

Multipass Ion Mobility Separation of GLP-1 Receptor Agonist Impurities Using the SELECT SERIES™ Cyclic™ IMS

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Abstract

In this work, Cyclic Ion Mobility Mass Spectrometry was employed to separate, detect, and quantitate selected D- impurities of glucagon-like peptide 1 receptor agonist (GLP-1 RA) semaglutide that are otherwise difficult to differentiate using ultra performance liquid chromatography (UPLC™). The multipass cyclic IM capabilities of the Cyclic Ion Mobility Separation (IMS) bring enhanced mobility separation capabilities that are orthogonal to the LC separation and allow better characterization of complex impurity profiles such as those seen for GLP-1 therapeutics.

Benefits

- Resolve challenging semaglutide impurity isomers that are difficult to distinguish by LC-MS alone
 - Achieve baseline separation of highly similar isomers using multipass ion mobility and advanced IMSⁿ experiments
 - Enable confident impurity identification and characterization for complex formulation samples
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Introduction

The use of GLP-1 RA as therapeutics for type 2 diabetes and obesity have gained significant attention, resulting in unprecedented demand. As a result, there is an increased pressure on manufacturers for continuous improvement of their production processes to ensure efficiency, safety, and meet growing demands of an already enormous market.

One of the most challenging aspects in the development and production of GLP-1 RAs is in the characterization of impurities. While the typical impurity profile, along with pertinent regulations, is dependent on the manufacturing process (biotechnology or synthetic), what remains constant is the difficulty in separating possible impurities. Many of these are single amino acid configuration substitutions (L to D) on a peptide with 30 amino acid residues or more (which are often already modified with long fatty acid chains), which can require lengthy LC gradients and specialized mobile phases to achieve separation. Even then, in some cases, there is co-elution of impurities, which further complicates characterization and calls for orthogonal separation methods.

In this work, cyclic ion mobility with scalable resolution was evaluated as an orthogonal separation technique and as means to further elucidate impurity profiles on a typical GLP-1 agonist API (semaglutide).

Experimental

Sample Description

Standards of semaglutide API (all L- amino acids), along with D-Phe6, D-Ala19, and D-Arg30 impurities

were obtained synthetically and previously characterized. These specific impurities were chosen because, while there's LC separation between them and the API, all three impurities co-elute using the 60-minute method developed for this assay and are very hard to resolve from one another utilizing LC alone. Figure 1 illustrates the structure of semaglutide and these impurities.

Direct infusion experiments were used to optimize Cyclic IMS. Peptides were solubilized in ultrapure water to 1 mg/mL (~243 µM), then diluted to 1 µM in water/acetonitrile with 0.1% formic acid. For LC-IMS-MS analysis, peptides were solubilized in 100 mM Tris HCl pH 7.5 to 1 mg/mL (~243 µM), then diluted to 0.1 mg/mL (~24.3 µM) and transferred to QuanRecovery™ Vials for injection.



B: 2-aminobutyric acid

X: Lys with linker and fatty acid

RED: possible sites of single D- amino acids

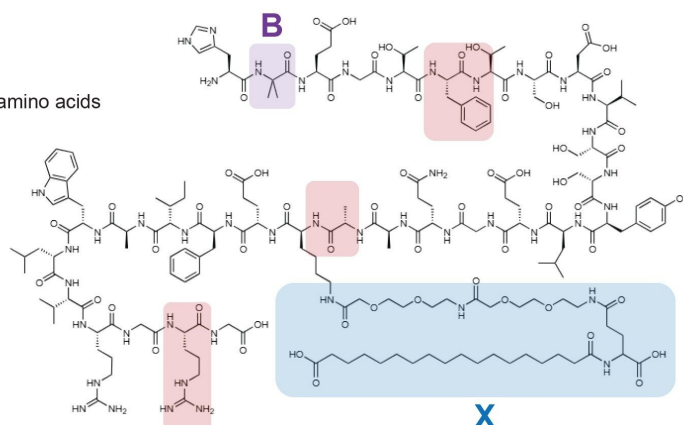


Figure 1. Structure of semaglutide API and D-amino acid impurities D-Phe6, D-Ala19, and D-Arg30.

Method Conditions

LC separations were carried out on an ACQUITY™ Premier UPLC System with an Quaternary Solvent Manager (QSM), using a ternary 60-minute gradient of water, acetonitrile, and methanol with 0.1% trifluoroacetic acid (TFA), each on an ACQUITY Premier CSH™ C18 Column 2.1 × 150 mm, 1.7 µm held at 60 °C.

MS Conditions

MS system:	SELECT SERIES Cyclic IMS
Ionization mode:	ESI+
Acquisition range:	<i>m/z</i> 50 – 2000
Capillary voltage:	2.50 kV
Cone voltage:	40
Quad profile:	600 (25/25) / 700 (25/25) / 800

All other remaining Cyclic IMS settings were installation defaults.

