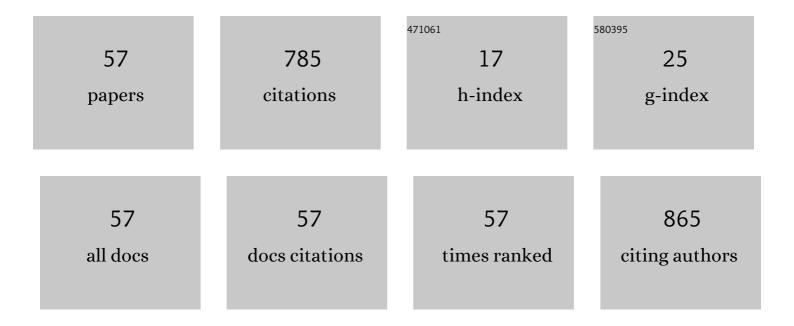
Anđelija M Malenović

List of Publications by Year in descending order

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#	Article	IF	CITATIONS
1	Properties of diclofenac sodium sorption onto natural zeolite modified with cetylpyridinium chloride. Colloids and Surfaces B: Biointerfaces, 2011, 83, 165-172.	2.5	105
2	Desirability-based optimization and its sensitivity analysis for the perindopril and its impurities analysis in a microemulsion LC system. Microchemical Journal, 2011, 99, 454-460.	2.3	53
3	Quantitation of pregabalin in dried blood spots and dried plasma spots by validated LC–MS/MS methods. Journal of Pharmaceutical and Biomedical Analysis, 2015, 109, 79-84.	1.4	34
4	Cationic surfactants-modified natural zeolites: improvement of the excipients functionality. Drug Development and Industrial Pharmacy, 2010, 36, 1215-1224.	0.9	29
5	Investigation of adsorption and release of diclofenac sodium by modified zeolites composites. Applied Clay Science, 2013, 83-84, 322-326.	2.6	29
6	Development of liquid chromatographic method for the analysis of dabigatran etexilate mesilate and its ten impurities supported by quality-by-design methodology. Journal of Pharmaceutical and Biomedical Analysis, 2015, 111, 7-13.	1.4	29
7	Microemulsion liquid chromatographic method for characterisation of fosinopril sodium and fosinoprilat separation with chemometrical support. Analytical and Bioanalytical Chemistry, 2005, 383, 687-694.	1.9	25
8	Chaotropic agents in liquid chromatographic method development for the simultaneous analysis of levodopa, carbidopa, entacapone and their impurities. Journal of Pharmaceutical and Biomedical Analysis, 2013, 77, 9-15.	1.4	23
9	Stepwise optimization approach for improving LCâ€MS/MS analysis of zwitterionic antiepileptic drugs with implementation of experimental design. Journal of Mass Spectrometry, 2013, 48, 875-884.	0.7	21
10	Chaotropic salts in liquid chromatographic method development for the determination of pramipexole and its impurities following quality-by-design principles. Journal of Pharmaceutical and Biomedical Analysis, 2015, 102, 314-320.	1.4	21
11	Effect of ibuprofen entrapment procedure on physicochemical and controlled drug release performances of chitosan/xanthan gum polyelectrolyte complexes. International Journal of Biological Macromolecules, 2021, 167, 547-558.	3.6	21
12	Monitoring of Impurity Level of Valsartan and Hydrochlorothiazide Employing an RP–HPLC Gradient Mode. Journal of Liquid Chromatography and Related Technologies, 2007, 30, 2879-2890.	0.5	20
13	Improved chromatographic response function in HILIC analysis: Application to mixture of antidepressants. Talanta, 2012, 98, 54-61.	2.9	20
14	Comparison of AQbD and grid point search methodology in the development of micellar HPLC method for the analysis of cilazapril and hydrochlorothiazide dosage form stability. Microchemical Journal, 2019, 145, 655-663.	2.3	20
15	Hematocrit effect on dried blood spots in adults: a computational study and theoretical considerations. Scandinavian Journal of Clinical and Laboratory Investigation, 2019, 79, 325-333.	0.6	19
16	Analytical quality by design development of an ecologically acceptable enantioselective HPLC method for timolol maleate enantiomeric purity testing on ovomucoid chiral stationary phase. Journal of Pharmaceutical and Biomedical Analysis, 2020, 180, 113034.	1.4	18
17	Optimization of Artificial Neural Networks for Modeling of Atorvastatin and Its Impurities Retention in Micellar Liquid Chromatography. Chromatographia, 2011, 73, 993-998.	0.7	17
18	Effects of derivatization reagents consisting of n-alkyl chloroformate/n-alcohol combinations in LC–ESI-MS/MS analysis of zwitterionic antiepileptic drugs. Talanta, 2013, 116, 91-99.	2.9	17

