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List of Publications by Year
in descending order

131 papers	5,263 citations	81839 39 h-index	98753 67 g-index
131 all docs	131 docs citations	131 times ranked	3528 citing authors

#	Article	IF	Citations
1	Harmonization of strategies for the validation of quantitative analytical procedures. Journal of Pharmaceutical and Biomedical Analysis, 2007, 45, 70-81.	1.4	339
2	Harmonization of strategies for the validation of quantitative analytical proceduresA SFSTP proposalâ€"part I. Journal of Pharmaceutical and Biomedical Analysis, 2004, 36, 579-586.	1.4	331
3	Harmonization of strategies for the validation of quantitative analytical procedures. Journal of Pharmaceutical and Biomedical Analysis, 2007, 45, 82-96.	1.4	312
4	The SFSTP guide on the validation of chromatographic methods for drug bioanalysis: from the Washington Conference to the laboratory. Analytica Chimica Acta, 1999, 391, 135-148.	2.6	235
5	Enantiomeric separations of drugs using mixtures of charged and neutral cyclodextrins. Journal of Chromatography A, 2000, 875, 123-134.	1.8	170
6	An analysis of the SFSTP guide on validation of chromatographic bioanalytical methods: progresses and limitations. Journal of Pharmaceutical and Biomedical Analysis, 2003, 32, 753-765.	1.4	143
7	Harmonization of strategies for the validation of quantitative analytical procedures: A SFSTP proposal. Journal of Pharmaceutical and Biomedical Analysis, 2008, 48, 760-771.	1.4	124
8	Advances in validation, risk and uncertainty assessment of bioanalytical methods. Journal of Pharmaceutical and Biomedical Analysis, 2011, 55, 848-858.	1.4	121
9	Critical review of near-infrared spectroscopic methods validations in pharmaceutical applications. Journal of Pharmaceutical and Biomedical Analysis, 2012, 69, 125-132.	1.4	101
10	An improved HPLC-UV method for the simultaneous quantification of triterpenic glycosides and aglycones in leaves of Centella asiatica (L.) Urb (APIACEAE). Journal of Chromatography B: Analytical Technologies in the Biomedical and Life Sciences, 2009, 877, 2396-2402.	1.2	96
11	Data processing of vibrational chemical imaging for pharmaceutical applications. Journal of Pharmaceutical and Biomedical Analysis, 2014, 101, 123-140.	1.4	95
12	Enantiomeric purity determination of propranolol by cyclodextrin-modified capillary electrophoresis. Journal of Chromatography A, 1995, 717, 203-209.	1.8	91
13	Moisture content determination of pharmaceutical pellets by near infrared spectroscopy: Method development and validation. Analytica Chimica Acta, 2009, 642, 186-192.	2.6	88
14	Innovative high-performance liquid chromatography method development for the screening of 19 antimalarial drugs based on a generic approach, using design of experiments, independent component analysis and design space. Journal of Chromatography A, 2011, 1218, 5205-5215.	1.8	86
15	Improvement of the decision efficiency of the accuracy profile by means of a desirability function for analytical methods validation. Analytica Chimica Acta, 2007, 591, 239-247.	2.6	76
16	Critical review of surface-enhanced Raman spectroscopy applications in the pharmaceutical field. Journal of Pharmaceutical and Biomedical Analysis, 2018, 147, 458-472.	1.4	71
17	Sensitive determination of buprenorphine and its N-dealkylated metabolite norbuprenorphine in human plasma by liquid chromatography coupled to tandem mass spectrometry. Journal of Pharmaceutical and Biomedical Analysis, 2003, 32, 619-631.	1.4	69
18	Harmonization of strategies for the validation of quantitative analytical procedures. Journal of Pharmaceutical and Biomedical Analysis, 2004, 36, 579-586.	1.4	69

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19	Performances of a multidimensional on-line SPE-LC-ECD method for the determination of three major catecholamines in native human urine: Validation, risk and uncertainty assessments. Journal of Chromatography B: Analytical Technologies in the Biomedical and Life Sciences, 2006, 844, 251-260.	1.2	65
