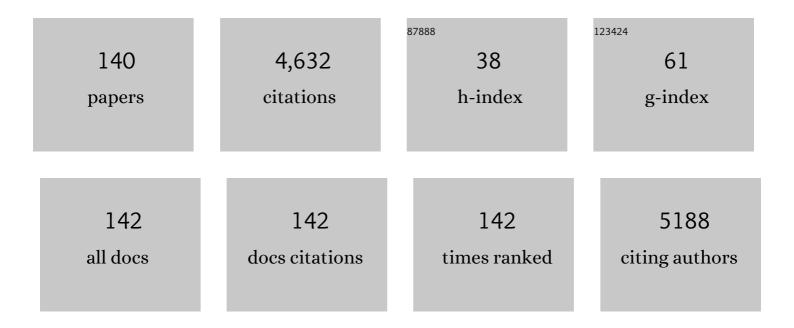
Rita Cortesi

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Cubosome Dispersions as Delivery Systems for Percutaneous Administration of Indomethacin. Pharmaceutical Research, 2005, 22, 2163-2173.	3.5	237
2	Production of lipospheres as carriers for bioactive compounds. Biomaterials, 2002, 23, 2283-2294.	11.4	179
3	Solid Lipid Nanoparticles as Delivery Systems for Bromocriptine. Pharmaceutical Research, 2008, 25, 1521-1530.	3.5	164
4	Gelatin microspheres: influence of preparation parameters and thermal treatment on chemico-physical and biopharmaceutical properties. Biomaterials, 1996, 17, 2009-2020.	11.4	152
5	Lipid-based supramolecular systems for topical application: A preformulatory study. AAPS PharmSci, 2003, 5, 62-76.	1.3	141
6	Sugar cross-linked gelatin for controlled release: microspheres and disks. Biomaterials, 1998, 19, 1641-1649.	11.4	133
7	Effect of cationic liposome composition on in vitro cytotoxicity and protective effect on carried DNA. International Journal of Pharmaceutics, 1996, 139, 69-78.	5.2	108
8	Nanoparticulate lipid dispersions for bromocriptine delivery: Characterization and in vivo study. European Journal of Pharmaceutics and Biopharmaceutics, 2012, 80, 306-314.	4.3	106
9	Formulation study for the antitumor drug camptothecin: liposomes, micellar solutions and a microemulsion. International Journal of Pharmaceutics, 1997, 159, 95-103.	5.2	105
10	Hyaluronan-based microspheres as tools for drug delivery: a comparative study. International Journal of Pharmaceutics, 2005, 288, 35-49.	5.2	97
11	Spray dried Eudragit microparticles as encapsulation devices for vitamin C. International Journal of Pharmaceutics, 2002, 242, 329-334.	5.2	90
12	Preparation of liposomes by reverse-phase evaporation using alternative organic solvents. Journal of Microencapsulation, 1999, 16, 251-256.	2.8	87
13	Production of Eudragit Microparticles by Spray-Drying Technique: Influence of Experimental Parameters on Morphological and Dimensional Characteristics. Pharmaceutical Development and Technology, 2000, 5, 267-278.	2.4	82
14	Effects of phospholipid based formulations on in vitro and in vivo percutaneous absorption of methyl nicotinate. Journal of Controlled Release, 1995, 34, 53-63.	9.9	74
15	In Vitro Antiproliferative Activity of Isothiocyanates and Nitriles Generated by Myrosinase-Mediated Hydrolysis of Glucosinolates from Seeds of Cruciferous Vegetables. Journal of Agricultural and Food Chemistry, 2000, 48, 3572-3575.	5.2	71
16	In VitroCytotoxic Activity of Some Glucosinolate-Derived Products Generated by Myrosinase Hydrolysis. Journal of Agricultural and Food Chemistry, 1996, 44, 1014-1021.	5.2	70
17	Clotrimazole-loaded nanostructured lipid carrier hydrogels: Thermal analysis and in vitro studies. International Journal of Pharmaceutics, 2013, 454, 695-702.	5.2	70
18	Preparation and characterisation of poly(vinyl alcohol)/cyclodextrin microspheres as matrix for inclusion and separation of drugs. International Journal of Pharmaceutics, 2004, 285, 87-96.	5.2	68

#	Article	IF	CITATIONS
19	Dextran cross-linked gelatin microspheres as a drug delivery system. European Journal of Pharmaceutics and Biopharmaceutics, 1999, 47, 153-160.	4.3	66
20	Myrosinase-generated isothiocyanate from glucosinolates: Isolation, characterization and in vitro antiproliferative studies. Bioorganic and Medicinal Chemistry, 1997, 5, 1799-1806.	3.0	65
