# Waters™



# Ropinirole On Oasis MCX

Waters Corporation

## Introduction

Ropinirole is a potent, basic drug with a low oral dose range. A bioanalytical method needs to be developed that can reach a lower limit of quantitation of 0.02 ng/mL, which is lower than what is typically required. The Oasis® MCX µElution 96-well plate was selected for its increased sensitivity through concentration, without drying down which allows reconstituting the extract.

# Experimental

#### **Test Conditions**

Oasis MCX µ Elution 96-well plate:

Condition: 200 µL MeOH

Equilibrate: 200 µL H2O

Load: 400 µL diluted plasma sample (200 µL plasma

diluted 1:1 with 4% H3P O4)

Wash 1: 200 μL 2% HCOOH Wash 2: 200 µL MeOH Elute: 2 X 25 µL 5% NH4OH in 90:10 MeOH:H2O (less MeOH used for a cleaner extract - less residual matrix components eluted.) Injection: 8 μL (direct injection) Column: ACQUITY UPLC® BEH C<sub>18</sub>, 2.1 x 50 mm, 1.7 μm Mobile Phase A: 10 mM NH4COOH, pH 9 Mobile P hase B: MeOH Flow Rate: 0.5 mL/min Injection Volume: 8.0 µL Column Temp: 45 °C 15 °C Sample Temp: Sample Diluent: 25:75 H2O:MeOH Strong Needle Wash: 60:40 ACN:IPA + 0.5% HCOOH (1200 μL) Weak Needle Wash: 95:5 H2O:MeOH (500 μL)

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### **Gradient Table**

Time	Profile		Curvo
	%A	%B	Curve
0.0	95	5	6
2.0	2	98	6
2.5	2	98	6
2.6	95	5	6
3.0	95	5	6

Quattro Premier™

Desolvation Temp: 350 °C

Cone Gas Flow: 50 L/Hr

Desolvation Gas Flow: 700 L/Hr

Collision C ell P ressure: 2.6 x 10 (-3) mbar

MRM transitions monitored: Ropinirole m/z 261.2 > 113.95 (ESI+)

Citalopram (IS) m/z 325.2 > 108.85 (ESI+)

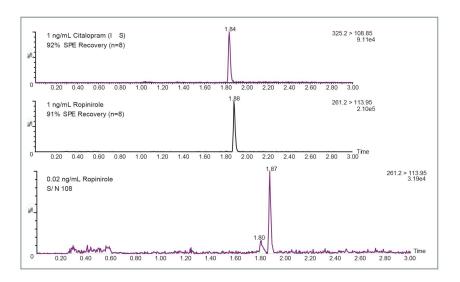
For both analytes, the optimal cone voltage was 40

V. For Ropinirole and Citalopram,

the collision energies were 18 eV and 25 eV,

respectively.

Ropinirole MW 260.4 pKa 9.7 Citalopram MW 324.4 pKa 9.5



## Results and Discussion

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