20	Acetaminophen determination in low-dose pharmaceutical syrup by NIR spectroscopy. Journal of Pharmaceutical and Biomedical Analysis, 2010, 53, 510-516.	1.4	65
21	Use of a novel cation-exchange restricted-access material for automated sample clean-up prior to the determination of basic drugs in plasma by liquid chromatography. Journal of Chromatography A, 2002, 975, 145-155.	1.8	64
22	Reliable low-cost capillary electrophoresis device for drug quality control and counterfeit medicines. Journal of Pharmaceutical and Biomedical Analysis, 2010, 53, 1278-1287.	1.4	64
23	Automated determination of tramadol enantiomers in human plasma using solid-phase extraction in combination with chiral liquid chromatography. Biomedical Applications, 1997, 698, 161-170.	1.7	63
24	Critical analysis of several analytical method validation strategies in the framework of the fit for purpose concept. Journal of Chromatography A, 2010, 1217, 3180-3192.	1.8	56
25	Active content determination of non-coated pharmaceutical pellets by near infrared spectroscopy: Method development, validation and reliability evaluation. Talanta, 2010, 80, 1750-1757.	2.9	55
26	Improvement of a stability-indicating method by Quality-by-Design versus Quality-by-Testing: A case of a learning process. Journal of Pharmaceutical and Biomedical Analysis, 2014, 88, 401-409.	1.4	54
27	Resolution improvement by use of carboxymethyl-Î²-cyclodextrin as chiral additive for the enantiomeric separation of basic drugs by capillary electrophoresis. Journal of Pharmaceutical and Biomedical Analysis, 1996, 14, 1107-1114.	1.4	52
28	Quality by Design Compliant Analytical Method Validation. Analytical Chemistry, 2012, 84, 106-112.	3.2	50
29	Integrated on-line sample clean-up using cation exchange restricted access sorbent for the LC determination of atropine in human plasma coupled to UV detection. Journal of Pharmaceutical and Biomedical Analysis, 2005, 36, 947-954.	1.4	49
30	LC method for the determination of R-timolol in S-timolol maleate: Validation of its ability to quantify and uncertainty assessment. Talanta, 2006, 68, 1166-1175.	2.9	49
31	Towards a full integration of optimization and validation phases: An analytical-quality-by-design approach. Journal of Chromatography A, 2015, 1395, 88-98.	1.8	49
32	Evaluation of a novel anion-exchange restricted-access sorbent for on-line sample clean-up prior to the determination of acidic compounds in plasma by liquid chromatography. Journal of Chromatography A, 2004, 1030, 95-102.	1.8	48
33	Comparing the qualitative performances of handheld NIR and Raman spectrophotometers for the detection of falsified pharmaceutical products. Talanta, 2019, 202, 469-478.	2.9	47
34	Determination of 4-aminophenol in a pharmaceutical formulation using surface enhanced Raman scattering: From development to method validation. Talanta, 2013, 116, 899-905.	2.9	46
35	Fully automated LC method for the determination of sotalol in human plasma using restricted access material with cation exchange properties for sample clean-up. Journal of Pharmaceutical and Biomedical Analysis, 2003, 32, 829-838.	1.4	45
36	Comparison of FT-NIR transmission and UVâ€“vis spectrophotometry to follow the mixing kinetics and to assay low-dose tablets containing riboflavin. Journal of Pharmaceutical and Biomedical Analysis, 2006, 41, 783-790.	1.4	45

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37	Quantitative analysis of N-acetylcysteine and its pharmacopeial impurities in a pharmaceutical formulation by liquid chromatographyâ€“UV detectionâ€“mass spectrometry. Journal of Chromatography A, 2000, 896, 191-199.	1.8	43
38	Determination of fenofibric acid in human plasma using automated solid-phase extraction coupled to liquid chromatography. Biomedical Applications, 2000, 742, 391-400.	1.7	43