21	Preparation and characterization of starch/cyclodextrin bioadhesive microspheres as platform for nasal administration of Gabexate Mesylate (Foy®) in allergic rhinitis treatment. Biomaterials, 2004, 25, 159-170.	11.4	62
22	Cationic liposomes as potential carriers for ocular administration of peptides with anti-herpetic activity. International Journal of Pharmaceutics, 2006, 317, 90-100.	5.2	60
23	Clotrimazole nanoparticle gel for mucosal administration. Materials Science and Engineering C, 2013, 33, 411-418.	7.3	58
24	Cellulose acetate butyrate microcapsules containing dextran ion-exchange resins as self-propelled drug release system. Biomaterials, 2005, 26, 4337-4347.	11.4	57
25	Nanostructured lipid systems modified with waste material of propolis for wound healing: Design, in vitro and in vivo evaluation. Colloids and Surfaces B: Biointerfaces, 2017, 158, 441-452.	5.0	57
26	Liposomes as carriers for DNA–PNA hybrids. Journal of Controlled Release, 2000, 68, 237-249.	9.9	56
27	Cationic liposomes as delivery systems for double-stranded PNA–DNA chimeras exhibiting decoy activity against NF-κB transcription factors. Biochemical Pharmacology, 2002, 64, 609-616.	4.4	54
28	Lipid-Based Nanosystems as a Tool to Overcome Skin Barrier. International Journal of Molecular Sciences, 2021, 22, 8319.	4.1	53
29	Nanostructured lipid dispersions for topical administration of crocin, a potent antioxidant from saffron (Crocus sativus L.). Materials Science and Engineering C, 2017, 71, 669-677.	7.3	49
30	Design and Characterization of Ethosomes for Transdermal Delivery of Caffeic Acid. Pharmaceutics, 2020, 12, 740.	4.5	46
31	Pullulan–cyclodextrin microspheres. Journal of Chromatography B: Analytical Technologies in the Biomedical and Life Sciences, 2003, 791, 407-419.	2.3	45
32	Challenges in the Physical Characterization of Lipid Nanoparticles. Pharmaceutics, 2021, 13, 549.	4.5	44
33	Ethosomes and Transethosomes for Mangiferin Transdermal Delivery. Antioxidants, 2021, 10, 768.	5.1	44
34	Cannabinoid antagonist in nanostructured lipid carriers (NLCs): design, characterization and in vivo study. Materials Science and Engineering C, 2015, 48, 328-336.	7.3	43
35	Curcumin containing monoolein aqueous dispersions: A preformulative study. Materials Science and Engineering C, 2013, 33, 4923-4934.	7.3	42
36	Evaluation of Monooleine Aqueous Dispersions as Tools for Topical Administration of Curcumin: Characterization, In Vitro and Ex-Vivo Studies. Journal of Pharmaceutical Sciences, 2013, 102, 2349-2361.	3.3	42

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37	Effect of nanostructured lipid vehicles on percutaneous absorption of curcumin. European Journal of Pharmaceutics and Biopharmaceutics, 2014, 86, 121-132.	4.3	41
38	Preparation and characterization of cationic microspheres for gene delivery. International Journal of Pharmaceutics, 1999, 189, 29-41.	5.2	40
39	Encapsulation of cannabinoid drugs in nanostructured lipid carriers. European Journal of Pharmaceutics and Biopharmaceutics, 2016, 102, 87-91.	4.3	39
40	Nanostructured lipid carriers (NLC) for the delivery of natural molecules with antimicrobial activity: production, characterisation and <i>in vitro</i> studies. Journal of Microencapsulation, 2017, 34, 63-72.	2.8	38
41	Preparation and Characterisation of Thermoresponsive		

