chemistry, 2018, 29, 1302-1311.	3.6	34

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19	Effect of alkyl chain on cellular uptake and antitumor activity of hydroxycamptothecin nanoparticles based on amphiphilic linear molecules. European Journal of Pharmaceutical Sciences, 2018, 124, 266-272.	4.0	4
20	Administration of raloxifene hydrochloride nanosuspensions partially attenuates bone loss in ovariectomized mice. RSC Advances, 2018, 8, 23748-23756.	3.6	1
21	Intracellular Delivery of Colloidally Stable Core-Cross-Linked Triblock Copolymer Micelles with Glutathione-Responsive Enhanced Drug Release for Cancer Therapy. Molecular Pharmaceutics, 2017, 14, 2518-2528.	4.6	24
22	High drug payload curcumin nanosuspensions stabilized by mPEG-DSPE and SPC: in vitro and in vivo evaluation. Drug Delivery, 2017, 24, 109-120.	5.7	34
23	Honokiol nanoparticles stabilized by oligoethylene glycols codendrimer: in vitro and in vivo investigations. Journal of Materials Chemistry B, 2017, 5, 697-706.	5.8	12
24	Hydroxycamptothecin Nanorods Prepared by Fluorescently Labeled Oligoethylene Glycols (OEG) Codendrimer: Antitumor Efficacy in Vitro and in Vivo. Bioconjugate Chemistry, 2017, 28, 390-399.	3.6	20
25	Regioselective glycosylation of novobiocin alters activity. Carbohydrate Research, 2017, 452, 116-121.	2.3	1
26	Genkwanin nanosuspensions: a novel and potential antitumor drug in breast carcinoma therapy. Drug Delivery, 2017, 24, 1491-1500.	5.7	24
27	Folate-modified Annonaceous acetogenins nanosuspensions and their improved antitumor efficacy. International Journal of Nanomedicine, 2017, Volume 12, 5053-5067.	6.7	23
28	10-Hydroxycamptothecin (HCPT) nanosuspensions stabilized by mPEG ₁₀₀₀ -HCPT conjugate: high stabilizing efficiency and improved antitumor efficacy. International Journal of Nanomedicine, 2017, Volume 12, 3681-3695.	6.7	27
29	A stabilizer-free and organic solvent-free method to prepare 10-hydroxycamptothecin nanocrystals: in vitro and in vivo evaluation. International Journal of Nanomedicine, 2016, 11, 2979.	6.7	27
30	Annonaceous acetogenins nanosuspensions stabilized by PCL–PEG block polymer: significantly improved antitumor efficacy. International Journal of Nanomedicine, 2016, Volume 11, 3239-3253.	6.7	24
31	Annonaceous acetogenins (ACCs) nanosuspensions based on a self-assembly stabilizer and the significantly improved anti-tumor efficacy. Colloids and Surfaces B: Biointerfaces, 2016, 145, 319-327.	5.0	37
32	Honokiol nanosuspensions: Preparation, increased oral bioavailability and dramatically enhanced biodistribution in the cardio-cerebro-vascular system. Colloids and Surfaces B: Biointerfaces, 2014, 116, 114-120.	5.0	45
33	Codendrimer (PAG) from polyamidoamine (PAMAM) and oligoethylene glycols (OEG) dendron: evaluation as drug carrier. Journal of Materials Chemistry B, 2013, 1, 6078.	5.8	10
34	Preparation, characterization, biodistribution and antitumor efficacy of hydroxycamptothecin nanosuspensions. International Journal of Pharmaceutics, 2013, 455, 85-92.	5.2	38