39	A rapid validated UHPLCâ€“PDA method for anthocyanins quantification from Euterpe oleracea fruits. Journal of Chromatography B: Analytical Technologies in the Biomedical and Life Sciences, 2012, 907, 108-116.	1.2	41
40	Optimization and validation of a fast supercritical fluid chromatography method for the quantitative determination of vitamin D3 and its related impurities. Journal of Chromatography A, 2017, 1491, 171-181.	1.8	41
41	Using total error as decision criterion in analytical method transfer. Chemometrics and Intelligent Laboratory Systems, 2007, 85, 262-268.	1.8	39
42	Application of an innovative design space optimization strategy to the development of liquid chromatographic methods to combat potentially counterfeit nonsteroidal anti-inflammatory drugs. Journal of Chromatography A, 2012, 1263, 113-124.	1.8	39
43	Quantitative analysis of non-steroidal anti-inflammatory drugs by capillary zone electrophoresis. Journal of Pharmaceutical and Biomedical Analysis, 1995, 13, 497-503.	1.4	38
44	Development of response models for optimising HPLC methods. Chemometrics and Intelligent Laboratory Systems, 2004, 74, 263-268.	1.8	37
45	Validation of manufacturing process of Diltiazem HCl tablets by NIR spectrophotometry (NIRS). Journal of Pharmaceutical and Biomedical Analysis, 2007, 45, 356-361.	1.4	37
46	Development of a nano-liquid chromatography on chip tandem mass spectrometry method for high-sensitivity hepcidin quantitation. Journal of Chromatography A, 2011, 1218, 9046-9054.	1.8	36
47	Determination of albendazole and its main metabolites in ovine plasma by liquid chromatography with dialysis as an integrated sample preparation technique. Journal of Chromatography A, 2000, 870, 121-134.	1.8	33
48	Methodologies for the transfer of analytical methods: A review. Journal of Chromatography B: Analytical Technologies in the Biomedical and Life Sciences, 2009, 877, 2214-2223.	1.2	32
49	Validation of an automated method for the liquid chromatographic determination of atenolol in plasma: application of a new validation protocol. Analytica Chimica Acta, 1999, 391, 227-238.	2.6	30
50	Determination of uncertainty in analytical measurements from collaborative study results on the analysis of a phenoxymethylpenicillin sample. Analytica Chimica Acta, 2003, 481, 261-272.	2.6	30
51	Liquid chromatographic determination of enrofloxacin in nasal secretions and plasma of healthy pigs using restricted access material for on-line sample clean-up. Journal of Chromatography A, 2008, 1189, 456-466.	1.8	30
52	Near infrared and Raman spectroscopy as Process Analytical Technology tools for the manufacturing of silicone-based drug reservoirs. Analytica Chimica Acta, 2011, 699, 96-106.	2.6	30
53	Knowledge-based system for the automated solid-phase extraction of basic drugs from plasma coupled with their liquid chromatographic determination. Journal of Chromatography A, 1994, 665, 87-99.	1.8	29
54	Automated determination of verapamil and norverapamil in human plasma with on-line coupling of dialysis to high-performance liquid chromatography and fluorometric detection. Journal of Chromatography A, 1996, 750, 351-360.	1.8	29

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55	Total error and uncertainty: Friends or foes?. TrAC - Trends in Analytical Chemistry, 2011, 30, 797-806.	5.8	29
56	Fully automated method for the liquid chromatographicâ€“tandem mass spectrometric determination of cyproterone acetate in human plasma using restricted access material for on-line sample clean-up. Journal of Chromatography A, 2004, 1056, 105-110.	1.8	28
57	Automated determination of pirlindole enantiomers in plasma by on-line coupling of a pre-column packed with restricted access material to a chiral liquid chromatographic column. Journal of Pharmaceutical and Biomedical Analysis, 2002, 27, 447-455.	1.4	27
