Paola Mura

List of Publications by Year in descending order

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74163 34105 8,021 187 52 75 citations h-index g-index papers 188 188 188 7016 docs citations times ranked citing authors all docs

#	Article	IF	CITATIONS
1	Role of Cyclodextrins and Drug Solid State Properties on Flufenamic Acid Dissolution Performance from Tablets. Pharmaceutics, 2022, 14, 284.	4.5	6
2	Multiple Roles of Chitosan in Mucosal Drug Delivery: An Updated Review. Marine Drugs, 2022, 20, 335.	4.6	40
3	Evaluation and Comparison of Solid Lipid Nanoparticles (SLNs) and Nanostructured Lipid Carriers (NLCs) as Vectors to Develop Hydrochlorothiazide Effective and Safe Pediatric Oral Liquid Formulations. Pharmaceutics, 2021, 13, 437.	4.5	53
4	Development and microbiological evaluation of chitosan and chitosan-alginate microspheres for vaginal administration of metronidazole. International Journal of Pharmaceutics, 2021, 598, 120375.	5.2	27
5	Improvement of Butamben Anesthetic Efficacy by the Development of Deformable Liposomes Bearing the Drug as Cyclodextrin Complex. Pharmaceutics, 2021, 13, 872.	4. 5	8
6	Development of a Cyclodextrin-Based Mucoadhesive-Thermosensitive In Situ Gel for Clonazepam Intranasal Delivery. Pharmaceutics, 2021, 13, 969.	4. 5	20
7	Development of a Near Infrared Spectroscopy method for the in-line quantitative bilastine drug determination during pharmaceutical powders blending. Journal of Pharmaceutical and Biomedical Analysis, 2021, 204, 114277.	2.8	11
8	Combined Use of Cyclodextrins and Amino Acids for the Development of Cefixime Oral Solutions for Pediatric Use. Pharmaceutics, 2021, 13, 1923.	4. 5	7
9	Preparation, Characterization and Evaluation of the Anti-Inflammatory Activity of Epichlorohydrin-Î ² -Cyclodextrin/Curcumin Binary Systems Embedded in a Pluronic®/Hyaluronate Hydrogel. International Journal of Molecular Sciences, 2021, 22, 13566.	4.1	8
10	The role of solid state properties on the dissolution performance of flufenamic acid. Journal of Pharmaceutical and Biomedical Analysis, 2020, 180 , 113058 .	2.8	10
11	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.	5.2	8
12	Development and Characterization of Liquisolid Tablets Based on Mesoporous Clays or Silicas for Improving Glyburide Dissolution. Pharmaceutics, 2020, 12, 503.	4. 5	9
13	Advantages of the combined use of cyclodextrins and nanocarriers in drug delivery: A review. International Journal of Pharmaceutics, 2020, 579, 119181.	5.2	53
14	Tablets of "Hydrochlorothiazide in Cyclodextrin in Nanoclayâ€. A New Nanohybrid System with Enhanced Dissolution Properties. Pharmaceutics, 2020, 12, 104.	4. 5	10
15	Curcumin-in-Cyclodextrins-in-Liposomes: An Alternative for Osteoarthritis Treatment. Proceedings (mdpi), 2020, 78, .	0.2	1
16	Characterization and evaluation of the performance of different calcium and magnesium salts as excipients for direct compression. International Journal of Pharmaceutics, 2019, 567, 118454.	5. 2	6
17	Characterization and evaluation of different mesoporous silica kinds as carriers for the development of effective oral dosage forms of glibenclamide. International Journal of Pharmaceutics, 2019, 563, 43-52.	5. 2	18
18	Characterization and microbiological evaluation of chitosan-alginate microspheres for cefixime vaginal administration. Carbohydrate Polymers, 2018, 192, 176-183.	10.2	32

#	Article	IF	CITATIONS
19	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.	4.3	95
20	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.	5.7	86
21	Novel Findings about Double-Loaded Curcumin-in-HPβcyclodextrin-in Liposomes: Effects on the Lipid Bilayer and Drug Release. Pharmaceutics, 2018, 10, 256.	4.5	32
22	Combined Approach of Cyclodextrin Complexationand Nanostructured Lipid Carriers for the Development of a Pediatric Liquid Oral Dosage Form of Hydrochlorothiazide. Pharmaceutics, 2018, 10, 287.	4.5	17
23	Grinding as Solvent-Free Green Chemistry Approach for Cyclodextrin Inclusion Complex Preparation in the Solid State. Pharmaceutics, 2018, 10, 189.	4.5	56