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55	Gelatin microspheres as a new approach for the controlled delivery of synthetic oligonucleotides and PCR-generated DNA fragments. International Journal of Pharmaceutics, 1994, 105, 181-186.	5.2	27
56	Acyclovir delivery systems. Expert Opinion on Drug Delivery, 2008, 5, 1217-1230.	5.0	27
57	Non-phospholipid vesicles as carriers for peptides and proteins: Production, characterization and stability studies. International Journal of Pharmaceutics, 2007, 339, 52-60.	5.2	26
58	Liposomes- and ethosomes-associated distamycins: a comparative study. Journal of Liposome Research, 2010, 20, 277-285.	3.3	26
59	Ethosomes and organogels for cutaneous administration of crocin. Biomedical Microdevices, 2016, 18, 108.	2.8	26
60	The Potential of Caffeic Acid Lipid Nanoparticulate Systems for Skin Application: In Vitro Assays to Assess Delivery and Antioxidant Effect. Nanomaterials, 2021, 11, 171.	4.1	26
61	Liposomes, micelles and microemulsions as new delivery systems for cytotoxic alkaloids. Pharmaceutical Science & Technology Today, 1999, 2, 288-298.	0.7	25
62	Gallic acid loaded poloxamer gel as new adjuvant strategy for melanoma: A preliminary study. Colloids and Surfaces B: Biointerfaces, 2020, 185, 110613.	5.0	25
63	Macrophages loaded with doxorubicin by ATP-mediated permeabilization: Potential carriers for antitumor therapy. Biochimica Et Biophysica Acta - Molecular Cell Research, 1994, 1224, 269-276.	4.1	24
64	Nanosystems for skin hydration: a comparative study. International Journal of Cosmetic Science, 2007, 29, 39-47.	2.6	24
65	Pharmaceutical films made from the waste material from the preparation of propolis extracts: development and characterization. Brazilian Journal of Pharmaceutical Sciences, 2015, 51, 847-859.	1.2	24
66	Ethosomes and Transethosomes as Cutaneous Delivery Systems for Quercetin: A Preliminary Study on Melanoma Cells. Pharmaceutics, 2022, 14, 1038.	4.5	24
67	Pectinâ€Based Microspheres. Annals of the New York Academy of Sciences, 2001, 944, 160-179.	3.8	23
68	A novel multicompartimental system based on aminated poly(vinyl alcohol) microspheres/succinoylated pullulan microspheres for oral delivery of anionic drugs. International Journal of Pharmaceutics, 2007, 330, 129-137.	5.2	23
69	Thermal Magnetic Field Activated Propolis Release From Liquid Crystalline System Based on Magnetic Nanoparticles. AAPS PharmSciTech, 2018, 19, 3258-3271.	3.3	23
70	Nanoparticulate Gels for Cutaneous Administration of Caffeic Acid. Nanomaterials, 2020, 10, 961.	4.1	23
71	Lipid nanoparticles for administration of poorly water soluble neuroactive drugs. Biomedical Microdevices, 2017, 19, 44.	2.8	22
72	Poly[(N-isopropylacrylamide-co-acrylamide-co-(hydroxyethylmethacrylate))] thermoresponsive microspheres: An accurate method based on solute exclusion technique to determine the volume phase transition temperature. European Polymer Journal, 2007, 43, 3500-3509.	5.4	21

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73	Effect of new curcuminâ€containing nanostructured lipid dispersions on human keratinocytes proliferative responses. Experimental Dermatology, 2015, 24, 449-454.	2.9	21
74	Amphiphilic association systems for Amphotericin B delivery. International Journal of Pharmaceutics, 2003, 260, 249-260.	5.2	20
75	L-dopa co-drugs in nanostructured lipid carriers: A comparative study. Materials Science and Engineering C, 2017, 72, 168-176.	7.3	20
