

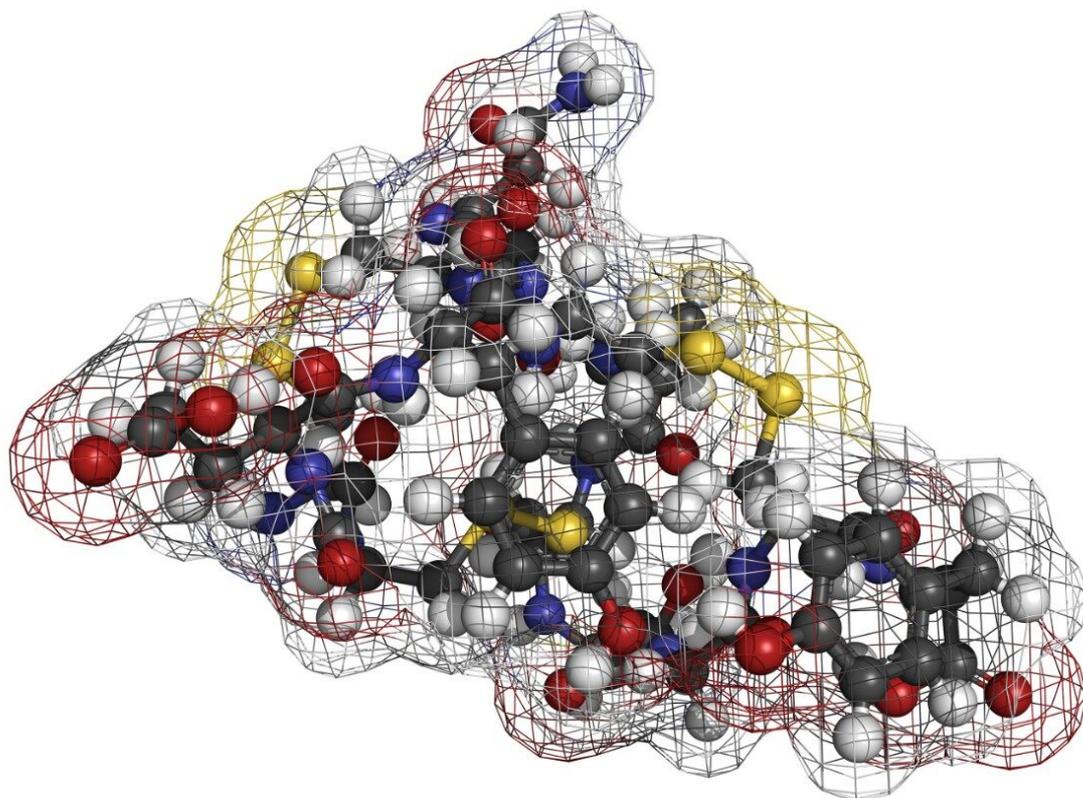
Nota applicativa

## UPLC-MS/MS Bioanalytical Quantification of Linaclotide from Plasma

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Bhaskar K, Raghu T., Taposh Gorella, Rajesh PMN, Dr. Padmakar Wagh

Waters Corporation



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Abstract

This application highlights the development of a highly sensitive and robust LC-MS assay for the quantification of linaclotide extracted from plasma.

The method described herein achieves an LLOQ of 10.0 pg/mL with a linear dynamic range of 10–4000 pg/mL. The high sensitivity and linearity of method was attributed to extraction specificity using mixed-mode  $\mu$ Elution SPE sample preparation, high-resolution UPLC chromatographic separation with the ACQUITY UPLC I-Class System using a sub-2-micron UPLC column, and sensitivity of the Xevo TQ-XS Mass Spectrometer. This developed method has demonstrated its fit-for-purpose use in support of drug discovery and research.

## Benefits

- Use of ACQUITY UPLC Columns for excellent resolution of linaclotide from endogenous interferences
- Fast, simple, and selective sample preparation using Oasis MAX SPE
- Linear, accurate, and precise quantitative performance
- High analytical sensitivity detection, achieving 10.0 pg/mL, using the Xevo TQ-XS equipped with ESI ion source

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## Introduction

Peptide therapeutics represents an emerging class of drugs with great pharmacologic potential because of their high specificity, affinity, and molecular target recognition. Linaclotide (LINZESS), shown in Figure 1, is a novel, potent 14 amino acid (CCEYCCNPACTGCV) synthetic peptide containing three disulfide bonds (1–6, 2–10, 5–13) with a molecular weight of 1526.80. It is a guanylate cyclase-C (GC-C) agonist used for treating irritable bowel syndrome with constipation and chronic idiopathic constipation.<sup>1,2,3</sup> Following oral dosing, linaclotide elicits its potent pharmacologic response locally in the gastrointestinal (GI) tract. It is minimally adsorbed with low systemic circulating plasma levels (<50 pg/mL).