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19	Chemometrically assisted development and validation of LC–MS/MS method for the analysis of potential genotoxic impurities in meropenem active pharmaceutical ingredient. Journal of Pharmaceutical and Biomedical Analysis, 2017, 145, 307-314.	1.4	17
20	Analysis of potential genotoxic impurities in rabeprazole active pharmaceutical ingredient via Liquid Chromatography-tandem Mass Spectrometry, following quality-by-design principles for method development. Journal of Pharmaceutical and Biomedical Analysis, 2018, 149, 410-418.	1.4	17
21	Using a Combination of Experimental and Mathematical Method To Explore Critical Micelle Concentration of a Cationic Surfactant. Journal of Chemical Education, 2016, 93, 1277-1281.	1.1	14
22	FORCED DEGRADATION STUDIES OF SIMVASTATIN USING MICROEMULSION LIQUID CHROMATOGRAPHY. Journal of Liquid Chromatography and Related Technologies, 2010, 33, 536-547.	0.5	13
23	The influence of inorganic salts with chaotropic properties on the chromatographic behavior of ropinirole and its two impurities. Talanta, 2014, 123, 122-127.	2.9	13
24	Vigabatrin in dried plasma spots: Validation of a novel LC–MS/MS method and application to clinical practice. Journal of Chromatography B: Analytical Technologies in the Biomedical and Life Sciences, 2014, 962, 102-108.	1.2	12
25	Assessment of βâ€lactams retention in hydrophilic interaction chromatography applying <scp>B</scp> ox– <scp>B</scp> ehnken <scp>D</scp> esign. Journal of Separation Science, 2012, 35, 1424-1431.	1.3	11
26	Chemometrically assissted optimization and validation of RP-HPLC method for the analysis of itraconazole and its impurities. Acta Pharmaceutica, 2013, 63, 159-173.	0.9	11
27	CRITICAL REVIEW ON THE ANALYTICAL METHODS FOR THE DETERMINATION OF ZWITTERIONIC ANTIEPILEPTIC DRUGS—VIGABATRIN, PREGABALIN, AND GABAPENTIN—IN BULK AND FORMULATIONS. Instrumentation Science and Technology, 2014, 42, 486-512.	0.9	11
28	Investigation into the phenomena affecting the retention behavior of basic analytes in chaotropic chromatography: Joint effects of the most relevant chromatographic factors and analytes' molecular properties. Journal of Chromatography A, 2015, 1425, 150-157.	1.8	11
29	Chaotropic chromatography method development for the determination of aripiprazole and its impurities following analytical quality by design principles. Journal of Separation Science, 2020, 43, 3242-3250.	1.3	11
30	Validation of an Oil-in-Water Microemulsion Liquid Chromatography Method for Analysis of Perindopril tert-Butylamine and Its Impurities. Journal of AOAC INTERNATIONAL, 2011, 94, 723-734.	0.7	10
31	Quantitation of brinzolamide in dried blood spots by a novel LC-QTOF-MS/MS method. Journal of Pharmaceutical and Biomedical Analysis, 2016, 119, 84-90.	1.4	10
32	Design of Experiments–Design Space Approach for Development of Chaotropic Chromatography Method for Determination of Trimetazidine Dihydrochloride and Two Impurities. Chromatographia, 2017, 80, 585-592.	0.7	9
33	Modified aqueous mobile phases: A way to improve retention behavior of active pharmaceutical compounds and their impurities in liquid chromatography. Journal of Chromatography Open, 2022, 2, 100023.	0.8	9
34	Chitosan/Sodium Dodecyl Sulfate Complexes for Microencapsulation of Vitamin E and Its Release Profile—Understanding the Effect of Anionic Surfactant. Pharmaceuticals, 2022, 15, 54.	1.7	8
35	The influence of salt chaotropicity, column hydrophobicity and analytes' molecular properties on the retention of pramipexole and its impurities. Journal of Chromatography A, 2015, 1386, 39-46.	1.8	7
36	Robust Optimization of Chaotropic Chromatography Assay for Lamotrigine and its Two Impurities in Tablets. Chromatographia, 2019, 82, 565-577.	0.7	7