76	Monoolein liquid crystalline phases for topical delivery of crocetin. Colloids and Surfaces B: Biointerfaces, 2018, 171, 67-74.	5.0	20
77	Lipid Nanoparticles and Active Natural Compounds: A Perfect Combination for Pharmaceutical Applications. Current Medicinal Chemistry, 2019, 26, 4681-4696.	2.4	19
78	In vitro effect on human leukemic K562 cells of co-administration of liposome-associated retinoids and cytosine arabinoside (ara-C). , 1999, 62, 33-43.		18
79	Lipid nanostructures for antioxidant delivery: a comparative preformulation study. Beilstein Journal of Nanotechnology, 2019, 10, 1789-1801.	2.8	17
80	Design of propolis-loaded film forming systems for topical administration: The effect of acrylic acid derivative polymers. Journal of Molecular Liquids, 2021, 322, 114514.	4.9	17
81	Eudragit [®] microparticles for the release of budesonide: A comparative study. Indian Journal of Pharmaceutical Sciences, 2012, 74, 403.	1.0	17
82	Colloidal dispersions for the delivery of acyclovir: A comparative study. Indian Journal of Pharmaceutical Sciences, 2011, 73, 687.	1.0	16
83	Eudragit® microparticles as a possible tool for ophthalmic administration of acyclovir. Journal of Microencapsulation, 2007, 24, 445-456.	2.8	15
84	Monoolein aqueous dispersions as a delivery system for quercetin. Biomedical Microdevices, 2017, 19, 41.	2.8	15
85	Design of Nanosystems for the Delivery of Quorum Sensing Inhibitors: A Preliminary Study. Molecules, 2020, 25, 5655.	3.8	15
86	Design and characterization of fenretinide containing organogels. Materials Science and Engineering C, 2013, 33, 383-389.	7.3	14
87	A Correlative Imaging Study of in vivo and ex vivo Biodistribution of Solid Lipid Nanoparticles. International Journal of Nanomedicine, 2020, Volume 15, 1745-1758.	6.7	14
88	Liposome-associated retinoids: production, characterization and antiproliferative activity on neoplastic cells. European Journal of Pharmaceutical Sciences, 1994, 2, 281-291.	4.0	13
89	Production and characterization of biodegradable microparticles for the controlled delivery of proteinase inhibitors. International Journal of Pharmaceutics, 1996, 129, 263-273.	5.2	13
90	Cross-Enzyme Inhibition by Gabexate Mesylate: Formulation and Reactivity Study. Journal of Pharmaceutical Sciences, 1998, 87, 1335-1340.	3.3	13

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91	Formulations for natural and peptide nucleic acids based on cationic polymeric submicron particles. AAPS PharmSci, 2004, 6, 10-21.	1.3	13
92	Formulation and Bioequivalence of Two Valsartan Tablets After a Single Oral Administration. Scientia Pharmaceutica, 2011, 79, 123-135.	2.0	13
93	Cubic Phases, Cubosomes and Ethosomes for Cutaneous Application. Current Pharmaceutical Design, 2016, 22, 5382-5399.	1.9	13
94	Production and Characterization of Nanoparticle Based Hyaluronate Gel Containing Retinyl Palmitate for Wound Healing. Current Drug Delivery, 2018, 15, 1172-1182.	1.6	13
95	Influence of liposomal formulation parameters on the in vitro absorption of methyl nicotinate. International Journal of Pharmaceutics, 1998, 172, 255-260.	5.2	12
96	Natural antimicrobials in spray-dried microparticles based on cellulose derivatives as potential eco-compatible agrochemicals. Journal of Plant Diseases and Protection, 2017, 124, 269-278.	2.9	12
97	Controlled release of 1-β-D-arabinofuranosylcytosine (ara-C) from hydrophilic gelatin microspheres: in vitro studies. International Journal of Pharmaceutics, 1995, 117, 151-158.	5.2	11
