Green, eco-friendly bio-analytical techniques for pharmaceutical analysis.

Shaaban H*

Department of Pharmaceutical Chemistry, College of Clinical Pharmacy, University of Dammam, Dammam, Saudi Arabia

Accepted on September 22, 2017

Introduction

Analytical chemistry plays an important role in pharmaceutical research. It is involved in different stages of drug manufacturing such as analysis of active ingredients, separation of enantiomers or impurities in bulk drugs and pharmaceutical formulations [1]. Bio analytical methods play an important role in the quantitative evaluation of drugs and their metabolites in physiological matrices such as blood, plasma, serum, or urine. Greening the analytical methods is gaining high interest among researchers. Because of the monetary and ecological impact of using large amounts of organic solvents and waste disposal, the analytical community are directed to implement the principles of green analytical chemistry (GAC) in analytical laboratories and to substitute polluting analytical methodologies with green ones [2]. However, various techniques can be used for the bio analysis of pharmaceuticals, liquid chromatography is still the most popular technique used in commercial and research laboratories [3]. Chromatographic techniques have the potential to be greener at all steps of the analysis, from sample collection to separation. Different approaches could be used for making liquid chromatographic methods more eco-friendly such as using high speed liquid chromatography, superheated water chromatography, enhanced fluidity liquid chromatography, micellar liquid chromatography, miniaturized instruments, direct liquid chromatography and replacing toxic reagents with green alternatives [4].

Successful trials for greening bio-analytical methods are well documented in the literature. For example, green, eco-friendly UPLC methods were developed for the analysis of various classes of pharmaceuticals in biological fluids using columns packed with fully porous sub-2 μ m particles and superficially porous particles. The use of small particle size packing (sub 2 μ m) could enhance the chromatographic performance and reduce the analysis time [5]. The high column backpressure induced by these particles could be overcome by using these particles in combination with UPLC systems.

Several applications of using UPLC–UV, UPLC–MS and UPLC-MS/MS (Quadrople and time-of-flight mass analyzer) methods with columns packed with sub-2 μ m particles in bio analysis of pharmaceuticals are reported [6,7]. Another approach for greening bio analysis is the use of superficially porous particles (fused-core particles). These particles could reduce the analysis times while maintaining column efficiencies with relatively low back pressures. The use of fused-core particles for the analysis of pharmaceuticals in biological fluids is well documented in the literature [8-10].

Direct analysis is another approach for greening liquid chromatography and the used direct LC methods for the bioanalysis of pharmaceuticals is also reported [11-13]. For example, Strano-Rossi et al. developed a UHPLC–ESI-MS/MS method for the direct analysis of 14 drugs of abuse in oral fluids with no sample preparation [14]. Miniaturization is also one of the recent methods for greening bio analytical techniques and its applications in bio analysis is well documented [15,16]. In addition, two-dimensional liquid chromatographic separations are applied for the identification and quantification of drugs and their metabolites in biological fluids such as plasma or urine [17].

Greening bio analytical techniques and replacing the existed methods with green ones are expected in the coming years with the aim of minimizing the ecological impacts of the consumption of large amounts of solvents and waste disposal.

References

- 1. Gorog S. The changing face of pharmaceutical analysis. Trac Trend Anal Chem. 2007;26(1):7-12.
- 2. Shaaban H, Gorecki T. Current trends in green liquid chromatography for the analysis of pharmaceutically active compounds in the environmental water compartments. Talanta. 2015;132:739-52.
- 3. Nunez O, Gallart-Ayala H, Martins CPB. New trends in fast liquid chromatography for food and environmental analysis. J Chromatogr. 2012;1228:298-323.
- 4. Shaaban H. New insights into liquid chromatography for more eco-friendly analysis of pharmaceuticals. Anal Bioanal Chem. 2016;408(25):6929-44.
- 5. Shaaban H, Górecki T. Fused core particles as an alternative to fully porous sub-2 μm particles in pharmaceutical analysis using coupled columns at elevated temperature. Anal Methods. 2012;4:2735-43.
- 6. Keith LH, Gron LU, Young JL. Green analytical methodologies. Chem Rev. 2007;107(6):2695-708.
- 7. El-Bagary R, Azzazy HME, ElKady EF, et al. Simultaneous determination of sildenafil citrate and some nitric oxide releasing drugs in human plasma using UPLC MS/MS. Clin Biochem. 2014;47(7):654-6.
- 8. Tolgyesi A, Sharma VK, Fekete S, et al. Development of a rapid method for the determination and confirmation of nitroimidazoles in six matrices by fast liquid chromatography-tandem mass spectrometry. J Pharmaceut Biomed Anal. 2012;64: 40-8.

- Cunliffe JM, Shen JX, Wei XR, et al. Implementation of high-temperature superficially porous technologies for rapid LC-MS/MS diastereomer bioanalysis. Bioanalysis. 2011;3:735-43.
- Tolgyesi A, Sharma VK, Fekete J. Confirmatory analysis of stanozolol metabolites in bovine, pig and sheep urines using an optimized clean-up and liquid chromatography-tandem mass spectrometry. J Pharm Biomed Anal. 2014;88:45-52.
- 11. Yamamoto E, Sakaguchi T, Kajima T, et al. Novel methylcellulose-immobilized cation-exchange precolumn for online enrichment of cationic drugs in plasma. J Chromatogr. 2004;807(2):327-34.
- Santos-Neto AJ, Markides KE, Sjoberg PJR, et al. Capillary column switching restricted-access media-liquid chromatography-electrospray ionization-tandem mass spectrometry system for simultaneous and direct analysis of drugs in biofluids. Anal Chem. 2007;79(16):6359-67.
- 13. Capella-Peiro ME, Gil-Agusti M, Martinavarro-Dominguez A, et al. Determination in serum of some barbiturates using

*Correspondence to:

Shaaban H Department of Pharmaceutical Chemistry College of Clinical Pharmacy University of Dammam Dammam Saudi Arabia Tel: +966546262270; Fax: + 966133330290; E-mail: hsmohammed79@gmail.com micellar liquid chromatography with direct injection. Anal Biochem. 2002;309(2):261-8.

- 14. Strano-Rossi S, Anzillotti L, Castrignano E, et al. UHPLC-ESI-MS/MS method for direct analysis of drugs of abuse in oral fluid for DUID assessment. Anal Bioanal Chem. 2011;401(2):609-24.
- 15. Otsuki Y, Kotani A, Kusu F. Capillary liquid chromatography with UV detection using N, N-diethyldithiocarbamate for determining platinum-based antitumor drugs in plasma. Chem Pharm Bull. 2012;60(5):665-9.
- Liu J, Zhao Z, Teffera Y. Application of on-line nanoliquid chromatography/mass spectrometry in metabolite identification studies. Rapid Commun Mass Spectrom. 2012;26(3):320-6
- Komaba J, Masuda Y, Hashimoto Y, et al. Ultrasensitive determination of limaprost, a prostaglandin E-1 analogue, in human plasma using on-line twodimensional reversedphase liquid chromatography-tandem mass spectrometry. J Chromatogr B. 2012;852(1):590-7.