58	Development and validation of a fully automated LC method for the determination of cloxacillin in human plasma using anion exchange restricted access material for sample clean-up. Journal of Pharmaceutical and Biomedical Analysis, 2005, 36, 961-968.	1.4	27
59	Models to estimate overall analytical measurements uncertainty: Assumptions, comparisons and applications. Analytica Chimica Acta, 2011, 702, 160-171.	2.6	27
60	Development of an analytical method for crystalline content determination in amorphous solid dispersions produced by hot-melt extrusion using transmission Raman spectroscopy: A feasibility study. International Journal of Pharmaceutics, 2017, 530, 249-255.	2.6	27
61	Simultaneous determination of methylphenobarbital enantiomers and phenobarbital in human plasma by on-line coupling of an achiral precolumn to a chiral liquid chromatographic column. Journal of Chromatography A, 1998, 819, 143-153.	1.8	26
62	Determination of verapamil and norverapamil in human plasma by liquid chromatography: Comparison between a liquidâ€”liquid extraction procedure and an automated liquidâ€”solid extraction method for sample preparation. Journal of Pharmaceutical and Biomedical Analysis, 1992, 10, 937-942.	1.4	25
63	Validation of a liquid chromatographic-tandem mass spectrometric method for the determination of loperamide in human plasma. Journal of Chromatography B: Analytical Technologies in the Biomedical and Life Sciences, 2005, 814, 263-273.	1.2	25
64	Interlaboratory study of a liquid chromatography method for erythromycin: determination of uncertainty. Journal of Chromatography A, 2003, 1010, 63-74.	1.8	24
65	The transfer of a LC-UV method for the determination of fenofibrate and fenofibric acid in Lidoses: Use of total error as decision criterion. Journal of Pharmaceutical and Biomedical Analysis, 2006, 42, 64-70.	1.4	24
66	Quantitation of active pharmaceutical ingredient through the packaging using Raman handheld spectrophotometers: A comparison study. Talanta, 2020, 207, 120306.	2.9	24
67	Fully automated method for the liquid chromatographic determination of cyproterone acetate in plasma using restricted access material for sample pre-treatment. Journal of Chromatography B: Analytical Technologies in the Biomedical and Life Sciences, 2003, 795, 73-82.	1.2	23
68	Development of a quantitative approach using surface-enhanced Raman chemical imaging: First step for the determination of an impurity in a pharmaceutical model. Journal of Pharmaceutical and Biomedical Analysis, 2014, 90, 111-118.	1.4	23
69	Development, validation and comparison of NIR and Raman methods for the identification and assay of poor-quality oral quinine drops. Journal of Pharmaceutical and Biomedical Analysis, 2015, 111, 21-27.	1.4	23
70	Automatic Determination of Indomethacin in Human Plasma Using Liquid-Solid Extraction on Disposable Cartridges in Combination with HPLC. Journal of Liquid Chromatography and Related Technologies, 1990, 13, 3891-3907.	0.9	22
71	Robustness testing of a chiral NACE method for R-timolol determination in S-timolol maleate and uncertainty assessment from quantitative data. Journal of Pharmaceutical and Biomedical Analysis, 2007, 44, 640-651.	1.4	22
72	Development and validation of a sensitive solid phase extraction/hydrophilic interaction liquid chromatography/mass spectrometry method for the accurate determination of glucosamine in dog plasma. Journal of Chromatography A, 2010, 1217, 3275-3281.	1.8	21

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73	Enantioselective determination of oxprenolol in human plasma using dialysis coupled on-line to reversed-phase chiral liquid chromatography. Journal of Pharmaceutical and Biomedical Analysis, 1997, 15, 1365-1374.	1.4	19
