



# Simultaneous detection and quantification of gastrin 17 and 34 sulfated and nonsulfated forms by LC-MS/MS in human plasma

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### Introduction:

Gastrin, secreted by G cells plays a crucial role in digestion and has diverse functions including regulation of the intestinal epithelium and stomach growth. Gastrin peptides are derived from progastrin. Peptides G17 and G34 are the most abundant in the blood. Both of them may be sulfated. Current gastrin measurement relies on the DIAsource RIA kit. However, it display cross-reactivity issues.

## • Materials and Methods:



ColumnBEH Peptide C18 130Å column<br/> $100 \times 2,1 \text{ mm}, 1.7 \mu \text{m}$  - WatersMobile<br/>phasesA:  $H_2O + 0,1\% \text{ NH}_4OH$ <br/>B: ACNSignal<br/>modeElectrospray – Negative

60 Gradient % 40 20





### Q1 Scan – MRM – Post column flow injection

	G34	G34S	G17	G17S
Transitions	768,8 > 716,0	784,9 > 732,4	523,4 > 610,8	434,5 > 478,0
	DP: -122,5	DP: -122,5	DP: -122,5	DP: -122,5
Compounds	EP: -7,0	EP: -10,0	EP: -8,0	EP: -10,0
parameters	CXP: -12,0	CXP: -12,0	CXP: -12,0	CXP: -12,0
	CE: -28,0	CE: -22,0	CE: -22,0	CE: -21,0
Source parameters	CUR: 20,0	IS: -3500	GS1: 40,0	
	CAD: Medium	TEM: 650	GS2: 50,0	

#### Validation steps

#### Change to ESI +

Colum lifetime issues >< high pH of the mobile phases</li>

• Derivatization of the carboxylic acids

43-62% Matrix effect

Recoveries

## • Conclusion:

- $\,\circ\,$  Choose the detection mode: ESI -/ ESI +
- Validation planned according CLSI guidelines





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