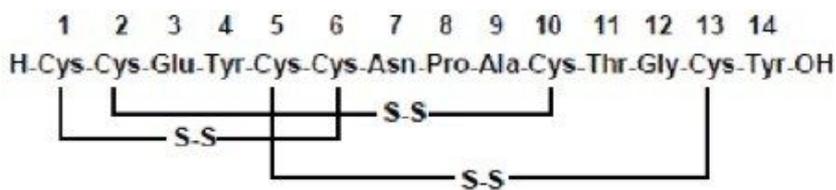


Figure 1. Amino acid sequence of linaclotide with disulfide bonds.

Versus traditional small molecule therapeutics, peptides are often more difficult to analyze by LC-MS/MS, as MS sensitivity is low due to poor transfer into the gas phase and poor fragmentation. In the case of linaclotide, it has 14 chiral centers and exhibits stereoisomerism. Additionally, due to the presence of disulphide bonds, it is a cyclic peptide (Figure 2). Thus, achieving MS sensitivity and selectivity adds additional challenges, as the protonation sites are less accessible, and fragmentation is limited due to the presence of disulphide bonds. With its cyclic nature and low circulating levels developing a sensitive and robust LC-MS analytical method for linaclotide is quite challenging.

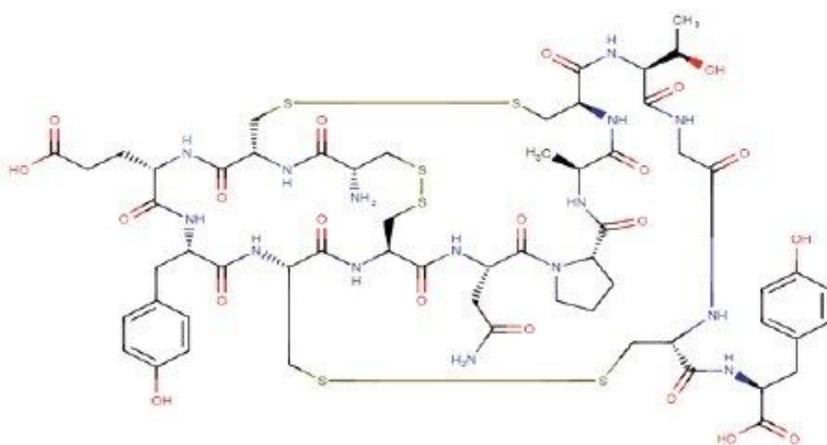


Figure 2. Cyclic structure of linaclotide.

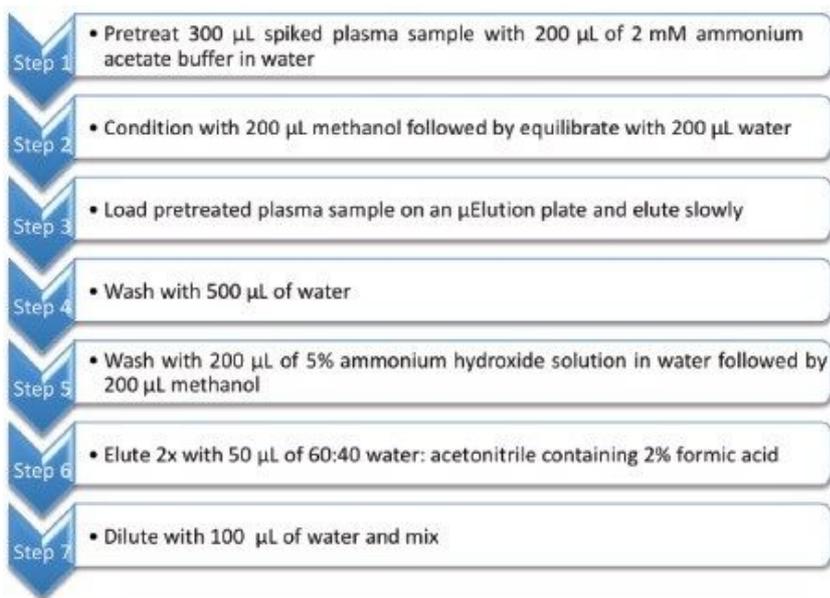
In this work, a selective, sensitive, and robust quantification method has been developed for the accurate quantification of linaclotide in plasma. This method uses UPLC separation, tandem quadrupole MS with ESI (electrospray ionization), and selective mixed-mode solid-phase (SPE) extraction to achieve a lower limit of quantitation (LLOQ) of 10.0 pg/mL from 300  $\mu$ L of plasma.