Data Management

Chromatography software:	MassLynx™ Software
MS software:	MassLynx Software
Informatics:	DriftScope™ Software

Results and Discussion

To investigate the mobility separation of the target impurity isomers, diluted (1 μM) solutions of the API and each impurity were first analyzed by direct infusion or as a mixture containing all three impurities. Direct infusion of standards is typically the preferred approach, as it enables optimization of mobility separations and advanced ion mobility experiments under a stable analyte signal. Features such as the pass calculator can help streamline method development and mobility optimization, providing an effective alternative when standards are not available.

Both API and D- impurities were observed mainly as $[M+3H]^{3+}$ species at m/z 1372, with some $[M+4H]^{4+}$ present as well at m/z 1029. Both charge states were investigated for IM separations, but much better results were observed for the $[M+4H]^{4+}$ ions so the discussion will focus on them.

For an equimolar mixture of D-Phe₆, D-Ala₁₉, and D-Arg₃₀ impurities, a multipass experiment with seven passes generated a distribution (Figure 2) containing two unresolved peaks (50.40/51.42 ms) and an almost fully resolved peak (52.27 ms). Analysis of the isolated impurities confirmed the identity of the first two unresolved peaks as D-Arg₃₀ (50.40 ms) and D-Ala₁₉ (51.42 ms), with D-Phe₆ at 52.27 ms.

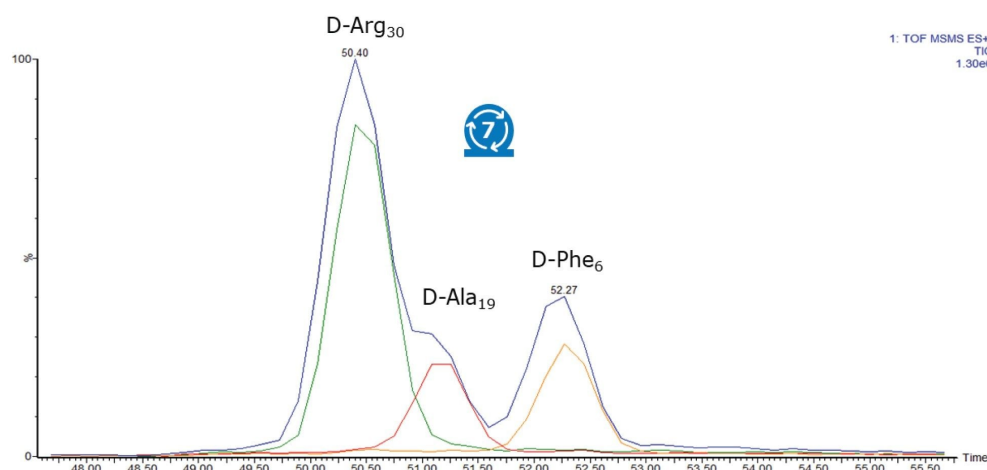


Figure 2. Multipass separation (7 passes) of an equimolar mix of the D-amino acid impurities, generating a distribution with two unresolved peaks (50.40/51.42 ms) and an almost fully resolved peak (52.27 ms).

Increasing the number of passes beyond seven didn't improve the separation of the unresolved isomers (data not shown). To achieve greater separation and distinguish the two unresolved peaks, a slicing IMSn experiment was performed. Following an initial seven-pass separation, the mobility region containing the first two peaks was isolated and stored in the pre-array before the mobility racetrack. After the remaining ions were ejected, the selected ion population was reinjected and subjected to additional multipass ion mobility separation. This targeted approach enabled baseline separation of the D-Arg₃₀ and D-Ala₁₉ isomers after a further 18 passes (Figure 3). This illustrates the power of IMSn and flexibility of the Cyclic IMS.

Having demonstrated separation of all three impurity isomers by infusion, the workflow was transferred to LC-IM/MS to show that the same mobility methods can be applied to more complex analytes required for samples representative of real formulations. A MS method was created for the Cyclic IMS containing two functions: a seven-pass function to resolve D-Phe6, and a slicing/18-pass function to resolve the D-Arg30 and D-Ala19 isomers. To better visualize data, the DriftScope Software was used to extract the mobility chromatogram for a single LC peak containing all three impurities at around 42 minutes. The results closely reproduced what was observed for direct infusion (Figure 4), confirming that same performance is obtained with an LC-IM/MS experiment.