74	Automatic determination of diltiazem and desacetyldiltiazem in human plasma using liquid-solid extraction on disposable cartridges coupled to HPLC â€" Part I: Optimization of the HPLC system and method validation. Journal of Pharmaceutical and Biomedical Analysis, 1991, 9, 877-882.	1.4	18
75	HPLC Determination of Verapamil and Norverapamil in Plasma Using Automated Solid Phase Extraction for Sample Preparation and Fluorometric Detection. Journal of Liquid Chromatography and Related Technologies, 1994, 17, 2147-2170.	0.9	18
76	Comparison of the quantitative performances and measurement uncertainty estimates obtained during method validation versus routine applications of a novel hydrophilic interaction chromatography method for the determination of cidofovir in human plasma. Journal of Pharmaceutical and Biomedical Analysis, 2012, 57, 153-165.	1.4	18
77	A simple approach for ultrasensitive detection of bisphenols by multiplexed surface-enhanced Raman scattering. Analytica Chimica Acta, 2015, 888, 118-125.	2.6	18
78	Collaborative study of an liquid chromatographic method for the determination of R-timolol and other related substances in S-timolol maleate. Analytica Chimica Acta, 2005, 546, 182-192.	2.6	17
79	Determination of binary polymorphic mixtures of fluconazole using near infrared spectroscopy and X-ray powder diffraction: A comparative study based on the pre-validation stage results. Journal of Pharmaceutical and Biomedical Analysis, 2011, 55, 1208-1212.	1.4	17
80	Automated liquid chromatographic determination of atenolol in plasma using dialysis and trace enrichment on a cation-exchange precolumn for sample handling. Biomedical Applications, 2000, 739, 205-217.	1.7	16
81	Uncertainty assessment from robustness testing applied on an LC assay for R-timolol and other related substances in S-timolol maleate. Analytica Chimica Acta, 2005, 531, 131-140.	2.6	16
82	Application of an innovative design space optimization strategy to the development of LC methods for the simultaneous screening of antibiotics to combat poor quality medicines. Journal of Pharmaceutical and Biomedical Analysis, 2013, 85, 83-92.	1.4	16
83	Vibrational spectroscopy and microspectroscopy analyzing qualitatively and quantitatively pharmaceutical hot melt extrudates. Journal of Pharmaceutical and Biomedical Analysis, 2015, 113, 21-33.	1.4	16
84	Global approach for the validation of an in-line Raman spectroscopic method to determine the API content in real-time during a hot-melt extrusion process. Talanta, 2017, 171, 45-52.	2.9	16
85	Risk-based approach for the transfer of quantitative methods: Bioanalytical applications. Journal of Chromatography A, 2008, 1189, 32-41.	1.8	15
86	Is supercritical fluid chromatography hyphenated to mass spectrometry suitable for the quality control of vitamin D3 oily formulations?. Journal of Chromatography A, 2017, 1515, 209-217.	1.8	15
87	Development of a SERS strategy to overcome the nanoparticle stabilisation effect in serum-containing samples: Application to the quantification of dopamine in the culture medium of PC-12 cells. Talanta, 2018, 186, 8-16.	2.9	15
88	Composition analysis of falsified chloroquine phosphate samples seized during the COVID-19 pandemic. Journal of Pharmaceutical and Biomedical Analysis, 2021, 194, 113761.	1.4	15
89	Determination of the enantiomers of 3-tert.-butylamino-1,2-propanediol by high-performance liquid chromatography coupled to evaporative light scattering detection. Journal of Chromatography A, 2000, 890, 239-249.	1.8	14
90	Fingerprinting and validation of a LC-DAD method for the analysis of biflavanones in Garcinia kola -based antimalarial improved traditional medicines. Journal of Pharmaceutical and Biomedical Analysis, 2016, 128, 382-390.	1.4	14

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91	A fully automated high-performance liquid chromatographic method for the determination of indomethacin in plasma. Journal of Pharmaceutical and Biomedical Analysis, 1989, 7, 1819-1827.	1.4	13