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## Experimental

### Sample preparation

Standards and quality control (QC) samples were prepared by spiking linaclotide into commercially available human plasma at various concentrations (10–4000 pg/mL). Calibration curve standards were prepared in duplicate, while 6 replicates of each quality control (QC) level were prepared. For each of the prepared samples, 300  $\mu$ L of plasma sample was mixed with 200  $\mu$ L of a 2 mM ammonium acetate solution containing 0.2% ammonium hydroxide. The pretreated plasma samples were extracted using the Waters Oasis MAX 96-well  $\mu$ Elution plate. For this extraction, the entirety of the pretreated plasma samples was loaded onto the plate. Following sample loading, samples were washed with 200  $\mu$ L of 5% Ammonium hydroxide in water, followed by 200  $\mu$ L of methanol solution. Linaclotide was then eluted from the sorbent using 100  $\mu$ L of the elution solvent (two aliquots of 50  $\mu$ L). The extracted samples were then diluted with 100  $\mu$ L water for a final sample volume of 200  $\mu$ L and injected onto the LC-MS/MS system. This protocol is shown in Figure 3.



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Figure 3. Oasis MAX 96-well  $\mu$ Elution plate extraction protocol for linaclotide from human plasma samples.

### LC conditions

LC system:	ACQUITY UPLC I-Class
Column:	ACQUITY HSS PFP 130Å, 1.7 µm, 2.1 mm × 100 mm
Column temp.:	42 °C
Sample temp.:	15 °C
Injection volume:	10 µL
Mobile phase A:	0.2% Formic acid in water
Mobile phase B:	0.2% Formic acid in acetonitrile
Purge and wash solvent:	200:10:10:10 Acetonitrile:Methanol:Isopropanol:Water with 0.1% formic acid

Time (min)	Flow (mL/min)	Mobile phase A	Mobile phase B	Curve
Initial	0.200	10.0	90.0	6
0.20	0.200	10.0	90.0	6
5.00	0.200	30.0	70.0	6
5.10	0.400	32.0	68.0	6
6.50	0.400	80.0	20.0	6
8.00	0.400	10.0	90.0	6
9.00	0.400	10.0	90.0	6
9.10	0.200	10.0	90.0	6
10.00	0.200	10.0	90.0	6

Table 1. LC gradient for linaclotide separation.

## MS conditions

Mass spectrometer:	Xevo TQ-XS Tandem Quadrupole
Ionization:	ESI+
Capillary voltage:	3.5 kV
Source temp.:	150 °C
Desolvation temp.:	550 °C
Cone gas flow:	150 L/h
Desolvation gas flow:	1000 L/h

Collision cell pressure:  $3.6 \times 10^{-3}$  mbar

## Data management

Instrument control software: MassLynx (v 4.2)

Quantification software: TargetLynx

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## Results and Discussion

In this work, we have developed a complete sample preparation and UPLC LC-MS/MS method for sensitive and accurate quantification of linaclotide from plasma. SPE extraction of linaclotide from plasma was performed using the Oasis MAX, a mixed-mode sorbent in the 96-well  $\mu$ Elution plate format, to enhance selectivity of the extraction. Oasis MAX is a polymeric ion exchange sorbent which relies on both reversed-phase and ion-exchange retention mechanisms to selectively separate linaclotide from other plasma matrix components.

LC-MS/MS quantification was performed using a Waters Xevo TQ-XS Tandem/Triple Quadrupole MS coupled to an ACQUITY UPLC I-Class System. Reversed phase chromatographic separation was achieved with an ACQUITY UPLC HSS PFP 1.8  $\mu$ m Column, at an initial flow rate of 0.2 mL/min using a linear flow gradient (Table 1) with 0.2% formic acid in water and 0.2% formic acid in acetonitrile mobile phases here optimized and finalized saw tooth gradient to reduce the column carryover for this cyclic peptide. Linaclotide is a polar Basic compound. During initial method development, several reversed phase columns was evaluated for overall chromatographic performance. Best chromatographic performance for linaclotide (retention, peak shape, resolution from endogenous interferences, etc.) had achieved by using the HSS PFP column.

