Mario GonzÃ;lez-de la Parra

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Reference Datasets for Studies in a Replicate Design Intended for Average Bioequivalence with Expanding Limits. AAPS Journal, 2020, 22, 44.	2.2	0
2	Application of Design of Experiments (DOE) to the Development and Validation of a Swab Sampling Method for Cleaning Validation. Asian Journal of Chemistry and Pharmaceutical Sciences, 2017, 2, 16.	0.0	0
3	Application of Sequential Design of Experiments to Develop Ibuprofen (400 mg) Tablets by Direct Compression. Asian Journal of Chemistry and Pharmaceutical Sciences, 2017, 2, 10.	0.0	2
4	Quantification of 4-aminopyridine in plasma by capillary electrophoresis with electrokinetic injection. Journal of Chromatography B: Analytical Technologies in the Biomedical and Life Sciences, 2010, 878, 290-294.	1.2	9
5	Relative bioavailability of two oral formulations of piroxicam 20 mg: A single-dose, randomized-sequence, open-label, two-period crossover comparison in healthy Mexican adult volunteers. Clinical Therapeutics, 2010, 32, 357-364.	1.1	12
6	Bioavailability of two oral-tablet and two oral-suspension formulations of naproxen sodium/paracetamol (acetaminophen): Single-dose, randomized, open-label, two-period crossover comparisons in healthy Mexican adult subjects. Clinical Therapeutics, 2009, 31, 399-410.	1.1	5
7	Bioavailability of two oral formulations of a single dose of levofloxacin 500 mg: An open-label, randomized, two-period crossover comparison in healthy mexican volunteers. Clinical Therapeutics, 2009, 31, 1796-1803.	1.1	12
8	Bioavailability of Two Single-Dose Oral Formulations of Omeprazole 20 mg: An Open-Label, Randomized Sequence, Two-Period Crossover Comparison in Healthy Mexican Adult Volunteers. Clinical Therapeutics, 2008, 30, 693-699.	1.1	9
9	Bioavailability of two sublingual formulations of ketorolac tromethamine 30 mg: A randomized, open-label, single-dose, two-period crossover comparison in healthy mexican adult volunteers. Clinical Therapeutics, 2008, 30, 1667-1674.	1.1	15
10	Development and validation of a densitometric HPTLC method for quantitative analysis of levofloxacin in human plasma. Journal of Planar Chromatography - Modern TLC, 2008, 21, 209-212.	0.6	6
11	Development and Validation of a High-Performance Thin-Layer Chromatographic Method, with Densitometry, for Quantitative Analysis of Ketorolac Tromethamine in Human Plasma. Journal of AOAC INTERNATIONAL, 2008, 91, 1191-1195.	0.7	7
12	Bioequivalence of Two Commercial Preparations of trimethoprim/sulfamethoxazole: A randomized, single-dose, single-blind, crossover trial. Clinical Therapeutics, 2007, 29, 326-333.	1.1	6
13	Bioavailability of two oral suspension and two oral tablet formulations of acyclovir 400 mg: Two single-dose, open-label, randomized, two-period crossover comparisons in healthy Mexican adult subjects. Clinical Therapeutics, 2007, 29, 1146-1152.	1.1	8
14	Development and validation of a high-performance thin-layer chromatographic method, with densitometry, for quantitative analysis of tizoxanide (a Metabolite of Nitazoxanide) in human plasma. Journal of Planar Chromatography - Modern TLC, 2007, 20, 331-334.	0.6	4
15	Bioavailability of two oral formulations of loratadine 20 mg with concomitant ketoconazole: An open-label, randomized, two-period crossover comparison in healthy Mexican adult volunteers. Clinical Therapeutics, 2006, 28, 110-115.	1.1	6
16	Using Structural Equation Modeling (SEM) for the Study of Impurity Profiles of Drug Substances. Quality Engineering, 2006, 18, 225-235.	0.7	6
17	Bioavailability of two oral formulations of azithromycin 500 mg: A randomized, open-label, two-period crossover comparison in healthy Mexican adult subjects. Clinical Therapeutics, 2005, 27, 1607-1611.	1.1	13
18	Preliminary study on the synthesis and high‒resolution NMR analysis of Naproxen and Ibuprofen esters. Spectroscopy, 2004, 18, 485-500.	0.8	0

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19	Studying Unidentified Impurities in Drug Substances by the Application of Multivariate (Principal) Tj ETQq1 1 0.78	4314 rgB1 0.7	∫4Overloc≷
20	Application of Analysis of Means (ANOM) to Nested Designs for Improving the Visualization and Understanding of the Sources of Variation of Chemical and Pharmaceutical Processes. Quality Engineering, 2003, 15, 663-670.	0.7	7
21	Application of the Multivariate T2Control Chart and the Mason–Tracy–Young Decomposition Procedure to the Study of the Consistency of Impurity Profiles of Drug Substances. Quality Engineering, 2003, 16, 127-142.	0.7	14
22	The Use of a Germination Bioassay as a Toxicological Screening System to Study the Potential Drug–Drug Interactions of (S)-Naproxen and (S)-Ibuprofen and their corresponding Oxidation State Analogues, (S)-Naproxol and (S)-Ibuprofen Alcohol. ATLA Alternatives To Laboratory Animals, 1999, 27, 461-469.	0.7	0
23	Acaricidal potential of piquerols A and B againstBoophilus microplus. Pest Management Science, 1991, 33, 73-80.	0.6	10
24	Macrolide biosynthesis: Stereochemistry of the hydroxylation of brefelding C Journal of Antibiotics, 1987, 40, 1170-1174.	1.0	8