24	Improving the therapeutic efficacy of prilocaine by PLGA microparticles: Preparation, characterization and in vivo evaluation. International Journal of Pharmaceutics, 2018, 547, 24-30.	5 . 2	24
25	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.	5. 2	9
26	Development and Optimization by Quality by Design Strategies of Frovatriptan Orally Disintegrating Tablets for Migraine Management. Current Drug Delivery, 2018, 15, 436-445.	1.6	3
27	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.	5. 2	50
28	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.	5.2	12
29	Calcium alginate microspheres containing metformin hydrochloride niosomes and chitosomes aimed for oral therapy of type 2 diabetes mellitus. International Journal of Pharmaceutics, 2017, 530, 430-439.	5. 2	43
30	Development of cyclodextrin hydrogels for vaginal delivery of dehydroepiandrosterone. Journal of Pharmacy and Pharmacology, 2016, 68, 762-771.	2.4	13
31	Development and characterization of fast-dissolving tablet formulations of glyburide based on solid self-microemulsifying systems. European Journal of Pharmaceutics and Biopharmaceutics, 2016, 104, 19-29.	4.3	23
32	Polymeric mucoadhesive tablets for topical or systemic buccal delivery of clonazepam: Effect of cyclodextrin complexation. Carbohydrate Polymers, 2016, 152, 755-763.	10.2	33
33	Analysis of physicochemical properties of ternary systems of oxaprozin with randomly methylated-ÃY-cyclodextrin and I -arginine aimed to improve the drug solubility. Journal of Pharmaceutical and Biomedical Analysis, 2016, 129, 350-358.	2.8	42
34	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.	5.2	44
35	Hybrid systems based on "drug – in cyclodextrin – in nanoclays―for improving oxaprozin dissolution properties. International Journal of Pharmaceutics, 2016, 509, 8-15.	5.2	36
36	Development of solid lipid nanoparticles as carriers for improving oral bioavailability of glibenclamide. European Journal of Pharmaceutics and Biopharmaceutics, 2016, 102, 41-50.	4.3	80

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37	Advanced formulations for improving therapies with anti-inflammatory or anaesthetic drugs: A review. Journal of Drug Delivery Science and Technology, 2016, 32, 192-205.	3.0	20
38	Comparative evaluation of polymeric and waxy microspheres for combined colon delivery of ascorbic acid and ketoprofen. International Journal of Pharmaceutics, 2015, 485, 365-373.	5.2	30
39	Amidated pectin-based wafers for econazole buccal delivery: Formulation optimization and antimicrobial efficacy estimation. Carbohydrate Polymers, 2015, 121, 231-240.	10.2	35
40	Analytical techniques for characterization of cyclodextrin complexes in the solid state: A review. Journal of Pharmaceutical and Biomedical Analysis, 2015, 113, 226-238.	2.8	215
41	Combined use of bile acids and aminoacids to improve permeation properties of acyclovir. International Journal of Pharmaceutics, 2015, 490, 351-359.	5.2	7
42	Fast analysis of glibenclamide and its impurities: quality by design framework in capillary electrophoresis method development. Analytical and Bioanalytical Chemistry, 2015, 407, 7637-7646.	3.7	16
43	Development and ex vivo evaluation of 5-aminolevulinic acid-loaded niosomal formulations for topical photodynamic therapy. International Journal of Pharmaceutics, 2015, 494, 258-263.	5.2	27
44	Cyclodextrin complexation highly enhances efficacy of arylsulfonylureido benzenesulfonamide carbonic anhydrase inhibitors as a topical antiglaucoma agents. Bioorganic and Medicinal Chemistry, 2015, 23, 6223-6227.	3.0	10
45	Injectable liposomal formulations of opiorphin as a new therapeutic strategy in pain management. Future Science OA, 2015, 1, FSO2.	1.9	11
46	Comparative analysis of binary and ternary cyclodextrin complexes with econazole nitrate in solution and in solid state. Journal of Pharmaceutical and Biomedical Analysis, 2014, 91, 81-91.	2.8	44
