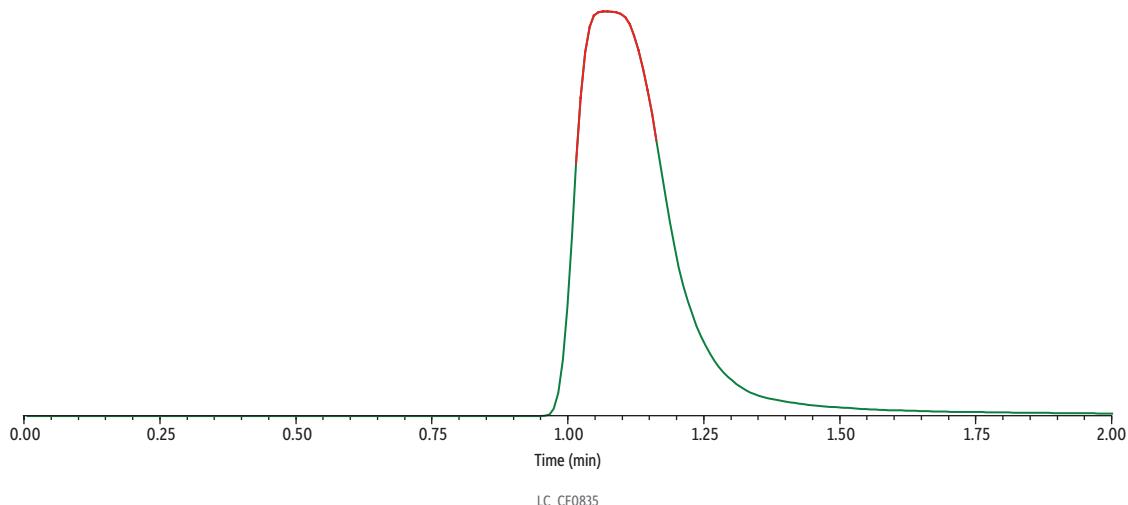


## 500 µg/mL of Gabapentin in Urine Analyzed on Raptor Biphenyl Using Method 1



Peaks	$t_r$ (min)	Precursor	Product 1	Product 2
1. Gabapentin	1.09	172.1	154.1	136.9

<b>Column</b>	Raptor Biphenyl (cat.# 9309A52)		
<b>Dimensions:</b>	50 mm x 2.1 mm ID		
<b>Particle Size:</b>	2.7 µm		
<b>Pore Size:</b>	90 Å		
<b>Guard Column:</b>	Raptor Biphenyl EXP guard column cartridge 5 mm, 2.1 mm ID, 2.7 µm (cat.# 9309A0252)		
<b>Temp.:</b>	45 °C		
<b>Standard/Sample</b>			
<b>Diluent:</b>	90:10 Water:methanol, both with 0.1% formic acid		
<b>Conc.:</b>	500 µg/mL		
<b>Inj. Vol.:</b>	5 µL		
<b>Mobile Phase</b>			
A:	Water, 0.1% formic acid		
B:	Methanol, 0.1% formic acid		
Time (min)	Flow (mL/min)	%A	%B
0.00	0.6	90	10
6.00	0.6	25	75
7.00	0.6	0	100
8.00	0.6	0	100
8.01	0.6	90	10
9.00	0.6	90	10
<b>Max Pressure:</b>	300 bar		
<b>Detector</b>	Shimadzu 8045 LC-MS/MS		
<b>Ion Mode:</b>	ESI+		
<b>Mode:</b>	MRM		
<b>Instrument</b>	Shimadzu Nexera X2		
<b>Sample Preparation</b>	Control urine (20 µL) was added to a 1.5 mL microcentrifuge tube along with 20 µL of a premade enzyme hydrolysis master mix. The sample was vortexed for 10 seconds and left to incubate at room temperature for 20 minutes. After the incubation, 260 µL of the diluent (water, 0.1% formic acid:methanol, 0.1% formic acid 90:10 [v/v]) was added. A 100 µL aliquot was added to a vial insert (cat.# 21776) in a 2.0 mL, amber, short-cap vial (cat.# 21142) and capped with a 9 mm short cap (cat.# 24497) and injected on the LC-MS/MS for analysis.		