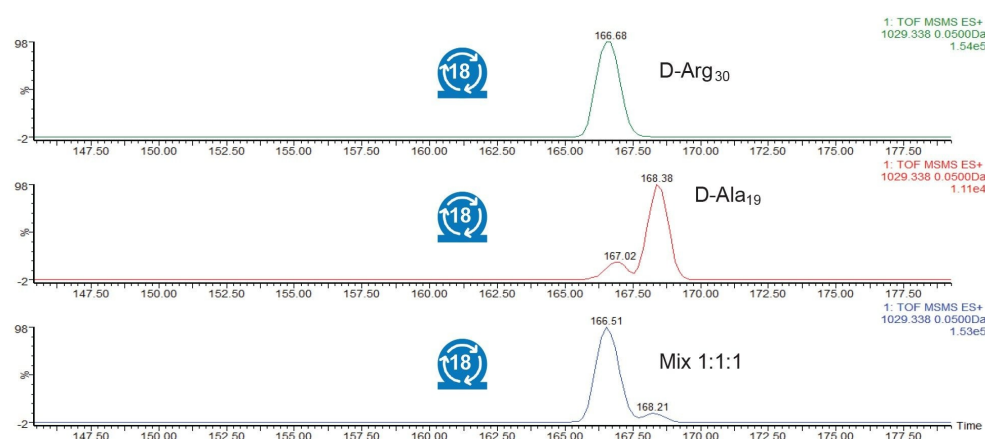


Figure 3. "Slicing" mobility experiment to further baseline separate D-Arg30 and D-Ala19. The mobility region containing these two isomers was isolated and stored in the pre-array, then reinjected and subsequently separated with 18 passes.

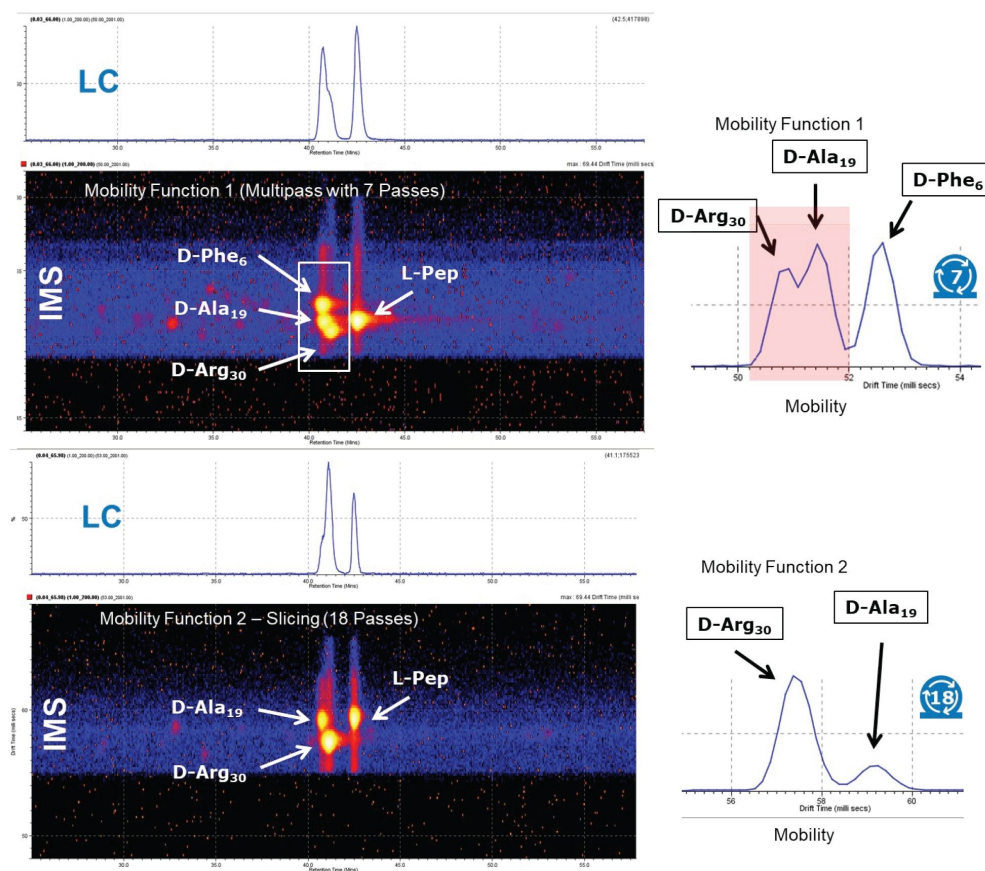


Figure 4. Advanced Cyclic IMS separation of semaglutide D-amino acid impurities co-eluting in the LC dimension. Top shows a multipass (7 pass) separation of a mixture of D-Arg30, D-Ala19, and D-Phe6 impurities of semaglutide. Bottom shows a subsequent slicing (18 pass) experiment on the unresolved distribution obtained for D-Arg30/D-Ala19 (highlighted on the top panel), yielding baseline separation. Both mobility experiments are acquired simultaneously during the LC-IM/MS run in different channels, and XICs can then be used to quantitate the impurities if desired.

Conclusion

The SELECT SERIES Cyclic IMS enabled the separation and identification of single D-amino acid

substitution impurities of semaglutide that are highly similar in structure and difficult to distinguish using LC-MS alone. By combining multipass ion mobility with slicing IMSⁿ experiments, baseline-resolved mobility peaks were obtained for all three isomers, providing a robust approach for confident impurity characterization in complex formulation samples.

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