Antonio M Rabasco

List of Publications by Year in descending order

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74 papers 2,345 citations

218677 26 h-index 214800 47 g-index

75 all docs

75 docs citations

75 times ranked 2690 citing authors

#	Article	IF	CITATIONS
1	Central composite design optimization for a controlled valsartan release from polycaprolactone microspheres. Journal of Applied Polymer Science, 2022, 139, 51584.	2.6	1
2	pH-temperature dual-sensitive nucleolipid-containing stealth liposomes anchored with PEGylated AuNPs for triggering delivery of doxorubicin. International Journal of Pharmaceutics, 2022, 619, 121691.	5.2	10
3	Preparation and In Vivo Evaluation of Rosmarinic Acid-Loaded Transethosomes After Percutaneous Application on a Psoriasis Animal Model. AAPS PharmSciTech, 2021, 22, 103.	3.3	18
4	Cholesterol Levels Affect the Performance of AuNPs-Decorated Thermo-Sensitive Liposomes as Nanocarriers for Controlled Doxorubicin Delivery. Pharmaceutics, 2021, 13, 973.	4.5	7
5	Synergistic Effect of Acetazolamide-(2-hydroxy)propyl Î ² -Cyclodextrin in Timolol Liposomes for Decreasing and Prolonging Intraocular Pressure Levels. Pharmaceutics, 2021, 13, 2010.	4.5	1
6	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
7	Curcumin-in-Cyclodextrins-in-Liposomes: An Alternative for Osteoarthritis Treatment. Proceedings (mdpi), 2020, 78, .	0.2	1
8	Didodecyldimethylammonium Bromide Role in Anchoring Gold Nanoparticles onto Liposome Surface for Triggering the Drug Release. AAPS PharmSciTech, 2019, 20, 294.	3.3	6
9	Electrochemical characterization of a mixed lipid monolayer supported on Au(111) electrodes with implications for doxorubicin delivery. Journal of Electroanalytical Chemistry, 2018, 815, 246-254.	3.8	10
10	Ophthalmic administration of a 10-fold-lower dose of conventional nanoliposome formulations caused levels of intraocular pressure similar to those induced by marketed eye drops. European Journal of Pharmaceutical Sciences, 2018, 111, 186-194.	4.0	16
11	Fucoxanthin-Containing Cream Prevents Epidermal Hyperplasia and UVB-Induced Skin Erythema in Mice. Marine Drugs, 2018, 16, 378.	4.6	62
12	Topical Application of Glycolipids from Isochrysis galbana Prevents Epidermal Hyperplasia in Mice. Marine Drugs, 2018, 16, 2.	4.6	22
13	A comparative study of stabilising effect and antioxidant activity of different antioxidants on levodopa-loaded liposomes. Journal of Microencapsulation, 2018, 35, 357-371.	2.8	19
14	Deformability properties of timolol-loaded transfersomes based on the extrusion mechanism. Statistical optimization of the process. Drug Development and Industrial Pharmacy, 2016, 42, 1683-1694.	2.0	41
15	Specific requirements regarding module 5. Pharmaceuticals Policy and Law, 2015, 17, 279-281.	0.1	O
16	Specific requirements for somatic cell therapy medicinal products and tissue engineered products. Pharmaceuticals Policy and Law, 2015, 17, 271-277.	0.1	0
17	Surface functionalizing of a lipid nanosystem to promote brain targeting: step-by-step design and physico-chemical characterization. Pharmaceutical Development and Technology, 2015, 21, 1-9.	2.4	2
18	Método Avenzoarâ€,para la implantación racional de la atención farmacéutica en la farmacia comunitaria. FarmacÉuticos Comunitarios, 2015, 7, 37-44.	0.0	0