During MS optimization, several multiply charge precursors were observed. Of these, the 2+ ( $m/z$  764) precursor was the most predominant species. The 764 $\rightarrow$ 182 MRM transition was used as the primary quantification transition. In this assay, the use of the specific y-ion 182 fragment, derived from the doubly charged precursor, combined with the mixed-mode SPE methodology significantly improved assay specificity. Optimized MS conditions used for linaclotide quantification are listed in Table 2.

Analyte	Precursor ion (m/z)	Product ion (m/z)	Cone voltage (V)	Collision energy (eV)
Linaclotide	764.05	182.03	25	15

Table 2. MRM transitions, collision energies, and cone voltages for linaclotide.

The Xevo TQ-XS Mass Spectrometer, equipped with a novel step-wave ion guide and tool free ESI ion source, enabled improved ion sampling in the source and better ion transfer efficiency, while use of the method events within the software to divert flow to waste, reduced system contamination. Use of a low dispersion UPLC I-Class System and sub-2-micron ACQUITY UPLC Column with reverse gradient mechanism, coupled to the Xevo TQ-XS Mass Spectrometer, afforded excellent resolution from endogenous matrix components, enhancing selectivity and sensitivity. This sensitivity and selectivity of this SPE LC-MS method is illustrated in Figure 4 for the LLOQ, 10 pg/mL extracted plasma sample, achieving a signal to noise (S/N) ratio of 122.82.

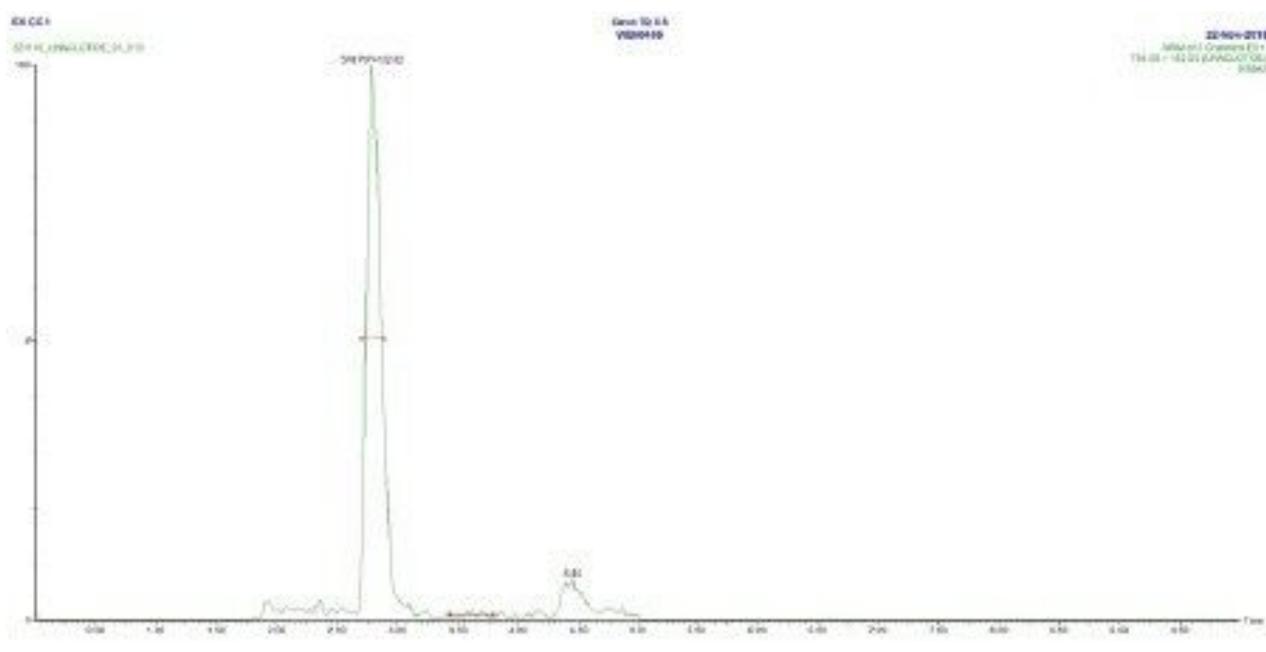


Figure 4. Representative linaclotide LLOQ (10.0 pg/mL) chromatogram, extracted from plasma, demonstrating excellent sensitivity and selectivity with S/N ratio of 122.82.