92	Determination of the enantiomers of 3-tert.-butylamino-1,2-propanediol by high-performance liquid chromatography using mass spectrometric detection. Journal of Chromatography A, 2000, 896, 201-207.	1.8	13
93	Application of a new optimization strategy for the separation of tertiary alkaloids extracted from Strychnos usambarensis leaves. Journal of Pharmaceutical and Biomedical Analysis, 2011, 56, 30-37.	1.4	13
94	Usefulness of capability indices in the framework of analytical methods validation. Analytica Chimica Acta, 2012, 714, 47-56.	2.6	13
95	Usefulness of Information Criteria for the Selection of Calibration Curves. Analytical Chemistry, 2013, 85, 6327-6335.	3.2	13
96	Automatic determination of diltiazem and desacetyldiltiazem in human plasma using liquidâ€" solid extraction on disposable cartridges coupled to HPLC â€" Part II: Optimization of liquidâ€"solid extraction. Journal of Pharmaceutical and Biomedical Analysis, 1991, 9, 883-887.	1.4	12
97	Enantiomeric separation of pirlindole by liquid chromatography using different types of chiral stationary phases. Journal of Pharmaceutical and Biomedical Analysis, 1998, 18, 605-614.	1.4	11
98	Optimisation and validation of a generic method for the LC assay of six corticosteroids and salicylic acid in dermopharmaceutical forms. Chromatographia, 2002, 55, 263-269.	0.7	11
99	Automated method for the determination of a new matrix metalloproteinase inhibitor in ovine plasma and serum by coupling of restricted access material for on-line sample clean-up to liquid chromatography. Journal of Chromatography B: Analytical Technologies in the Biomedical and Life Sciences, 2005, 817, 109-117.	1.2	11
100	VALIDATED QUANTITATIVE SIMULTANEOUS DETERMINATION OF COCAINE, OPIATES AND AMPHETAMINES IN SERUM BY U-HPLC COUPLED TO TANDEM MASS SPECTROMETRY. Acta Clinica Belgica, 2010, 65, 75-84.	0.5	11
101	Determination of l-lysine N-acetylcysteinate and its mono- and dimeric related compounds by liquid chromatographyâ€"mass spectrometry. Journal of Chromatography A, 1998, 819, 161-170.	1.8	10
102	Strategy for the development of automated methods involving dialysis and trace enrichment as on-line sample preparation for the determination of basic drugs in plasma by liquid chromatography. Journal of Chromatography A, 2002, 948, 151-161.	1.8	10
103	Direct determination of tagitinin C in Tithonia diversifolia leaves by on-line coupling of supercritical carbon dioxide extraction to FT-IR spectroscopy by means of optical fibres. Talanta, 2007, 71, 911-917.	2.9	10
104	Simultaneous determination of pirlindole enantiomers and dehydropirlindole by chiral liquid chromatography. Journal of Pharmaceutical and Biomedical Analysis, 1998, 17, 1071-1079.	1.4	9
105	Validation of analytical methods involved in dissolution assays: Acceptance limits and decision methodologies. Analytica Chimica Acta, 2012, 751, 44-51.	2.6	9
106	Application of NIR handheld transmission spectroscopy and chemometrics to assess the quality of locally produced antimalarial medicines in the Democratic Republic of Congo. Talanta Open, 2021, 3, 100025.	1.7	9
107	Fully automated determination of sulfamethazine in ovine plasma using solid-phase extraction on disposable cartridges and liquid chromatography. Biomedical Applications, 1993, 622, 53-60.	1.7	8
108	Enantioseparation of acidic drugs by capillary electrophoresis using dual systems with mixtures of charged and neutral cyclodextrins. , 1998, 12, 131-132.		8

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109	Liquid chromatographic analysis of local anesthetics in human plasma after sample preparation by on-line dialysis. Optimization by use of experimental design. Chromatographia, 2001, 53, 678-686.	0.7	8
110	Development and validation of a quantitative method for the selective determination of tin species in tin octoate by differential pulse polarography. Talanta, 2010, 80, 1413-1420.	2.9	8
