Marzia Cirri

List of Publications by Year in descending order

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87888 175258 3,064 76 38 52 h-index citations g-index papers 77 77 77 3346 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 | Combined Use of Cyclodextrins and Amino Acids for the Development of Cefixime Oral Solutions for Pediatric Use. Pharmaceutics, 2021, 13, 1923. | 4.5 | 7 |
| 8 | Amine- and Amino Acid-Based Compounds as Carbonic Anhydrase Activators. Molecules, 2021, 26, 7331. | 3.8 | 9 |
| 9 | 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 |
| 10 | Development and Characterization of Liquisolid Tablets Based on Mesoporous Clays or Silicas for Improving Glyburide Dissolution. Pharmaceutics, 2020, 12, 503. | 4. 5 | 9 |
| 11 | Tablets of "Hydrochlorothiazide in Cyclodextrin in Nanoclay― A New Nanohybrid System with Enhanced Dissolution Properties. Pharmaceutics, 2020, 12, 104. | 4.5 | 10 |
| 12 | Î ² -Sitosterol Loaded Nanostructured Lipid Carrier: Physical and Oxidative Stability, In Vitro Simulated Digestion and Hypocholesterolemic Activity. Pharmaceutics, 2020, 12, 386. | 4.5 | 13 |
| 13 | 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 |
| 14 | Characterization and microbiological evaluation of chitosan-alginate microspheres for cefixime vaginal administration. Carbohydrate Polymers, 2018, 192, 176-183. | 10.2 | 32 |
| 15 | 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 |
| 16 | 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 |
| 17 | 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 |
| 18 | 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 |

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| 19 | 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 |
| 20 | 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 |
| 21 | Development of cyclodextrin hydrogels for vaginal delivery of dehydroepiandrosterone. Journal of Pharmacy and Pharmacology, 2016, 68, 762-771. | 2.4 | 13 |
| 22 | 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 |
| 23 | Polymeric mucoadhesive tablets for topical or systemic buccal delivery of clonazepam: Effect of cyclodextrin complexation. Carbohydrate Polymers, 2016, 152, 755-763. | 10.2 | 33 |
| 24 | Analysis of physicochemical properties of ternary systems of oxaprozin with randomly methylated-ß-cyclodextrin and I -arginine aimed to improve the drug solubility. Journal of Pharmaceutical and Biomedical Analysis, 2016, 129, 350-358. | 2.8 | 42 |
| 25 | 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 |
| 26 | 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 |
| 27 | Combined use of bile acids and aminoacids to improve permeation properties of acyclovir. International Journal of Pharmaceutics, 2015, 490, 351-359. | 5. 2 | 7 |
| 28 | 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 |
| 29 | Comparative study of liposomes, transfersomes and ethosomes as carriers for improving topical delivery of celecoxib. Drug Delivery, 2012, 19, 354-361. | 5.7 | 106 |
| 30 | 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 |
| 31 | Quality by design approach for developing chitosan-Ca-alginate microspheres for colon delivery of celecoxib-hydroxypropyl- 12 -cyclodextrin-PVP complex. European Journal of Pharmaceutics and Biopharmaceutics, 2012, 80, 67-75. | 4.3 | 99 |
| 32 | 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 |
| 33 | 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 |
| 34 | 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 |
| 35 | Mixture experiment methods in the development and optimization of microemulsion formulations. Journal of Pharmaceutical and Biomedical Analysis, 2011, 55, 610-617. | 2.8 | 44 |
| 36 | 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 |

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| 37 | 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 |
| 38 | 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 |
| 39 | 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 |
| 40 | 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 |
| 41 | Microspheres for colonic delivery of ketoprofen-hydroxypropyl- \hat{l}^2 -cyclodextrin complex. European Journal of Pharmaceutical Sciences, 2008, 34, 1-11. | 4.0 | 57 |
| 42 | 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 |
| 43 | 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 |
| 44 | Dissolution and Permeation Properties of Naproxen From Solid-State Systems With Chitosan. Drug Delivery, 2008, 15, 303-312. | 5.7 | 18 |
| 45 | Fast-Dissolving Tablets of Glyburide Based on Ternary Solid Dispersions with PEG 6000 and Surfactants. Drug Delivery, 2007, 14, 247-255. | 5.7 | 25 |
| 46 | Liquid spray formulations of xibornol by using self-microemulsifying drug delivery systems. International Journal of Pharmaceutics, 2007, 340, 84-91. | 5.2 | 59 |
| 47 | 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 |
| 48 | 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 |
| 49 | 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 |
| 50 | 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 |
| 51 | 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 |
| 52 | 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 |
| 53 | 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 |
| 54 | 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 |

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| 55 | 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 |
| 56 | 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 |
| 57 | 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 |
| 58 | 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 |
| 59 | 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 |
| 60 | 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 |
| 61 | Development of Fast-Dissolving Tablets of Flurbiprofen-Cyclodextrin Complexes. Drug Development and Industrial Pharmacy, 2005, 31, 697-707. | 2.0 | 45 |
| 62 | 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 |
| 63 | 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 |
| 64 | 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 |
| 65 | Solid-state characterization of glyburide-cyclodextrin co-ground products. Journal of Thermal Analysis and Calorimetry, 2004, 77, 413-422. | 3.6 | 22 |
| 66 | 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 |
| 67 | Photostability studies on nicardipine–cyclodextrin complexes by capillary electrophoresis. Journal of Pharmaceutical and Biomedical Analysis, 2004, 35, 267-275. | 2.8 | 29 |
| 68 | 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 |
| 69 | Characterization of Ibuproxam Binary and Ternary Dispersions with Hydrophilic Carriers. Drug Development and Industrial Pharmacy, 2004, 30, 65-74. | 2.0 | 44 |
| 70 | Development of Enteric-coated Timed-release Matrix Tablets for Colon Targeting. Journal of Drug Targeting, 2004, 12, 607-612. | 4.4 | 43 |
| 71 | Development and Evaluation of Glyburide Fast Dissolving Tablets Using Solid Dispersion Technique. Drug Development and Industrial Pharmacy, 2004, 30, 525-534. | 2.0 | 77 |
| 72 | Title is missing!. Journal of Thermal Analysis and Calorimetry, 2003, 73, 635-646. | 3.6 | 50 |

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| 73 | Enhancement of Dehydroepiandrosterone Solubility and Bioavailability by Ternary Complexation with αâ€Cyclodextrin and Glycine. Journal of Pharmaceutical Sciences, 2003, 92, 2177-2184. | 3.3 | 31 |
| 74 | Ternary systems of naproxen with hydroxypropyl- \hat{l}^2 -cyclodextrin and aminoacids. International Journal of Pharmaceutics, 2003, 260, 293-302. | 5.2 | 105 |
| 75 | Development of Enteric-coated Pectin-based Matrix Tablets for Colonic Delivery of Theophylline. Journal of Drug Targeting, 2003, 11, 365-371. | 4.4 | 54 |
| 76 | 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 |