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19	Development and validation of a high performance chromatographic method for determining sumatriptan in niosomes. Journal of Pharmaceutical and Biomedical Analysis, 2013, 72, 251-260.	2.8	17
20	Thermal and 31P-NMR studies to elucidate sumatriptan succinate entrapment behavior in Phosphatidylcholine/Cholesterol liposomes. Comparative 31P-NMR analysis on negatively and positively-charged liposomes. Colloids and Surfaces B: Biointerfaces, 2013, 105, 14-23.	5. 0	26
21	Robust Optimization of Alginate-Carbopol 940 Bead Formulations. Scientific World Journal, The, 2012, 2012, 1-15.	2.1	7
22	Applying the taguchi method to optimize sumatriptan succinate niosomes as drug carriers for skin delivery. Journal of Pharmaceutical Sciences, 2012, 101, 3845-3863.	3.3	20
23	Charged liposomes as carriers to enhance the permeation through the skin. Expert Opinion on Drug Delivery, 2011, 8, 857-871.	5. O	73
24	Diclofenac Salts, Part 6: Release from Lipid Microspheres. Journal of Pharmaceutical Sciences, 2011, 100, 3482-3494.	3.3	11
25	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
26	Bimodal Release of Olanzapine from Lipid Microspheres. Journal of Pharmaceutical Sciences, 2010, 99, 4251-4260.	3.3	12
27	Positively and negatively charged liposomes as carriers for transdermal delivery of sumatriptan: in vitro characterization. Drug Development and Industrial Pharmacy, 2010, 36, 666-675.	2.0	44
28	Effect of preparation technique on the properties and in vivo in vivo in efficacy of benzocaine-loaded ethosomes. Journal of Liposome Research, 2009, 19, 253-260.	3.3	68
29	Development and validation of a reverse-phase liquid chromatographic method for the assay of lidocaine hydrochloride in alginate-Gantrez® microspheres. Journal of Pharmaceutical and Biomedical Analysis, 2008, 47, 501-507.	2.8	19
30	Development, characterization and in vivo evaluation of benzocaine-loaded liposomes. European Journal of Pharmaceutics and Biopharmaceutics, 2007, 67, 86-95.	4.3	137
31	Application of statistical experimental design to study the formulation variables influencing the coating process of lidocaine liposomes. International Journal of Pharmaceutics, 2007, 337, 336-345.	5.2	84
32	Effect of cholesterol and ethanol on dermal delivery from DPPC liposomes. International Journal of Pharmaceutics, 2005, 298, 1-12.	5.2	273
33	Release of indomethacin from ultrasound dry granules containing lactose-based excipients. Journal of Controlled Release, 2005, 102, 39-47.	9.9	12
34	Diclofenac salts, II. Solid dispersions in PEG6000 and Gelucire 50/13. European Journal of Pharmaceutics and Biopharmaceutics, 2005, 60, 99-111.	4.3	67
35	Modified Doxorubicin for Improved Encapsulation in PVA Polymeric Micelles. Drug Delivery, 2004, 12, 15-20.	5.7	19
36	Characterization of Ibuproxam Binary and Ternary Dispersions with Hydrophilic Carriers. Drug Development and Industrial Pharmacy, 2004, 30, 65-74.	2.0	44

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37	Development of Enteric-coated Pectin-based Matrix Tablets for Colonic Delivery of Theophylline. Journal of Drug Targeting, 2003, 11, 365-371.	4.4	54
38	Eudragit® RS-PM and Ethocel® 100 Premium: influence over the behavior of didanosine inert matrix system. Il Farmaco, 2002, 57, 649-656.	0.9	9
39	Alginate/chitosan particulate systems for sodium diclofenac release. International Journal of Pharmaceutics, 2002, 232, 225-234.	5.2	241
40	Didanosine extended-release matrix tablets: optimization of formulation variables using statistical experimental design. International Journal of Pharmaceutics, 2002, 237, 107-118.	5.2	69
41	Development of sustained release matrix tablets of didanosine containing methacrylic and ethylcellulose polymers. International Journal of Pharmaceutics, 2002, 234, 213-221.	5.2	47
42	Channeling Agent and Drug Release from a Central Core Matrix Tablet. Drug Development and Industrial Pharmacy, 2001, 27, 439-446.	2.0	17
43	Estimation of the percolation thresholds in dextromethorphan hydrobromide matrices. European Journal of Pharmaceutical Sciences, 2001, 12, 453-459.	4.0	28
44	Lipids in pharmaceutical and cosmetic preparations. Grasas Y Aceites, 2000, 51, .	0.9	51
45	Design of controlled release inert matrices of naltrexone hydrochloride based on percolation concepts. International Journal of Pharmaceutics, 1999, 181, 23-30.	5.2	32
46	Effect of the temperature on a hydrate diclofenac salt. International Journal of Pharmaceutics, 1999, 181, 95-106.	5.2	6
47	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
48	Evaluation of Eudragit® RS-PO and Ethocel® 100 Matrices for the Controlled Release of Lobenzarit Disodium. Drug Development and Industrial Pharmacy, 1999, 25, 229-233.	2.0	20
49	The role of the drug/excipient particle size ratio in the percolation model for tablets. Pharmaceutical Research, 1998, 15, 216-220.	3.5	54
50	Validation study of the conductometrical analysis. Application to the drug release studies from controlled release systems. Journal of Pharmaceutical and Biomedical Analysis, 1998, 18, 281-285.	2.8	10
51	Influence of the pH Value of the Dissolution Medium on the Release Profiles of a Morphine Polymeric Complex. Drug Development and Industrial Pharmacy, 1997, 23, 553-559.	2.0	3
52	Application of Percolation Theory to Characterize the Release Behavior of Carteolol Matrix Systems. Drug Development and Industrial Pharmacy, 1997, 23, 1-8.	2.0	10
53	Dissolution Behavior of Oxazepam in Presence of Cyclodextrins: Evaluation of Oxazepam-Dimeb Binary Systemxs. Drug Development and Industrial Pharmacy, 1997, 23, 379-385.	2.0	28
54	Influence of the Disintegrant on the Drug Percolation Threshold in Tablets. Drug Development and Industrial Pharmacy, 1997, 23, 665-669.	2.0	0

