Natascia Mennini

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	4-Heteroaryl Substituted Amino-3,5-Dicyanopyridines as New Adenosine Receptor Ligands: Novel Insights on Structure-Activity Relationships and Perspectives. Pharmaceuticals, 2022, 15, 478.	1.7	4
2	Multiple Roles of Chitosan in Mucosal Drug Delivery: An Updated Review. Marine Drugs, 2022, 20, 335.	2.2	40
3	Improvement of Butamben Anesthetic Efficacy by the Development of Deformable Liposomes Bearing the Drug as Cyclodextrin Complex. Pharmaceutics, 2021, 13, 872.	2.0	8
4	Development of a Cyclodextrin-Based Mucoadhesive-Thermosensitive In Situ Gel for Clonazepam Intranasal Delivery. Pharmaceutics, 2021, 13, 969.	2.0	20
5	Combined Use of Cyclodextrins and Amino Acids for the Development of Cefixime Oral Solutions for Pediatric Use. Pharmaceutics, 2021, 13, 1923.	2.0	7
6	Development of a stable oral pediatric solution of hydrochlorothiazide by the combined use of cyclodextrins and hydrophilic polymers. International Journal of Pharmaceutics, 2020, 587, 119692.	2.6	8
7	Characterization and evaluation of the performance of different calcium and magnesium salts as excipients for direct compression. International Journal of Pharmaceutics, 2019, 567, 118454.	2.6	6
8	In situ mucoadhesive-thermosensitive liposomal gel as a novel vehicle for nasal extended delivery of opiorphin. European Journal of Pharmaceutics and Biopharmaceutics, 2018, 122, 54-61.	2.0	95
9	Design, characterization and in vivo evaluation of nanostructured lipid carriers (NLC) as a new drug delivery system for hydrochlorothiazide oral administration in pediatric therapy. Drug Delivery, 2018, 25, 1910-1921.	2.5	86
10	A preliminary study for the development and optimization by experimental design of an in vitro method for prediction of drug buccal absorption. International Journal of Pharmaceutics, 2018, 547, 530-536.	2.6	9
11	Development and Optimization by Quality by Design Strategies of Frovatriptan Orally Disintegrating Tablets for Migraine Management. Current Drug Delivery, 2018, 15, 436-445.	0.8	3
12	Development and in vivo evaluation of an innovative "Hydrochlorothiazide-in Cyclodextrins-in Solid Lipid Nanoparticles―formulation with sustained release and enhanced oral bioavailability for potential hypertension treatment in pediatrics. International Journal of Pharmaceutics, 2017, 521, 73-83.	2.6	50
13	Development and characterization of fast dissolving tablets of oxaprozin based on hybrid systems of the drug with cyclodextrins and nanoclays. International Journal of Pharmaceutics, 2017, 531, 640-649.	2.6	12
14	Development of cyclodextrin hydrogels for vaginal delivery of dehydroepiandrosterone. Journal of Pharmacy and Pharmacology, 2016, 68, 762-771.	1.2	13
15	Polymeric mucoadhesive tablets for topical or systemic buccal delivery of clonazepam: Effect of cyclodextrin complexation. Carbohydrate Polymers, 2016, 152, 755-763.	5.1	33
16	Quality of wound dressings: a first step in establishing shared criteria and objective procedures to evaluate their performance. Journal of Wound Care, 2016, 25, 428-437.	0.5	23
17	Analysis of physicochemical properties of ternary systems of oxaprozin with randomly methylated-ĀŸ-cyclodextrin and l -arginine aimed to improve the drug solubility. Journal of Pharmaceutical and Biomedical Analysis, 2016, 129, 350-358.	1.4	42
18	Comparison of liposomal and NLC (nanostructured lipid carrier) formulations for improving the transdermal delivery of oxaprozin: Effect of cyclodextrin complexation. International Journal of Pharmaceutics, 2016, 515, 684-691.	2.6	44

