The Lewis acidity of zirconia-based supports for HPLC has historically presented problems in the analysis of analytes containing Lewis base moieties, such as carboxylates, particularly in LC/MS applications where volatile mobile phase additives are required. In this application note we demonstrate the utility of a new Lewis acid deactivated zirconia-based column, ZirChrom®-MS.

Introduction

Historically, the Lewis base carboxylic acid moiety on non-steroidal anti-inflammatory drugs required the use of a Lewis base mobile phase additive of a higher strength in the elutropic series (such as phosphate or fluoride) (1). While these types of additives work well in applications with UV/Vis detection, their use is almost entirely prohibited in LC/MS applications due to their relatively low volatility.

The deactivation of Lewis acid sites on the surface of the ZirChrom®-MS particle allows the chromatography of Lewis base analytes using mobile phase additives of the users choice including conventional LC/MS compatible buffers (such as acetate and formate) throughout the pH range of 1-10.

Experimental

Four non-steroidal anti-inflammatory drugs were separated at 35°C using a ZirChrom®-MS column. The separation conditions were as follows:

Column: ZirChrom®-MS, 50 mm x 4.6 mm i.d.  
(Part Number: MS01-0546)

Mobile Phase: Isocratic elution: 40/60 A/B  
A: acetonitrile  
B: 10mM ammonium acetate, pH 5.0

Temperature: 35 ºC
Flow Rate: 1.0 ml/min.
Injection Vol.: 5 µl
Pressure Drop: 68 bar
Detection: UV at 254 nm

Four non-steroidal anti-inflammatory drugs were separated using simple acetonitrile/water isocratic elution and a LC/MS friendly acetate buffer. The selectivity of all four compounds is excellent which allows for a very good separation using only a short 5 cm column.

References


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