47	Analytical techniques for characterization of cyclodextrin complexes in aqueous solution: A review. Journal of Pharmaceutical and Biomedical Analysis, 2014, 101, 238-250.	2.8	224
48	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.	2.8	42
49	Development of a chitosan-derivative micellar formulation to improve celecoxib solubility and bioavailability. Drug Development and Industrial Pharmacy, 2014, 40, 1494-1502.	2.0	18
50	Development of liposomal and microemulsion formulations for transdermal delivery of clonazepam: Effect of randomly methylated \hat{l}^2 -cyclodextrin. International Journal of Pharmaceutics, 2014, 475, 306-314.	5.2	47
51	Development and characterization of functionalized niosomes for brain targeting of dynorphin-B. European Journal of Pharmaceutics and Biopharmaceutics, 2014, 87, 73-79.	4.3	66
52	Selection of PLA polymers for the development of injectable prilocaine controlled release microparticles: Usefulness of thermal analysis. International Journal of Pharmaceutics, 2013, 441, 468-475.	5.2	28
53	Native and polymeric \hat{l}^2 -cyclodextrins in performance improvement of chitosan films aimed for buccal delivery of poorly soluble drugs. Journal of Inclusion Phenomena and Macrocyclic Chemistry, 2012, 74, 87-97.	1.6	21
54	Comparative study of liposomes, transfersomes and ethosomes as carriers for improving topical delivery of celecoxib. Drug Delivery, 2012, 19, 354-361.	5.7	106

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55	Development of low methoxy amidated pectin-based mucoadhesive patches for buccal delivery of triclosan: Effect of cyclodextrin complexation. Carbohydrate Polymers, 2012, 90, 1794-1803.	10.2	30
56	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.	4.3	123
57	Quality by design approach for developing chitosan-Ca-alginate microspheres for colon delivery of celecoxib-hydroxypropyl- \hat{l}^2 -cyclodextrin-PVP complex. European Journal of Pharmaceutics and Biopharmaceutics, 2012, 80, 67-75.	4.3	99
58	New solid self-microemulsifying systems to enhance dissolution rate of poorly water soluble drugs. Pharmaceutical Development and Technology, 2012, 17, 277-284.	2.4	46
59	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.	2.0	28
60	Development and Characterization of Niosomal Formulations of Doxorubicin Aimed at Brain Targeting. Journal of Pharmacy and Pharmaceutical Sciences, 2012, 15, 184.	2.1	66
61	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.	4.3	43
62	Analysis of triclosan inclusion complexes with \hat{l}^2 -cyclodextrin and its water-soluble polymeric derivative. Journal of Pharmaceutical and Biomedical Analysis, 2011, 54, 1030-1039.	2.8	73
63	Mixture experiment methods in the development and optimization of microemulsion formulations. Journal of Pharmaceutical and Biomedical Analysis, 2011, 55, 610-617.	2.8	44
64	New "drug-in cyclodextrin-in deformable liposomes―formulations to improve the therapeutic efficacy of local anaesthetics. International Journal of Pharmaceutics, 2010, 395, 222-231.	5.2	81
65	Physical chemical characterization of binary systems of prilocaine hydrochloride with triacetyl-Î ² -cyclodextrin. Journal of Inclusion Phenomena and Macrocyclic Chemistry, 2010, 68, 437-445.	1.6	22
66	Development of Mucoadhesive Films for Buccal Administration of Flufenamic Acid: Effect of Cyclodextrin Complexation. Journal of Pharmaceutical Sciences, 2010, 99, 3019-3029.	3.3	46
67	Preparation and solid-state characterization of bupivacaine hydrochloride cyclodextrin complexes aimed for buccal delivery. Journal of Pharmaceutical and Biomedical Analysis, 2010, 52, 9-18.	2.8	60
68	Phase solubility, 1H NMR and molecular modelling studies of bupivacaine hydrochloride complexation with different cyclodextrin derivates. Chemical Physics Letters, 2010, 500, 347-354.	2.6	21
69	Development of a new delivery system consisting in †drug†in cyclodextrin†in PLGA nanoparticlesâ€. Journal of Microencapsulation, 2010, 27, 479-486.	2.8	22