98	Lipid nanocarriers containing a levodopa prodrug with potential antiparkinsonian activity. Materials Science and Engineering C, 2015, 48, 294-300.	7.3	11
99	Microparticles containing gallic and ellagic acids for the biological control of bacterial diseases of kiwifruit plants. Journal of Plant Diseases and Protection, 2017, 124, 563-575.	2.9	11
100	Production and Characterization of a Clotrimazole Liposphere Gel for Candidiasis Treatment. Polymers, 2018, 10, 160.	4.5	11
101	New Strategies for the Delivery of Some Natural Anti-oxidants with Therapeutic Properties. Mini-Reviews in Medicinal Chemistry, 2019, 19, 1030-1039.	2.4	11
102	Liposomes and Micellar Dispersions For Delivery of Benzoheterocyclic Derivatives of Distamycin A. Drug Delivery, 2007, 14, 1-8.	5.7	10
103	Effect of charge and lipid concentration on in-vivo percutaneous absorption of methyl nicotinate from liposomal vesicles. Journal of Pharmacy and Pharmacology, 2010, 57, 1169-1176.	2.4	10
104	Distamycins: Strategies for Possible Enhancement of Activity and Specificity. Mini-Reviews in Medicinal Chemistry, 2010, 10, 218-231.	2.4	10
105	Long-chain cationic derivatives of PTA (1,3,5-triaza-7-phosphaadamantane) as new components of potential non-viral vectors. International Journal of Pharmaceutics, 2012, 431, 176-182.	5.2	10
106	"Plurethosome―as Vesicular System for Cutaneous Administration of Mangiferin: Formulative Study and 3D Skin Tissue Evaluation. Pharmaceutics, 2021, 13, 1124.	4.5	10
107	Complexation to cationic microspheres of double-stranded peptide nucleic acid-DNA chimeras exhibiting decoy activity. Journal of Biomedical Science, 2004, 11, 697-704.	7.0	9
108	Waste Material of Propolis as a Film Forming Agent Intended to Modify the Metronidazole Release: Preparation and Characterization. Current Drug Delivery, 2016, 13, 1152-1164.	1.6	9

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109	DNA binding activity and inhibition of DNA-protein interactions. Biochemical Pharmacology, 1992, 44, 1985-1994.	4.4	8
110	Effect of DNA Complexation and Freeze-Drying on the Physicochemical Characteristics of Cationic Liposomes. Oligonucleotides, 2000, 10, 205-215.	4.3	8
111	Liposomes Containing Distamycins: Preparation, Characterization and Antiproliferative Activity. Drug Delivery, 2004, 11, 83-88.	5.7	8
112	Delivery systems for DNA-binding drugs as gene expression modulators. Drug Discovery Today, 2001, 6, 893-904.	6.4	7
113	Lipid-based nanoparticles containing cationic derivatives of PTA (1,3,5-triaza-7-phosphaadamantane) as innovative vehicle for Pt complexes: Production, characterization and in vitro studies. International Journal of Pharmaceutics, 2015, 492, 291-300.	5.2	7
114	Design of Liposomes Carrying HelixComplex Snail Mucus: Preliminary Studies. Molecules, 2021, 26, 4709.	3.8	7
115	Solid lipid nanoparticles for the delivery of 1,3,5-triaza-7-phosphaadamantane (PTA) platinum (II) carboxylates. Materials Science and Engineering C, 2017, 74, 357-364.	7.3	6
116	A spectrofluorometric analysis to evaluate transcutaneous biodistribution of fluorescent nanoparticulate gel formulations. European Journal of Histochemistry, 2022, 66, .	1.5	6
117	Mangiferin-Loaded Smart Gels for HSV-1 Treatment. Pharmaceutics, 2021, 13, 1323.	4.5	5
118	Manganese in Diagnostics: A Preformulatory Study. Pharmaceutics, 2022, 14, 108.	4.5	5