111	Comments on â€œUncertainty profiles for the validation of analytical methodsâ€•by Saffaj and Ihssane. Talanta, 2012, 88, 769-771.	2.9	8
112	Methodology for the validation of analytical methods involved in uniformity of dosage units tests. Analytica Chimica Acta, 2013, 760, 46-52.	2.6	8
113	PLASMA LEVEL MONITORING OF THE MAJOR METABOLITES OF DIACETYLMORPHINE (HEROIN) BY THE â€œCHASING THE DRAGONâ€•ROUTE IN SEVERE HEROIN ADDICTS. Acta Clinica Belgica, 2013, 68, 359-367.	0.5	8
114	Evaluation of distributional homogeneity of pharmaceutical formulation using laser direct infrared imaging. International Journal of Pharmaceutics, 2022, 612, 121373.	2.6	8
115	Raman imaging as a new analytical tool for the quality control of the monitoring of osteogenic differentiation in forming 3D bone tissue. Journal of Pharmaceutical and Biomedical Analysis, 2020, 186, 113319.	1.4	7
116	Comparison of several strategies for the deployment of a multivariate regression model on several handheld NIR instruments. Application to the quality control of medicines. Journal of Pharmaceutical and Biomedical Analysis, 2022, 215, 114755.	1.4	7
117	Multivariate optimization approach for the separation of water-soluble vitamins and related compounds by capillary electrophoresis. , 2000, 14, 10-11.		6
118	Flexibility and applicability of Î²â€expectation tolerance interval approach to assess the fitness of purpose of pharmaceutical analytical methods. Drug Testing and Analysis, 2012, 4, 1014-1027.	1.6	6
119	In-line concentration and enantioseparation of clenbuterol by transient isotachophoresis-capillary zone electrophoresis-UV detection. , 2000, 14, 32-33.		5
120	Reply to the responses on the comments on â€œUncertainty profiles for the validation of analytical methodsâ€•by Saffaj and Ihssane. Talanta, 2012, 100, 290-292.	2.9	5
121	A simple calibration approach based on film-casting for confocal Raman microscopy to support the development of a hot-melt extrusion process. Talanta, 2016, 154, 392-399.	2.9	5
122	Do placebo based validation standards mimic real batch products behaviour? Case studies. Journal of Pharmaceutical and Biomedical Analysis, 2011, 55, 583-590.	1.4	4
123	Compromise in uncertainty estimation by modelling and validation approaches for an HPLC-UV method for measurement of biochemical indicators of vitamins A and E. Analyst, The, 2018, 143, 1492-1501.	1.7	4
124	Total error-based validation including the experimental design-based robustness evaluation of a stability-indicating method for the simultaneous quantification of hydrochlorothiazide and valsartan in tablet formulations. Acta Chromatographica, 2015, 27, 195-214.	0.7	3
125	HARMONISATION DES STRATÃ‰GIES DE VALIDATION ET ESTIMATION DE Lâ€™INCERTITUDE ASSOCIÃ‰E DANS LE CADRE DE Lâ€™ACCRÃ‰DITATION DES LABORATOIRES Dâ€™ESSAIS. Acta Clinica Belgica, 2006, 61, 52-54.	0.5	2
126	ESTIMATION OF UNCERTAINTY FROM THE TOTAL ERROR STRATEGY: APPLICATION TO INTERNAL AND NORMATIVE METHODS. Acta Clinica Belgica, 2010, 65, 100-104.	0.5	2

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127	Performance evaluation of a MIP for the MISPE-LC determination of p-[18F]MPPF and a potential metabolite in human plasma. Journal of Pharmaceutical and Biomedical Analysis, 2020, 180, 113015.	1.4	2
128	Fluorescence Detection of a Neurotoxic Astroglia-Released Factor using Gradient Liquid Chromatography after Precolumn Derivatization. , 1997, 11, 93-95.		1
129	LE TRANSFERT Dâ€™UNE MÃ‰THODE DE DOSAGE AUTOMATISÃ‰E DE LA NORADRÃ‰NALINE DANS Lâ€™URINE HUMAINE: UTILISATION DE Lâ€™ERREUR TOTALE COMME CRITÃˆRE DE DÃ‰CISION.. Acta Clinica Belgica, 2006, 61, 55-57.	0.5	1
130	Monitoring of anatabine release by methyl jasmonate elicited BY-2 cells using surface-enhanced Raman scattering. Talanta, 2016, 160, 754-760.	2.9	1
131	Process Analysisâ€”Overview. , 2018, , 396-396.		0