Nilufer Yuksel

List of Publications by Year in descending order

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NILLIFED YILKSEL

#	Article	IF	CITATIONS
1	Molecularly imprinted polymers: preparation, characterisation, and application in drug delivery systems. Journal of Microencapsulation, 2022, 39, 176-196.	1.2	7
2	A novel delivery system for enhancing bioavailability of S-adenosyl-l-methionine: Pectin nanoparticles-in-microparticles and their in vitro - in vivo evaluation'. Journal of Drug Delivery Science and Technology, 2021, 61, 102096.	1.4	11
3	A study to enhance the oral bioavailability of s-adenosyl-l-methionine (SAMe): SLN and SLN nand SLN nanocomposite particles. Chemistry and Physics of Lipids, 2021, 237, 105086.	1.5	10
4	Development and <i>In vitro</i> Evaluation of Nifedipine Gel Formulations for Anorectal Applications. Current Drug Delivery, 2020, 17, 126-139.	0.8	2
5	Characterization and optimization of colon targeted S-adenosyl-L-methionine loaded chitosan nanoparticles. Journal of Research in Pharmacy, 2019, 23, 914-926.	0.1	10
6	Investigations on clonazepam-loaded polymeric micelle-like nanoparticles for safe drug administration during pregnancy. Journal of Microencapsulation, 2018, 35, 149-164.	1.2	9
7	Evaluation of various block copolymers for micelle formation and brain drug delivery: InÂvitro characterization and cellular uptake studies. Journal of Drug Delivery Science and Technology, 2016, 36, 120-129.	1.4	36
8	In situ niosome forming maltodextrin proniosomes of candesartan cilexetil: In vitro and in vivo evaluations. International Journal of Biological Macromolecules, 2016, 82, 453-463.	3.6	39
9	Paclitaxel-loaded niosomes for intravenous administration: pharmacokineticsand tissue distribution in rats. Turkish Journal of Medical Sciences, 2015, 45, 1403-1412.	0.4	26
10	Development and Characterization of Mixed Niosomes for Oral Delivery Using Candesartan Cilexetil as a Model Poorly Water-Soluble Drug. AAPS PharmSciTech, 2015, 16, 108-117.	1.5	83
11	Provesicles as Novel Drug Delivery Systems. Current Pharmaceutical Biotechnology, 2015, 16, 344-364.	0.9	22
12	Stability Studies on Piroxicam Encapsulated Niosomes. Current Drug Delivery, 2015, 12, 192-199.	0.8	42
13	Niosomes encapsulating paclitaxel for oral bioavailability enhancement: preparation, characterization, pharmacokinetics and biodistribution. Journal of Microencapsulation, 2013, 30, 796-804.	1.2	42
14	Investigation of Formulation Variables and Excipient Interaction on the Production of Niosomes. AAPS PharmSciTech, 2012, 13, 826-835.	1.5	67
15	Investigation of triacetin effect on indomethacin release from poly(methyl methacrylate) microspheres: Evaluation of interactions using FT-IR and NMR spectroscopies. International Journal of Pharmaceutics, 2011, 404, 102-109.	2.6	38
16	Characterization of niosomes prepared with various nonionic surfactants for paclitaxel oral delivery. Journal of Pharmaceutical Sciences, 2010, 99, 2049-2060.	1.6	245
17	Preparation and optimization of superabsorbent hydrogel micromatrices based on poly(acrylic acid), partly sodium salt-g-poly(ethylene oxide) for modified release of indomethacin. Drug Development and Industrial Pharmacy, 2009, 35, 756-767.	0.9	3
18	Investigation of pluronic and PEG-PE micelles as carriers of meso-tetraphenyl porphine for oral administration. International Journal of Pharmaceutics, 2007, 332, 161-167.	2.6	67

NILUFER YUKSEL

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19	Preparation and characterization of polymeric micelles for solubilization of poorly soluble anticancer drugs. European Journal of Pharmaceutics and Biopharmaceutics, 2006, 64, 261-268.	2.0	290
20	Influence of aluminum tristearate and sucrose stearate as the dispersing agents on physical properties and release characteristics of Eudragit RS microspheres. AAPS PharmSciTech, 2006, 7, E111-E117.	1.5	18
21	Improved solubility and dissolution rate of piroxicam using gelucire 44/14 and labrasol. Il Farmaco, 2005, 60, 777-782.	0.9	70
22	Comparative Evaluation of Granules Made with Different Binders by a Fluidized Bed Method. Drug Development and Industrial Pharmacy, 2003, 29, 387-395.	0.9	11
23	Enhanced bioavailability of piroxicam using Gelucire 44/14 and Labrasol: in vitro and in vivo evaluation. European Journal of Pharmaceutics and Biopharmaceutics, 2003, 56, 453-459.	2.0	115
24	Comparison of in vitro dissolution profiles by ANOVA-based, model-dependent and -independent methods. International Journal of Pharmaceutics, 2000, 209, 57-67.	2.6	232
25	Influence of Swelling Degree on Release of Nicardipine Hydrochloride from Acrylic Microspheres Prepared by Solvent Evaporation Method. Pharmaceutical Development and Technology, 1998, 3, 115-121.	1.1	15
26	Interaction between nicardipine hydrochloride and polymeric microspheres for a controlled release system. International Journal of Pharmaceutics, 1996, 140, 145-154.	2.6	36
27	The changes in the Mechanic Properties of a direct tableting agent Microcrystalline Cellulose by Precompression. Drug Development and Industrial Pharmacy, 1994, 20, 2323-2331.	0.9	11
28	The preparation of prolonged action formulations in the form of semi solid matrix into hard gelatin capsules of oxprenolol II. Thixocap Method. Drug Development and Industrial Pharmacy, 1992, 18, 233-243.	0.9	4
29	The preparation of prolonged action formulations in the form of semi solid matrix into hard gelatin capsules of oxprenolol I. Thermocap method. Drug Development and Industrial Pharmacy, 1991, 17,	0.9	11