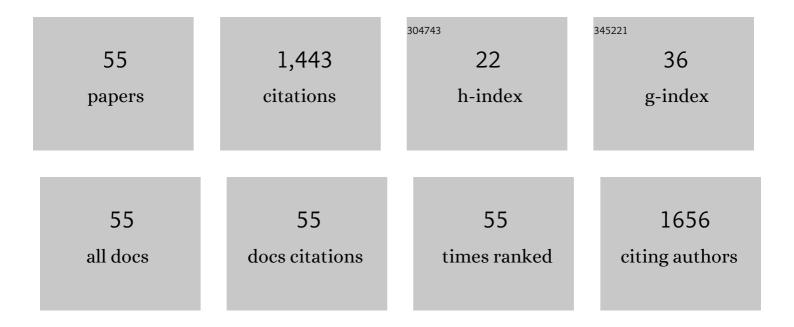
## Encarna Garcia\_Montoya

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Formulation of Direct Compression Zidovudine Tablets to Correlate the SeDeM Diagram Expert System and the Rotary Press Simulator Styl'ONE Results. AAPS PharmSciTech, 2020, 21, 1.	3.3	123
2	Coenzyme Q10 supplementation: Efficacy, safety, and formulation challenges. Comprehensive Reviews in Food Science and Food Safety, 2020, 19, 574-594.	11.7	94
3	A new expert systems (SeDeM Diagram) for control batch powder formulation and preformulation drug products. European Journal of Pharmaceutics and Biopharmaceutics, 2006, 64, 351-359.	4.3	80
4	Impact of physical parameters on particle size and reaction yield when using the ionic gelation method to obtain cationic polymeric chitosan–tripolyphosphate nanoparticles. International Journal of Pharmaceutics, 2013, 446, 199-204.	5.2	80
5	Quality by Design approach to understand the physicochemical phenomena involved in controlled release of captopril SR matrix tablets. International Journal of Pharmaceutics, 2014, 477, 431-441.	5.2	61
6	Trends in the food and sports nutrition industry: A review. Critical Reviews in Food Science and Nutrition, 2020, 60, 2405-2421.	10.3	60
7	Application of the SeDeM Diagram and a new mathematical equation in the design of direct compression tablet formulation. European Journal of Pharmaceutics and Biopharmaceutics, 2008, 69, 1029-1039.	4.3	57
8	The use of the SeDeM Diagram expert system to determine the suitability of diluents–disintegrants for direct compression and their use in formulation of ODT. European Journal of Pharmaceutics and Biopharmaceutics, 2009, 73, 414-423.	4.3	56
9	Predicting orally disintegrating tablets formulations of ibuprophen tablets: An application of the new SeDeM-ODT expert system. European Journal of Pharmaceutics and Biopharmaceutics, 2012, 80, 638-648.	4.3	55
10	Excipients in the Paediatric Population: A Review. Pharmaceutics, 2021, 13, 387.	4.5	51
11	An overview of microencapsulation in the food industry: opportunities, challenges, and innovations. European Food Research and Technology, 2020, 246, 1371-1382.	3.3	49
12	DNA delivery via cationic solid lipid nanoparticles (SLNs). European Journal of Pharmaceutical Sciences, 2013, 49, 157-165.	4.0	43
13	New classification of directly compressible (DC) excipients in function of the SeDeM Diagarm Expert System. International Journal of Pharmaceutics, 2014, 470, 15-27.	5.2	40
14	A new validated method for the simultaneous determination of benzocaine, propylparaben and benzyl alcohol in a bioadhesive gel by HPLC. Journal of Pharmaceutical and Biomedical Analysis, 2005, 39, 920-927.	2.8	38
15	Development and validation of a new HPLC analytical method for the determination of alprazolam in tablets. Journal of Pharmaceutical and Biomedical Analysis, 2004, 34, 979-987.	2.8	35
16	SeDeM expert system a new innovator tool to develop pharmaceutical forms. Drug Development and Industrial Pharmacy, 2014, 40, 222-236.	2.0	34
17	Relationships between surface free energy, surface texture parameters and controlled drug release in hydrophilic matrices. International Journal of Pharmaceutics, 2015, 478, 328-340.	5.2	34
18	Optimization of parameters of the SeDeM Diagram Expert System: Hausner index (IH) and relative humidity (%RH). European Journal of Pharmaceutics and Biopharmaceutics, 2011, 79, 464-472.	4.3	33