70	Liposomal formulations of prilocaine: effect of complexation with hydroxypropyl-ß-cyclodextrin on drug anesthetic efficacy. Journal of Liposome Research, 2010, 20, 315-322.	3.3	41
71	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.	2.8	31
72	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

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73	Physical–chemical characterization of binary and ternary systems of ketoprofen with cyclodextrins and phospholipids. Journal of Pharmaceutical and Biomedical Analysis, 2009, 50, 683-689.	2.8	20
74	Effect of preparation technique on the properties and <i>in vivo </i> efficacy of benzocaine-loaded ethosomes. Journal of Liposome Research, 2009, 19, 253-260.	3 . 3	68
75	Development of Glyburide Fast-Dissolving Tablets Based on the Combined Use of Cyclodextrins and Polymers. Drug Development and Industrial Pharmacy, 2009, 35, 73-82.	2.0	21
76	Microspheres for colonic delivery of ketoprofen-hydroxypropyl-β-cyclodextrin complex. European Journal of Pharmaceutical Sciences, 2008, 34, 1-11.	4.0	57
77	Response surface methodology in the optimization of chitosan–Ca pectinate bead formulations. European Journal of Pharmaceutical Sciences, 2008, 35, 318-325.	4.0	32
78	Sustained-release matrix tablets of metformin hydrochloride in combination with triacetyl-Î ² -cyclodextrin. European Journal of Pharmaceutics and Biopharmaceutics, 2008, 68, 303-309.	4.3	86
79	Development of enteric-coated calcium pectinate microspheres intended for colonic drug delivery. European Journal of Pharmaceutics and Biopharmaceutics, 2008, 69, 508-518.	4.3	93
80	Optimization of Formulation Variables of Benzocaine Liposomes using Experimental Design. Journal of Liposome Research, 2008, 18, 113-125.	3.3	25
81	Dissolution and Permeation Properties of Naproxen From Solid-State Systems With Chitosan. Drug Delivery, 2008, 15, 303-312.	5 . 7	18
82	Development, characterization and in vivo evaluation of benzocaine-loaded liposomes. European Journal of Pharmaceutics and Biopharmaceutics, 2007, 67, 86-95.	4.3	137
83	Fast-Dissolving Tablets of Glyburide Based on Ternary Solid Dispersions with PEG 6000 and Surfactants. Drug Delivery, 2007, 14, 247-255.	5 . 7	25
84	Liquid spray formulations of xibornol by using self-microemulsifying drug delivery systems. International Journal of Pharmaceutics, 2007, 340, 84-91.	5.2	59
85	Physical–chemical characterization of binary systems of metformin hydrochloride with triacetyl-β-cyclodextrin. Journal of Pharmaceutical and Biomedical Analysis, 2007, 45, 480-486.	2.8	44
86	Physicochemical characterization of drug-cyclodextrin complexes prepared by supercritical carbon dioxide and by conventional techniques. Journal of Inclusion Phenomena and Macrocyclic Chemistry, 2007, 57, 223-231.	1.6	28
87	Development of a sustained-release matrix tablet formulation of DHEA as ternary complex with \hat{l}_{\pm} -cyclodextrin and glycine. Journal of Inclusion Phenomena and Macrocyclic Chemistry, 2007, 57, 699-704.	1.6	2
88	The influence of chitosan on cyclodextrin complexing and solubilizing abilities towards drugs. Journal of Inclusion Phenomena and Macrocyclic Chemistry, 2007, 59, 307-313.	1.6	17
89	Evaluation of supercritical fluid technology as preparative technique of benzocaine–cyclodextrin complexes—Comparison with conventional methods. Journal of Pharmaceutical and Biomedical Analysis, 2007, 43, 566-574.	2.8	45
90	Study of formulation variables influencing the drug release rate from matrix tablets by experimental design. European Journal of Pharmaceutics and Biopharmaceutics, 2006, 62, 77-84.	4.3	55

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91	Influence of cyclodextrins and chitosan, separately or in combination, on glyburide solubility and permeability. European Journal of Pharmaceutics and Biopharmaceutics, 2006, 62, 241-246.	4.3	48
92	A new drug nanocarrier consisting of chitosan and hydoxypropylcyclodextrin. European Journal of Pharmaceutics and Biopharmaceutics, 2006, 63, 79-86.	4.3	113
