Pilar Pérez-Lozano

List of Publications by Year in descending order

Source: https://exaly.com/author-pdf/908774/publications.pdf

Version: 2024-02-01

516710 477307 35 863 16 citations h-index papers

g-index 36 36 36 934 docs citations times ranked citing authors all docs

29

#	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	Trends in the food and sports nutrition industry: A review. Critical Reviews in Food Science and Nutrition, 2020, 60, 2405-2421.	10.3	60
3	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
4	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
5	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
6	3D printed gummies: Personalized drug dosage in a safe and appealing way. International Journal of Pharmaceutics, 2020, 587, 119687.	5.2	51
7	Excipients in the Paediatric Population: A Review. Pharmaceutics, 2021, 13, 387.	4.5	51
8	DNA delivery via cationic solid lipid nanoparticles (SLNs). European Journal of Pharmaceutical Sciences, 2013, 49, 157-165.	4.0	43
9	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
10	SeDeM expert system a new innovator tool to develop pharmaceutical forms. Drug Development and Industrial Pharmacy, 2014, 40, 222-236.	2.0	34
11	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
12	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
13	Chitosan nanoparticles as non-viral gene delivery systems: Determination of loading efficiency. Biomedicine and Pharmacotherapy, 2014, 68, 775-783.	5 . 6	31
14	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
15	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
16	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
17	Bicyclic α-Iminophosphonates as High Affinity Imidazoline I ₂ Receptor Ligands for Alzheimer's Disease. Journal of Medicinal Chemistry, 2020, 63, 3610-3633.	6.4	17
18	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

#	Article	IF	CITATIONS
19	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
20	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
21	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
22	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
23	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
24	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
25	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
26	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
27	Approach to design space from retrospective quality data. Pharmaceutical Development and Technology, 2016, 21, 26-38.	2.4	6
28	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
29	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
30	Spatiotemporal Analysis of Hydration Mechanism in Sodium Alginate Matrix Tablets. Materials, 2021, 14, 646.	2.9	5
31	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
32	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
33	Robustness Optimization of an Existing Tablet Coating Process Applying Retrospective Knowledge (rQbD) and Validation. Pharmaceutics, 2020, 12, 743.	4.5	2
34	Application of a validated method in the stability study of colistin sulfate and methylparaben in a veterinary suspension formulation by high-performance liquid chromatography with a diode array detector. Journal of AOAC INTERNATIONAL, 2007, 90, 706-14.	1.5	2
35	(2-Imidazolin-4-yl)phosphonates: Green Chemistry and Biology Walk Together. Proceedings (mdpi), 2019, 22, 97.	0.2	0