Using this SPE LC-MS method, quantitative performance was excellent. Calibration curves were linear ( $r^2 > 0.997$ ) from 10–4000 pg/mL with accuracies between 85–115 % and CVs <15% for all points on the curve.

Figure 5 illustrates this performance. At the same time, QC statistics easily met recommended bioanalytical method development guidelines,<sup>4</sup> with average precision <8% and accuracy between 97–110%. This QC performance is highlighted in Table 3 for precision and accuracy (PA) batches, while QC chromatographic performance is illustrated in Figure 6.

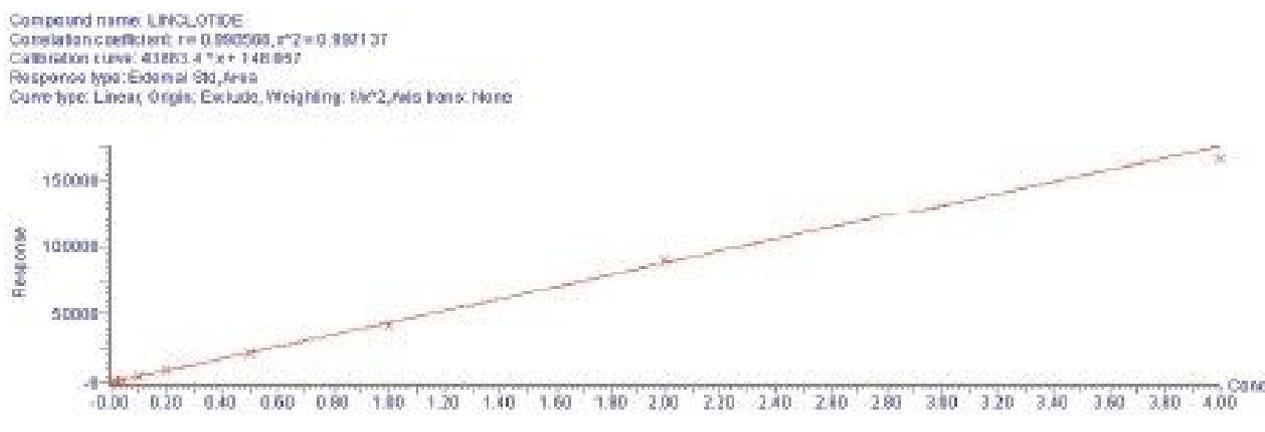


Figure 5. Representative lincolotide calibration curve (10.0 to 4000 pg/mL) extracted from plasma.

Batch name	Sample #	LLOQ (10.0 pg/mL)	LQC (20.0 pg/mL)	MQC2 (200 pg/mL)	HQC (3500 pg/mL)
P&A01	1	0.011	0.022	0.211	3.380
	2	0.010	0.021	0.224	3.387
	3	0.011	0.022	0.223	3.351
	4	0.010	0.021	0.221	3.461
	5	0.010	0.022	0.220	3.403
	6	0.009	0.022	0.219	3.407
	Mean conc. (pg/mL)	0.010	0.021	0.219	3.398
	STD	0.00075	0.00052	0.00463	0.03668
	% CV	7.40	2.38	2.10	1.07
	Mean % accuracy	102.27	107.78	100.83	97.10
P&A02	7	0.010	0.019	0.204	3.336
	8	0.011	0.021	0.206	3.434
	9	0.011	0.020	0.208	3.402
	10	0.010	0.023	0.209	3.483
	11	0.010	0.023	0.209	3.480
	12	0.010	0.022	0.216	3.485
	Mean conc. (pg/mL)	0.0103	0.0213	0.2087	3.4367
	STD	0.00052	0.00163	0.00408	0.05950
	% CV	4.99	7.65	1.95	1.73
	Mean % accuracy	100.30	107.07	104.38	98.18

Table 3. Summary of quality control results for linaclotide extracted from human plasma for P&A Batches 1 and 2.