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19	Amidated pectin-based wafers for econazole buccal delivery: Formulation optimization and antimicrobial efficacy estimation. Carbohydrate Polymers, 2015, 121, 231-240.	5.1	35
20	Combined use of bile acids and aminoacids to improve permeation properties of acyclovir. International Journal of Pharmaceutics, 2015, 490, 351-359.	2.6	7
21	Injectable liposomal formulations of opiorphin as a new therapeutic strategy in pain management. Future Science OA, 2015, 1, FSO2.	0.9	11
22	Comparative analysis of binary and ternary cyclodextrin complexes with econazole nitrate in solid state. Journal of Pharmaceutical and Biomedical Analysis, 2014, 91, 81-91.	1.4	44
23	Physico-chemical characterization in solution and in the solid state of clonazepam complexes with native and chemically-modified cyclodextrins. Journal of Pharmaceutical and Biomedical Analysis, 2014, 89, 142-149.	1.4	42
24	Development of a chitosan-derivative micellar formulation to improve celecoxib solubility and bioavailability. Drug Development and Industrial Pharmacy, 2014, 40, 1494-1502.	0.9	18
25	Development of liposomal and microemulsion formulations for transdermal delivery of clonazepam: Effect of randomly methylated β-cyclodextrin. International Journal of Pharmaceutics, 2014, 475, 306-314.	2.6	47
26	Development and characterization of functionalized niosomes for brain targeting of dynorphin-B. European Journal of Pharmaceutics and Biopharmaceutics, 2014, 87, 73-79.	2.0	66
27	Comparative study of liposomes, transfersomes and ethosomes as carriers for improving topical delivery of celecoxib. Drug Delivery, 2012, 19, 354-361.	2.5	106
28	Development of a new delivery system consisting in "drug – in cyclodextrin – in nanostructured lipid carriers―for ketoprofen topical delivery. European Journal of Pharmaceutics and Biopharmaceutics, 2012, 80, 46-53.	2.0	123
29	Quality by design approach for developing chitosan-Ca-alginate microspheres for colon delivery of celecoxib-hydroxypropyl-î²-cyclodextrin-PVP complex. European Journal of Pharmaceutics and Biopharmaceutics, 2012, 80, 67-75.	2.0	99
30	New solid self-microemulsifying systems to enhance dissolution rate of poorly water soluble drugs. Pharmaceutical Development and Technology, 2012, 17, 277-284.	1.1	46
31	Influence of cross-linking agent type and chitosan content on the performance of pectinate-chitosan beads aimed for colon-specific drug delivery. Drug Development and Industrial Pharmacy, 2012, 38, 1142-1151.	0.9	28
32	Development and Characterization of Niosomal Formulations of Doxorubicin Aimed at Brain Targeting. Journal of Pharmacy and Pharmaceutical Sciences, 2012, 15, 184.	0.9	66
33	Improvement of oxaprozin solubility and permeability by the combined use of cyclodextrin, chitosan, and bile components. European Journal of Pharmaceutics and Biopharmaceutics, 2011, 78, 385-393.	2.0	43
34	Mixture experiment methods in the development and optimization of microemulsion formulations. Journal of Pharmaceutical and Biomedical Analysis, 2011, 55, 610-617.	1.4	44
35	Development of Mucoadhesive Films for Buccal Administration of Flufenamic Acid: Effect of Cyclodextrin Complexation. Journal of Pharmaceutical Sciences, 2010, 99, 3019-3029.	1.6	46
36	Phase solubility, 1H NMR and molecular modelling studies of bupivacaine hydrochloride complexation with different cyclodextrin derivates. Chemical Physics Letters, 2010, 500, 347-354.	1.2	21

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37	Liposomal formulations of prilocaine: effect of complexation with hydroxypropyl-ß-cyclodextrin on drug anesthetic efficacy. Journal of Liposome Research, 2010, 20, 315-322.	1.5	41
38	Influence of the preparation method on the physical–chemical properties of ketoprofen–cyclodextrin–phosphatidylcholine ternary systems. Journal of Pharmaceutical and Biomedical Analysis, 2009, 50, 690-694.	1.4	31
39	Comparative study of oxaprozin complexation with natural and chemically-modified cyclodextrins in solution and in the solid state. Journal of Inclusion Phenomena and Macrocyclic Chemistry, 2009, 63, 17-25.	1.6	37
40	Physical–chemical characterization of binary and ternary systems of ketoprofen with cyclodextrins and phospholipids. Journal of Pharmaceutical and Biomedical Analysis, 2009, 50, 683-689.	1.4	20
41	Microspheres for colonic delivery of ketoprofen-hydroxypropyl-β-cyclodextrin complex. European Journal of Pharmaceutical Sciences, 2008, 34, 1-11.	1.9	57
42	Response surface methodology in the optimization of chitosan–Ca pectinate bead formulations. European Journal of Pharmaceutical Sciences, 2008, 35, 318-325.	1.9	32
43	Sustained-release matrix tablets of metformin hydrochloride in combination with triacetyl-β-cyclodextrin. European Journal of Pharmaceutics and Biopharmaceutics, 2008, 68, 303-309.	2.0	86
44	Development of enteric-coated calcium pectinate microspheres intended for colonic drug delivery. European Journal of Pharmaceutics and Biopharmaceutics, 2008, 69, 508-518.	2.0	93
45	Comparison of the effect of chitosan and polyvinylpyrrolidone on dissolution properties and analgesic effect of naproxen. European Journal of Pharmaceutics and Biopharmaceutics, 2004, 57, 93-99.	2.0	57
46	Development and characterization of naproxen–chitosan solid systems with improved drug dissolution properties. European Journal of Pharmaceutical Sciences, 2003, 19, 67-75.	1.9	77