93	Effect of preparation technique on the properties of liposomes encapsulating ketoprofen–cyclodextrin complexes aimed for transdermal delivery. International Journal of Pharmaceutics, 2006, 312, 53-60.	5.2	138
94	Simultaneous effect of cyclodextrin complexation, pH, and hydrophilic polymers on naproxen solubilization. Journal of Pharmaceutical and Biomedical Analysis, 2006, 42, 126-131.	2.8	63
95	Differential scanning calorimetry as a screening technique in compatibility studies of DHEA extended release formulations. Journal of Pharmaceutical and Biomedical Analysis, 2006, 42, 3-10.	2.8	41
96	Development and evaluation of an in vitro method for prediction of human drug absorption. European Journal of Pharmaceutical Sciences, 2006, 27, 346-353.	4.0	39
97	Development and evaluation of an in vitro method for prediction of human drug absorption. European Journal of Pharmaceutical Sciences, 2006, 27, 354-362.	4.0	88
98	Mixture design in the optimization of a microemulsion system for the electrokinetic chromatographic determination of ketorolac and its impurities: Method development and validation. Electrophoresis, 2006, 27, 805-818.	2.4	33
99	Interaction of naproxen with ionic cyclodextrins in aqueous solution and in the solid state. Journal of Pharmaceutical and Biomedical Analysis, 2005, 37, 987-994.	2.8	40
100	Determination of stability constant values of flurbiprofen–cyclodextrin complexes using different techniques. Journal of Pharmaceutical and Biomedical Analysis, 2005, 37, 995-1002.	2.8	43
101	Optimization of glibenclamide tablet composition through the combined use of differential scanning calorimetry and d-optimal mixture experimental design. Journal of Pharmaceutical and Biomedical Analysis, 2005, 37, 65-71.	2.8	47
102	Preparation and characterisation of liposomes encapsulating ketoprofen–cyclodextrin complexes for transdermal drug delivery. International Journal of Pharmaceutics, 2005, 298, 55-67.	5.2	181
103	Comparative Study on Triclosan Interactions in Solution and in the Solid State with Natural and Chemically Modified Cyclodextrins. Journal of Inclusion Phenomena and Macrocyclic Chemistry, 2005, 53, 77-83.	1.6	25
104	Development of Fast-Dissolving Tablets of Flurbiprofen-Cyclodextrin Complexes. Drug Development and Industrial Pharmacy, 2005, 31, 697-707.	2.0	45
105	Solid-state characterization and dissolution properties of Naproxen–Arginine–Hydroxypropyl-β-cyclodextrin ternary system. European Journal of Pharmaceutics and Biopharmaceutics, 2005, 59, 99-106.	4.3	83
106	Influence of formulation and process variables on in vitro release of theophylline from directly-compressed Eudragit matrix tablets. Il Farmaco, 2005, 60, 913-918.	0.9	66
107	Characterization and Dissolution Properties of Ketoprofen in Binary and Ternary Solid Dispersions with Polyethylene Glycol and Surfactants. Drug Development and Industrial Pharmacy, 2005, 31, 425-434.	2.0	43
108	Solid-state characterization of glyburide-cyclodextrin co-ground products. Journal of Thermal Analysis and Calorimetry, 2004, 77, 413-422.	3.6	22

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109	Characterization of the solid phases of paracetamol and fenamates at equilibrium in saturated solutions. Journal of Thermal Analysis and Calorimetry, 2004, 77, 541-554.	3.6	27
110	Photostability studies on nicardipine–cyclodextrin complexes by capillary electrophoresis. Journal of Pharmaceutical and Biomedical Analysis, 2004, 35, 267-275.	2.8	29
111	Influence of chitosan and its glutamate and hydrochloride salts on naproxen dissolution rate and permeation across Caco-2 cells. International Journal of Pharmaceutics, 2004, 271, 257-267.	5 . 2	58
112	Influence of solvent composition on the solid phase at equilibrium with saturated solutions of quinolones in different solvent mixtures. Journal of Pharmaceutical and Biomedical Analysis, 2004, 35, 715-726.	2.8	12