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37	INVESTIGATION OF TROPICAMIDE AND BENZALKONIUM CHLORIDE STABILITY USING LIQUID CHROMATOGRAPHY. Journal of Liquid Chromatography and Related Technologies, 2012, 35, 231-239.	0.5	6
38	Identification of the factors affecting the consistency of DBS formation via experimental design and image processing methodology. Microchemical Journal, 2019, 145, 1003-1010.	2.3	6
39	Quantitative structure retention relationship modeling as potential tool in chromatographic determination of stability constants and thermodynamic parameters of β-cyclodextrin complexation process. Journal of Chromatography A, 2020, 1619, 460971.	1.8	6
40	Five different columns in the analysis of basic drugs in hydrophilic interaction liquid chromatography. Open Chemistry, 2013, 11, 1150-1162.	1.0	5
41	OPTIMIZATION OF LIQUID CHROMATOGRAPHIC METHOD FOR THE SEPARATION OF FOLIC ACID AND ITS TWO IMPURITIES. Instrumentation Science and Technology, 2012, 40, 138-149.	0.9	4
42	Avoiding the False Negative Results in LC Method Robustness Testing by Modifications of the Algorithm of Dong and Dummy Factor Effects Approach. Chromatographia, 2012, 75, 397-401.	0.7	4
43	Testing the capability of a polynomialâ€modified gaussian model in the description and simulation of chromatographic peaks of amlodipine and its impurity in ionâ€interaction chromatography. Journal of Separation Science, 2014, 37, 1797-1804.	1.3	4
44	Simple and Efficient Solution for Robustness Testing in Gradient Elution Liquid Chromatographic Methods. Chromatographia, 2018, 81, 1135-1145.	0.7	4
45	Factorial Design in Optimization of Chromatographic Separation of Ramipril and Its Impurities. Chromatographia, 2010, 71, 799-804.	0.7	3
46	Physicochemical factors governing the partition of pramipexole and its five impurities in microemulsion liquid chromatographic systems. Journal of the Brazilian Chemical Society, 2012, , .	0.6	3
47	Evaluation of Seven Chromatographic Response Functions on Simulated and Experimentally Obtained Chromatograms in Hydrophilic Interaction Liquid Chromatography System. Analytical Letters, 2013, 46, 1198-1212.	1.0	3
48	Chemometrical Tools in the Study of the Retention Behavior of Azole Antifungals. Journal of Chromatographic Science, 2014, 52, 95-102.	0.7	3
49	Characterization of bonded stationary phase performance as a function of qualitative and quantitative chromatographic factors in chaotropic chromatography with risperidone and its impurities as model substances. Analytical and Bioanalytical Chemistry, 2018, 410, 4855-4866.	1.9	3
50	Influence of spray-drying process on properties of chitosan/xanthan gum polyelectrolyte complexes as carriers for oral delivery of ibuprofen. Arhiv Za Farmaciju, 2022, 72, 36-60.	0.2	3
51	Influence of the mobile phase and molecular structure parameters on the retention behavior of protonated basic solutes in chaotropic chromatography. Journal of Chromatography A, 2017, 1511, 68-76.	1.8	2
52	Evaluation of RP-HPLC Method Intended for the Analysis of Cefuroxime Axetil and ITS Impurities Supported by Experimental Design. Chromatographia, 2013, 76, 293-298.	0.7	1
53	Identification of the factors affecting the retention of weak acid solutes in hybrid micellar systems with cetyltrimethylammonium bromide. Journal of Liquid Chromatography and Related Technologies, 2019, 42, 45-53.	0.5	1
54	Corona Charged Aerosol Detector in studying retention and β-cyclodextrin complex stability using RP-HPLC. Journal of Pharmaceutical and Biomedical Analysis, 2021, 193, 113711.	1.4	1

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55	PDA-CAD method for the determination of magnesium, pyridoxine and thiamine in a dietary supplement supported by analytical quality by design methodology. Arhiv Za Farmaciju, 2021, 71, 378-392.	0.2	1
56	Generic Approach in a Gradient Elution HPLC Method Development that enables troubleshooting free method transfer. Journal of Pharmaceutical and Biomedical Analysis, 2021, 207, 114367.	1.4	0
57	Chaotropic effect of trifluoroacetic and perchloric acid on B-cyclodextrin inclusion complexation process with risperidone, olanzapine and their selected impurities. Arhiv Za Farmaciju, 2020, 70, 360-376.	0.2	ο