119	Production and antiproliferative activity of liposomes containing the antitumour drug chromomycin A ₃ . Journal of Microencapsulation, 1998, 15, 465-472.	2.8	4
120	Tailor-made core-shell nanospheres for antisense oligonucleotide delivery: IV.Adsorption/release behaviour. Journal of Biomaterials Science, Polymer Edition, 2001, 12, 1339-1357.	3.5	4
121	Analysis of the Drug Release Profiles from Formulations Based on Micro and Nano Systems. Current Analytical Chemistry, 2013, 9, 37-46.	1.2	4
122	Investigation of the Bioequivalence of Rosuvastatin 20 mg Tablets after a Single Oral Administration in Mediterranean Arabs Using a Validated LC-MS/MS Method. Scientia Pharmaceutica, 2016, 84, 536-546.	2.0	4
123	Nafion®-Containing Solid Lipid Nanoparticles as a Tool for Anticancer Pt Delivery: Preliminary Studies. Journal of Chemistry, 2017, 2017, 1-6.	1.9	4
124	Structural Studies of Lipid-Based Nanosystems for Drug Delivery: X-ray Diffraction (XRD) and Cryogenic Transmission Electron Microscopy (Cryo-TEM). , 2016, , 861-889.		4
125	Antisense Oligonucleotides Conjugated with Lipophilic Compounds: Synthesis and In Vitro Evaluation of Exon Skipping in Duchenne Muscular Dystrophy. International Journal of Molecular Sciences, 2022, 23, 4270.	4.1	4
126	Synthetic and Nanotechnological Approaches for a Diagnostic Use of Manganese. Molecules, 2022, 27, 3124.	3.8	4

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127	Gelified reverse micellar dispersions as percutaneous formulations. Journal of Drug Delivery Science and Technology, 2016, 32, 270-282.	3.0	3
128	Structural Studies of Lipid-Based Nanosystems for Drug Delivery: X-ray Diffraction (XRD) and Cryogenic Transmission Electron Microscopy (Cryo-TEM). , 2015, , 1-23.		3
129	Interchangeability of two 500 mg amoxicillin capsules with one 1000 mg amoxicillin tablet after a single oral administration. Indian Journal of Pharmaceutical Sciences, 2010, 72, 414.	1.0	3
130	Antioxidant-containing monoolein aqueous dispersions: a preliminary study. Drug Delivery and Translational Research, 2022, , 1.	5.8	3
131	Effect of long-term stabilization of cationic liposomes as defibrotide delivery system for antithrombotic activity. Drug Development Research, 2002, 55, 127-138.	2.9	2
132	Peptide-based cationic molecules for the production of positive charged liposomes and micelles. Journal of Microencapsulation, 2008, 25, 71-81.	2.8	2
133	Data on scaling up and in vivo human study of progesterone lipid nanoparticles. Data in Brief, 2017, 14, 639-642.	1.0	2
134	Production of Lipospheres for Bioactive Compound Delivery. , 2004, , 23-40.		2
135	Analysis of the Drug Release Profiles from Formulations Based on Micro and Nano Systems. Current Analytical Chemistry, 2012, 9, 37-46.	1.2	1
136	Monolein Aqueous Dispersions as a Tool to Increase Flavonoid Solubility: A Preliminary Study. Proceedings (mdpi), 2021, 78, 25.	0.2	1
137	Nanotechnological Strategies for Administration of Poorly Soluble Neuroactive Drugs. Proceedings (mdpi), 2020, 78, .	0.2	1
138	Polymeric microparticles for fenretinide administration. Macromolecular Symposia, 2014, 345, 14-23.	0.7	0
139	Cationic Lipospheres as Delivery Systems for Nucleic Acid Molecules. , 2004, , 143-159.		0
140	Complexation to Cationic Microspheres of Double-Stranded Peptide Nucleic Acid-DNA Chimeras Exhibiting Decoy Activity. Journal of Biomedical Science, 2004, 11, 697-704.	7.0	0