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55	Nuclear Magnetic Resonance Investigations of the Inclusion Complexation of Gliclazide with \hat{l}^2 -Cyclodextrin. Journal of Pharmaceutical Sciences, 1997, 86, 72-75.	3.3	31
56	Factors governing the dissolution of diclofenac salts. European Journal of Pharmaceutical Sciences, 1996, 4, 231-238.	4.0	22
57	Relationship between drug percolation threshold and particle size in matrix tablets. Pharmaceutical Research, 1996, 13, 387-390.	3.5	62
58	Thermal characterization of polyethylene glycols applied in the pharmaceutical technology using differential scanning calorimetry and hot stage microscopy. Journal of Thermal Analysis, 1996, 46, 291-304.	0.6	33
59	Dissolution properties and in vivo behaviour of triamterene in solid dispersions with polyethylene glycols. Pharmaceutica Acta Helvetiae, 1996, 71, 229-235.	1.2	28
60	Zero-order release periods in inert matrices. Influence of the distance to the percolation threshold. Pharmaceutica Acta Helvetiae, 1996, 71, 335-339.	1.2	17
61	Fractal Analysis of Sodium Cholate Particles. Journal of Pharmaceutical Sciences, 1996, 85, 971-975.	3.3	16
62	Using the Percolation Theory to Explain the Release Behavior from Inert Matrix Systems. Drug Development and Industrial Pharmacy, 1996, 22, 201-210.	2.0	3
63	Communications Simultaneous Hplc Determination of some Drugs Commonly Used in Cold Medications: Dextromethorphan, Dephenhydramine, Phenylephrine, Phenylpropanolamine and Pseudoephedrine. Drug Development and Industrial Pharmacy, 1995, 21, 605-613.	2.0	29
64	Morphine Polymeric Coprecipitates for Controlled Release: Elaboration and Characterization. Drug Development and Industrial Pharmacy, 1994, 20, 2409-2424.	2.0	16
65	The Application of Solid Dispersion Technique with D-mannitol to the Improvement in Oral Absorption of Triamterene. Journal of Drug Targeting, 1994, 2, 45-51.	4.4	24
66	Dissolution Rate Study of Fresh and Aging Triamterene-Urea Solid Dispersions. Drug Development and Industrial Pharmacy, 1994, 20, 2729-2740.	2.0	8
67	Thermal analysis of the system triamterene-d-mannitol. Journal of Thermal Analysis, 1994, 42, 143-158.	0.6	6
68	A Rapid HPLC Method for the Quantification of Tyrothricin, Menthol, and Benzocaine in Pharmaceutical Formulations. Journal of Pharmaceutical Sciences, 1994, 83, 1147-1149.	3.3	12
69	Improvement of the diuretic effect of triamterene via solid dispersion technique with PEG 4000. European Journal of Drug Metabolism and Pharmacokinetics, 1994, 19, 295-302.	1.6	8
70	Thermal study of the polyethyleneglycol 6000-triamterene system. Journal of Thermal Analysis, 1993, 40, 453-462.	0.6	8
71	Formulation Factors Affecting Thimerosal Stability. Drug Development and Industrial Pharmacy, 1993, 19, 1673-1691.	2.0	8
72	Blaboration and Technological Characterization of Inert Matrix Tables of Careolol Hydrochloride. Drug Development and Industrial Pharmacy, 1992, 18, 911-918.	2.0	6

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73	Rheological Study of Lactose Coated with Acrylic Resins. Drug Development and Industrial Pharmacy, 1990, 16, 295-313.	2.0	9
74	Elaboration and Characterization of the Diazepam-Polyethyleneglycol 6000 Solid Dispersions. Drug Development and Industrial Pharmacy, 1990, 16, 2283-2301.	2.0	14