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19	A new optimized formulation of cationic solid lipid nanoparticles intended for gene delivery: Development, characterization and DNA binding efficiency of TCERG1 expression plasmid. International Journal of Pharmaceutics, 2014, 473, 270-279.	5.2	31
20	Chitosan nanoparticles as non-viral gene delivery systems: Determination of loading efficiency. Biomedicine and Pharmacotherapy, 2014, 68, 775-783.	5.6	31
21	The use of the SeDeM diagram expert system for the formulation of Captopril SR matrix tablets by direct compression. International Journal of Pharmaceutics, 2014, 461, 38-45.	5.2	29
22	Modification of the morphology and particle size of pharmaceutical excipients by spray drying technique. Powder Technology, 2015, 270, 244-255.	4.2	29
23	Development and validation of a new RP-HPLC method for the simultaneous determination of hydroquinone, kojic acid, octinoxate, avobenzone, BHA and BHT in skin-whitening cream. Analytical Methods, 2016, 8, 1170-1180.	2.7	21
24	Cholesteryl oleate-loaded cationic solid lipid nanoparticles as carriers for efficient gene-silencing therapy. International Journal of Nanomedicine, 2018, Volume 13, 3223-3233.	6.7	20
25	Comparison between Microcrystalline Celluloses of different grades made by four manufacturers using the SeDeM diagram expert system as a pharmaceutical characterization tool. Powder Technology, 2019, 342, 780-788.	4.2	20
26	Characterization of alginate beads loaded with ibuprofen lysine salt and optimization of the preparation method. International Journal of Pharmaceutics, 2014, 460, 181-188.	5.2	18
27	Formulation of Sustained Release Hydrophilic Matrix Tablets of Tolcapone with the Application of Sedem Diagram: Influence of Tolcapone's Particle Size on Sustained Release. Pharmaceutics, 2020, 12, 674.	4.5	17
28	Improved formulation of cationic solid lipid nanoparticles displays cellular uptake and biological activity of nucleic acids. International Journal of Pharmaceutics, 2017, 516, 39-44.	5.2	16
29	Application of an Experimental Design for the Optimization and Validation of a New HPLC Method for the Determination of Vancomycin in an Extemporaneous Ophthalmic Solution. Journal of Chromatographic Science, 2008, 46, 828-834.	1.4	15
30	Therapeutic potential of nicotinamide adenine dinucleotide (NAD). European Journal of Pharmacology, 2020, 879, 173158.	3.5	15
31	Stability evaluation of amoxicillin in a solid premix veterinary formulation by monitoring the degradation products through a new HPLC analytical method. Journal of Pharmaceutical and Biomedical Analysis, 2006, 42, 192-199.	2.8	14
32	Optimization of the Cohesion Index in the SeDeM Diagram Expert System and application of SeDeM Diagram: An improved methodology to determine the Cohesion Index. PLoS ONE, 2018, 13, e0203846.	2.5	12
33	Quality assurance in research: incorporating ISO9001:2000 into a GMP quality management system in a pharmaceutical R+D+I center. Accreditation and Quality Assurance, 2010, 15, 297-304.	0.8	11
34	Development and validation of a simple highâ€performance liquid chromatography analytical method for simultaneous determination of phytosterols, cholesterol and squalene in parenteral lipid emulsions. Biomedical Chromatography, 2018, 32, e4084.	1.7	11
35	SeDeM Diagram: an expert system for preformation, characterization and optimization of tablets obtained by direct compression. , 2013, , 109-135.		10
36	Preformulation and characterization of a lidocaine hydrochloride and dexamethasone sodium phosphate thermo-reversible and bioadhesive long-acting gel for intraperitoneal administration. International Journal of Pharmaceutics, 2016, 498, 142-152.	5.2	10