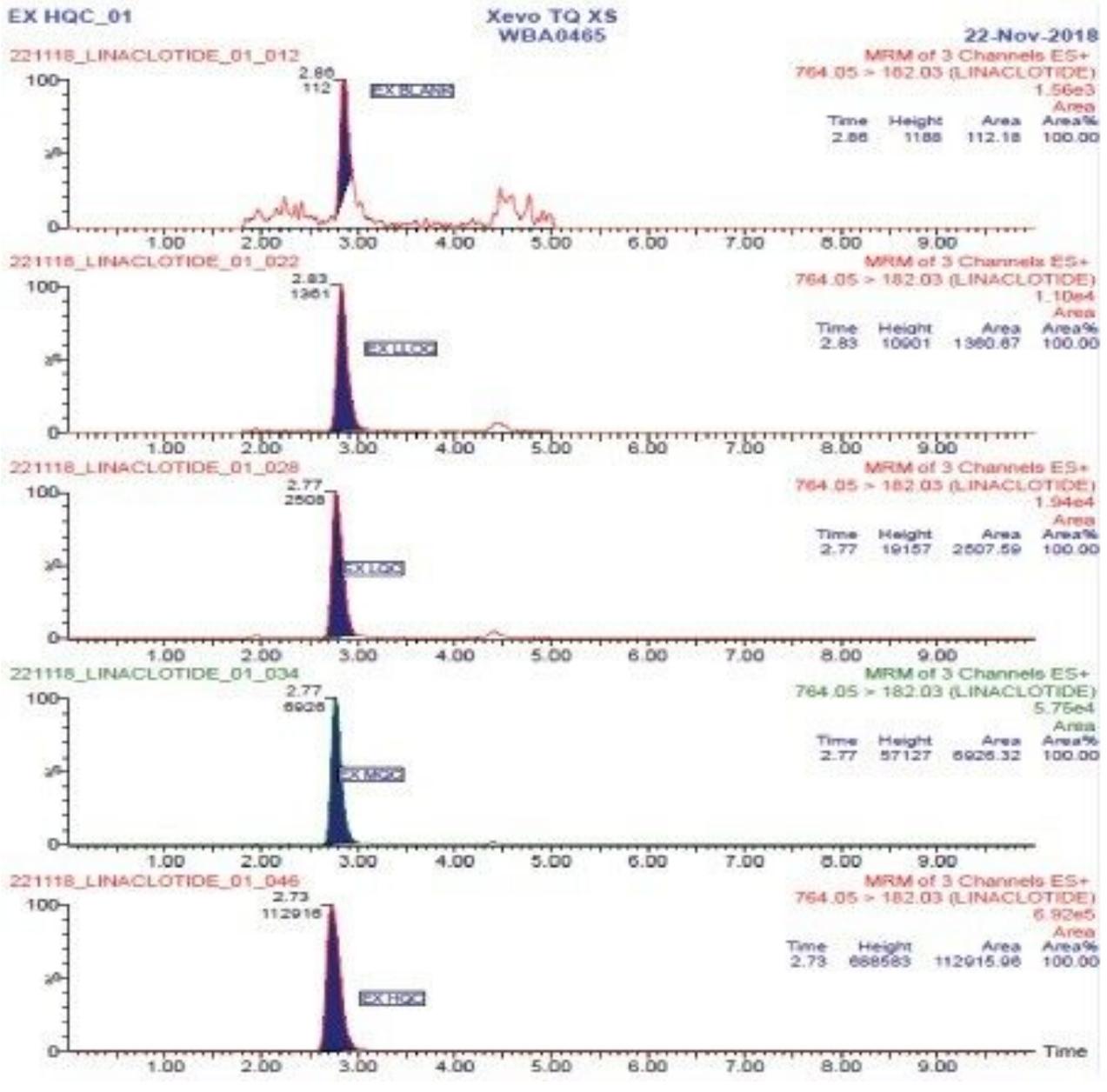


Figure 6. Representative QC's chromatograms for linaclootide extract from plasma, highlighting sensitivity and specificity.

## Conclusion

This application highlights the development of a highly sensitive and robust LC-MS assay for the

quantification of linaclotide extracted from plasma. The method described herein achieves an LLOQ of 10.0 pg/mL with a linear dynamic range of 10–4000 pg/mL. The high sensitivity and linearity of method was attributed to extraction specificity using mixed-mode  $\mu$ Elution SPE sample preparation, high-resolution UPLC chromatographic separation with the ACQUITY UPLC I-Class System using a sub-2-micron UPLC column, and sensitivity of the Xevo TQ-XS Mass Spectrometer. This developed method has demonstrated its fit-for-purpose use in support of drug discovery and research.

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