113	Characterization of Ibuproxam Binary and Ternary Dispersions with Hydrophilic Carriers. Drug Development and Industrial Pharmacy, 2004, 30, 65-74.	2.0	44
114	Development of Enteric-coated Timed-release Matrix Tablets for Colon Targeting. Journal of Drug Targeting, 2004, 12, 607-612.	4.4	43
115	Formulation and characterization of triclosan sub-micron emulsions and nanocapsules. Journal of Microencapsulation, 2004, 21, 857-864.	2.8	36
116	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.	4.3	57
117	Development and Evaluation of Glyburide Fast Dissolving Tablets Using Solid Dispersion Technique. Drug Development and Industrial Pharmacy, 2004, 30, 525-534.	2.0	77
118	Title is missing!. Journal of Thermal Analysis and Calorimetry, 2003, 73, 635-646.	3.6	50
119	How experimental design can improve the validation process. Studies in pharmaceutical analysis. Analytical and Bioanalytical Chemistry, 2003, 377, 937-944.	3.7	37
120	Development and characterization of naproxen–chitosan solid systems with improved drug dissolution properties. European Journal of Pharmaceutical Sciences, 2003, 19, 67-75.	4.0	77
121	In vitro release of sodium diclofenac from a central core matrix tablet aimed for colonic drug delivery. European Journal of Pharmaceutical Sciences, 2003, 20, 125-131.	4.0	40
122	Enhancement of Dehydroepiandrosterone Solubility and Bioavailability by Ternary Complexation with $\hat{l}\pm\hat{a}\in\mathbb{C}$ yclodextrin and Glycine. Journal of Pharmaceutical Sciences, 2003, 92, 2177-2184.	3.3	31
123	New docking CFF91 parameters specific for cyclodextrin inclusion complexes. Chemical Physics Letters, 2003, 370, 280-292.	2.6	14
124	Optimization of dissolution test precision for a ketoprofen oral extended-release product. Journal of Pharmaceutical and Biomedical Analysis, 2003, 32, 159-165.	2.8	30
125	Ternary systems of naproxen with hydroxypropyl- \hat{l}^2 -cyclodextrin and aminoacids. International Journal of Pharmaceutics, 2003, 260, 293-302.	5.2	105
126	Development of Enteric-coated Pectin-based Matrix Tablets for Colonic Delivery of Theophylline. Journal of Drug Targeting, 2003, 11, 365-371.	4.4	54

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127	Comparative study of ibuproxam complexation with amorphous \hat{l}^2 -cyclodextrin derivatives in solution and in the solid state. European Journal of Pharmaceutics and Biopharmaceutics, 2002, 54, 181-191.	4.3	32
128	Interaction of naproxen with noncrystalline acetyl \hat{l}^2 - and acetyl \hat{l}^3 -cyclodextrins in the solid and liquid state. European Journal of Pharmaceutical Sciences, 2002, 15, 21-29.	4.0	31
129	Cyclodextrin Complexes of Sulfonamide Carbonic Anhydrase IOnhibitors As Longâ€lasting Topically Acting Antiglaucoma Agents. Journal of Pharmaceutical Sciences, 2002, 91, 2211-2219.	3.3	16
130	Characterization of physicochemical properties of naproxen systems with amorphous \hat{l}^2 -cyclodextrin-epichlorohydrin polymers. Journal of Pharmaceutical and Biomedical Analysis, 2002, 29, 1015-1024.	2.8	79
131	Investigation of the effects of grinding and co-grinding on physicochemical properties of glisentide. Journal of Pharmaceutical and Biomedical Analysis, 2002, 30, 227-237.	2.8	74
132	Assessment of solid-state interactions of naproxen with amorphous cyclodextrin derivatives by DSC. Journal of Pharmaceutical and Biomedical Analysis, 2002, 30, 1173-1179.	2.8	24
133	Didanosine extended-release matrix tablets: optimization of formulation variables using statistical experimental design. International Journal of Pharmaceutics, 2002, 237, 107-118.	5.2	69
134	Development of sustained release matrix tablets of didanosine containing methacrylic and ethylcellulose polymers. International Journal of Pharmaceutics, 2002, 234, 213-221.	5.2	47
135	Computer-aided molecular modeling techniques for predicting the stability of drug–cyclodextrin inclusion complexes in aqueous solutions. Chemical Physics Letters, 2002, 358, 383-390.	2.6	54
136	Compatibility Studies of Multicomponent Tablet Formulations. DSC and experimental mixture design. Magyar AprÁ³vad Közlemények, 2002, 68, 541-551.	1.4	38
