

ANALYSIS OF IBUPROFEN AND PREDNISOLONE WITH THE QUATTRO PREMIER™ XE

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INTRODUCTION

As part of the pharmaceutical development process, it is a regulatory requirement to determine the toxicity and efficacy of a candidate compound prior to its release to market. These safety assessment and clinical trial studies require monitoring for compound exposure. The key criteria in these analyses are sensitivity, throughput and robustness. The accepted technology for these analyses is LC/MS/MS operating in MRM mode. Waters® ACQUITY UPLC™ System technology brings faster analyses with lower levels of detection. UPLC™ offers improved resolution over HPLC, rapid gradient analysis with optimized injection cycles. To complement, the Waters Micromass® Quattro Premier™ XE Mass Spectrometer has been specifically designed for unsurpassed productivity and sensitivity, applying enhanced electronics and a new detector to yield significantly enhanced performance.

This technical note compares the sensitivity of the Quattro Premier with that of the the Quattro Premier XE tandem quadrupole mass spectrometer in negative ion mode, using 2 test compounds: the non steroidal anti-inflammatory (NSAID) drug Ibuprofen, and the steroid Prednisolone.



Waters ACQUITY UPLC System with the Waters Micromass Quattro Premier XE Mass Spectrometer.

EXPERIMENTAL CONDITIONS

UPLC Conditions

LC System:	ACQUITY UPLC system		
Column:	ACQUITY UPLC BEH C ₁₈ 2.1 x 50 mm, 1.7 µm		
Flow Rate:	0.6 mL/min		
Mobile Phase:	A = 10 mM NH ₄ OAc in 20% MeOH, pH 5.0 B = 10 mM NH ₄ OAc in 80/20 MeOH/ACN		
Gradient:	Time	%A	Curve
	0.0	95	6
	0.8	5	6
	1.0	5	6
	3.0	55	1

MS Conditions

MS System:	Quattro Premier and Quattro Premier XE tandem quadrupole mass spectrometers
ESI Capillary Voltage:	3.0 kV
Source Temp:	130 °C
Desolvation Temp:	380 °C
Desolvation Gas Flow:	800 L/hr
Cone Gas Flow:	50 L/hr
Ionization Mode:	ESI ⁻
Dwell Time:	100 ms
MRM Transitions:	Prednisolone = 359.5→329.1 Ibuprofen = 205.2→161.2

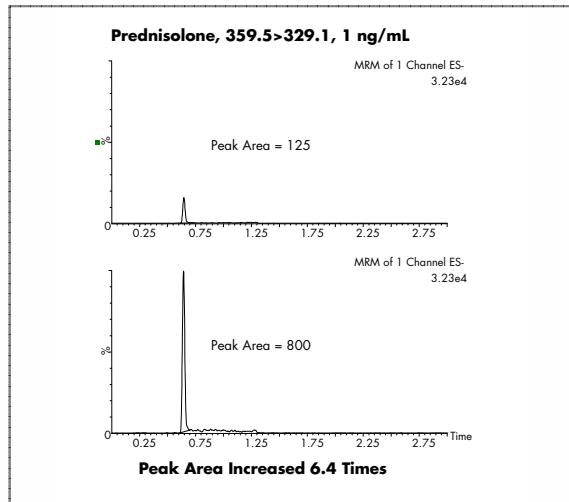


Figure 1. Comparative LC/MS/MS analysis for prednisolone on the Quattro Premier (top) and Quattro Premier XE (bottom) in negative electrospray.

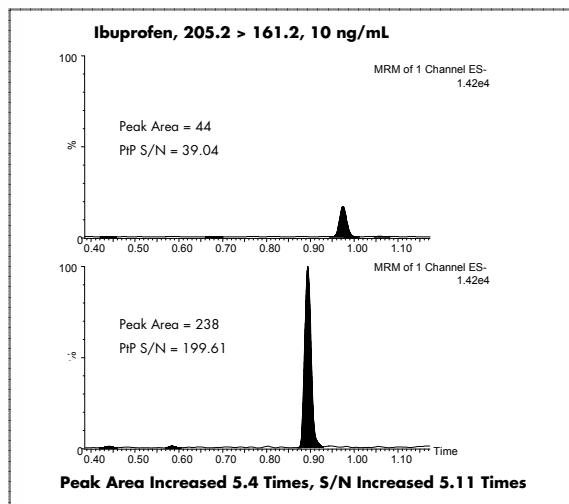


Figure 2. Comparison of the LC/MS/MS response for ibuprofen on the Quattro Premier (top) and Quattro Premier XE (bottom).

RESULTS AND DISCUSSION

The sensitivity of the Quattro Premier XE in negative ion mode was compared to that of the Quattro Premier. The data obtained from the analysis of the steroid prednisolone is displayed in Figure 1, using the MRM transition of 359.5→329.1. As demonstrated, with the same on-column loading and the same MRM parameters, a peak area increase of 6.4-fold was observed. In the second experiment, the NSAID compound ibuprofen was used as the test analyte. The results obtained are displayed in Figure 2. Here again we can see that with the same on-column loading and MS/MS parameters, a peak area increase of 5.4-fold was observed and the signal-to-noise ratio was increased by a factor of 5.

CONCLUSION

The Quattro Premier XE tandem mass spectrometer provides unmatched sensitivity in both positive and negative ionization modes. Additionally, the ACQUITY UPLC system produces the ultimate in chromatographic throughput and sensitivity, generating sharp peaks, 2 seconds at the base, with run times on the order of 1 minute. Combining these results in a premium analytical platform for increased data quality and productivity in quantitative bioanalysis.

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