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37	The role of SeDeM for characterizing the active substance and polyvinyilpyrrolidone eliminating metastable forms in an oral lyophilizate—A preformulation study. PLoS ONE, 2018, 13, e0196049.	2.5	10
38	Optimization and Validation of a Fast UPLC Method for Simultaneous Determination of Hydroquinone, Kojic Acid, Octinoxate, Avobenzone, BHA, and BHT. Journal of AOAC INTERNATIONAL, 2017, 100, 1-7.	1.5	10
39	Formulation and characterization of mucoadhesive controlled release matrix tablets of captopril. Journal of Drug Delivery Science and Technology, 2017, 42, 215-226.	3.0	9
40	Osmolality predictive models of different polymers as tools in parenteral and ophthalmic formulation development. International Journal of Pharmaceutics, 2018, 543, 190-200.	5.2	7
41	Improved synthesis and characterization of cholesteryl oleate-loaded cationic solid lipid nanoparticles with high transfection efficiency for gene therapy applications. Colloids and Surfaces B: Biointerfaces, 2019, 180, 159-167.	5.0	7
42	New SeDeM-ODT expert system: an expert system for formulation of orodispersible tablets obtained by direct compression. , 2013, , 137-154.		6
43	Approach to design space from retrospective quality data. Pharmaceutical Development and Technology, 2016, 21, 26-38.	2.4	6
44	Study of Formulation Parameters by Factorial Design in Metoprolol Tartrate Matrix Systems. Drug Development and Industrial Pharmacy, 2001, 27, 965-973.	2.0	5
45	Development and Validation of a Stability Indicating RP-HPLC Method for Hydrocortisone Acetate Active Ingredient, Propyl Parahydroxybenzoate and Methyl Parahydroxybenzoate Preservatives, Butylhydroxyanisole Antioxidant, and Their Degradation Products in a Rectal Gel Formulation. Journal of AOAC INTERNATIONAL, 2015, 98, 27-34.	1.5	5
46	A new design for the review and appraisal of semi-solid dosage forms: Semi-solid Control Diagram (SSCD). PLoS ONE, 2018, 13, e0201643.	2.5	5
47	Spatiotemporal Analysis of Hydration Mechanism in Sodium Alginate Matrix Tablets. Materials, 2021, 14, 646.	2.9	5
48	Hydration Patterns in Sodium Alginate Polymeric Matrix Tablets—The Result of Drug Substance Incorporation. Materials, 2021, 14, 6531.	2.9	4
49	Improving tablet coating robustness by selecting critical process parameters from retrospective data. Pharmaceutical Development and Technology, 2015, 21, 1-10.	2.4	3
50	Method for the development of topical medicinal aerosols using liquified hydrocarbon gas. International Journal of Pharmaceutics, 2008, 355, 126-130.	5.2	2
51	Determination of stress-induced degradation products of cetirizine dihydrochloride by a stability-indicating RP-HPLC method. DARU, Journal of Pharmaceutical Sciences, 2014, 22, 82.	2.0	2
52	Development and Validation of a New High-Performance Liquid Chromatography Method for the Simultaneous Quantification of Coenzyme Q10, Phosphatidylserine, and Vitamin C from a Cutting-Edge Liposomal Vehiculization. ACS Omega, 2019, 4, 19710-19715.	3.5	2
53	Robustness Optimization of an Existing Tablet Coating Process Applying Retrospective Knowledge (rQbD) and Validation. Pharmaceutics, 2020, 12, 743.	4.5	2
54	Editing of a Job Description. Drug Development and Industrial Pharmacy, 1997, 23, 351-358.	2.0	0

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55	Quality and Integrity of Data in Research, Development, and Innovation: A Risk Analysis Method Applied to Laboratory Notebooks in a University Pilot Plant. PDA Journal of Pharmaceutical Science and Technology, 2011, 65, 207-216.	0.5	0