137	Effects of Grinding with Microcrystalline Cellulose and Cyclodextrins on the Ketoprofen Physicochemical Properties. Drug Development and Industrial Pharmacy, 2001, 27, 119-128.	2.0	66
138	Interaction of Naproxen with Crystalline and Amorphous Methylated \hat{l}^2 -Cyclodextrin in the Liquid and Solid State. Supramolecular Chemistry, 2001, 12, 379-389.	1.2	17
139	Multicomponent Systems of Econazole with Hydroxyacids and Cyclodextrins. Journal of Inclusion Phenomena and Macrocyclic Chemistry, 2001, 39, 131-138.	1.6	38
140	The influence of polyvinylpyrrolidone on naproxen complexation with hydroxypropyl- \hat{l}^2 -cyclodextrin. European Journal of Pharmaceutical Sciences, 2001, 13, 187-194.	4.0	138
141	1H-NMR and molecular modelling techniques for the investigation of the inclusion complex of econazole with î±-cyclodextrin in the presence of malic acid. Journal of Pharmaceutical and Biomedical Analysis, 2000, 23, 25-31.	2.8	41
142	Evaluation of transcutol as a clonazepam transdermal permeation enhancer from hydrophilic gel formulations. European Journal of Pharmaceutical Sciences, 2000, 9, 365-372.	4.0	122
143	Differential Scanning Calorimetry Analysis of Crystallinity Changes of Naproxen in Ground Mixtures with Maltohexaose, the Non Cyclic Analog of Alpha-Cyclodextrin. , 1999, , 367-370.		1
144	Influence of the preparation method on the physicochemical properties of ketoprofen–cyclodextrin binary systems. International Journal of Pharmaceutics, 1999, 179, 117-128.	5.2	88

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145	Influence of the preparation method on the physicochemical properties of binary systems of econazole with cyclodextrins. International Journal of Pharmaceutics, 1999, 193, 85-95.	5.2	25
146	Physical characterization of picotamide monohydrate and anhydrous picotamide. Journal of Pharmaceutical Sciences, 1999, 88, 1133-1139.	3.3	18
147	Interaction of naproxen with alpha-cyclodextrin and its noncyclic analog maltohexaose. Pharmaceutical Research, 1999, 16, 689-694.	3.5	26
148	Thermal Behavior and Dissolution Properties of Naproxen From Binary and Ternary Solid Dispersions. Drug Development and Industrial Pharmacy, 1999, 25, 257-264.	2.0	73
149	Effects of the Host Cavity Size and the Preparation Method on the Physicochemical Properties of Ibuproxam-Cyclodextrin Systems. Drug Development and Industrial Pharmacy, 1999, 25, 279-287.	2.0	68
150	Dissolution Rate and Thermal Properties of Naproxen in Mixtures with Amorphous or Crystalline Dimethyl Beta-Cyclodextrin., 1999,, 371-374.		1
151	In vitro studies of simulated percutaneous absorption: Influence of various enhancers in the release of clonazepam from 2-hydroxyethyl acetate patches. Pharmaceutica Acta Helvetiae, 1998, 72, 263-269.	1.2	6
152	Differential scanning calorimetry in compatibility testing of picotamide with pharmaceutical excipients. Thermochimica Acta, 1998, 321, 59-65.	2.7	33
153	Interactions of ketoprofen and ibuprofen with \hat{l}^2 -cyclodextrins in solution and in the solid state. International Journal of Pharmaceutics, 1998, 166, 189-203.	5.2	166
154	Development and validation of a differential pulse polarographic method for quinolinic acid determination in human plasma and urine after solid-phase extraction: a chemometric approach. Journal of Pharmaceutical and Biomedical Analysis, 1998, 17, 1015-1028.	2.8	25
155	Compatibility study between ibuproxam and pharmaceutical excipients using differential scanning calorimetry, hot-stage microscopy and scanning electron microscopy. Journal of Pharmaceutical and Biomedical Analysis, 1998, 18, 151-163.	2.8	93
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157	Near-infrared Reflectance Spectrometry in the Studyof AtopyPart 2. Interactions Between the Skin and Polyethylene Glycol 400, Isopropyl Myristate and Hydrogel. Analyst, The, 1997, 122, 771-776.	3.5	10
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