Innovations on the QTRAP® 4500, 5500, 6500 and 6500+ LC-MS/MS Systems

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Mass spectrometry has transformed quantitative analysis to become the method of choice for many assays. More recently, LC-MS/MS has revolutionized quantitative bioanalysis. While single MS filtering offers advantages over non-mass selective techniques, the use of tandem mass spectrometry (MS/MS, or MS2) eliminates interferences and results in a dramatic increase in selectivity which yields a very low baseline, excellent limits of quantification, and very good linearity. As a result, the Multiple Reaction Monitoring (MRM) experiment performed on triple quadrupole mass spectrometers has become the technique of choice for highly sensitive and selective quantification in biological matrices.

In some cases, interferences cannot be eliminated using MRM. More elaborate sample cleanup and chromatography is required to eliminate these interferences. If a high baseline or matrix interference cannot be eliminated, the result is a compromised Lower Limit of Quantification (LOQ) as the detection of compounds in complex matrices is limited by signal-to-noise rather than by raw instrument response. In such cases, the addition of a third MS stage has been shown to greatly increase selectivity and eliminate the high baseline or chromatographic interference. The result is a lower LOQ and better chromatographic peak shape.

Key Features of the QTRAP® 4500, 5500, 6500 and 6500+ Systems for MRM³ Quantification

- MRM³ quantification - An MS³ scan is performed with a fast cycle time and using a narrow scan range centered at the second generation product ion m/z which is being used for quantification. This type of scan is referred to as MRM³ (Figure 1).
- Faster linear ion trap scan speeds – Scan speeds up to 20,000 Da/sec enable MS³ scans with an HPLC compatible cycle time such that extracted ion chromatograms (XICs) of second generation product ions can be extracted and integrated with a sufficient number of data points across the chromatographic peak.
- Better in-trap fragmentation – The new Linear Accelerator™ Trap with pulsed gas valve implemented in the QTRAP® systems provides faster, more efficient in-trap fragmentation (Figure 2)
- High ion trap sensitivity – The QTRAP systems feature high sensitivity when operating in linear ion trap mode
- High selectivity – Unit isolation of precursor ion in Q1 followed by excitation and fragmentation at unit resolution in the ion trap provides the very high selectivity in MRM³ analysis (Figure 3).
### Speed

The speed and efficiency of ion-trap fragmentation has also been greatly enhanced on the QTRAP® 4500, 5500, 6500 and 6500+ systems. Collision gas is now introduced through a high speed pulsed gas valve that enables a rapid increase in pressure in the LIT (Figure 2). Together with an increase in the RF drive frequency, this pulsed gas results in increased fragmentation efficiency and reduced excitation time of 25 ms or less.

In addition, the scan speed of the linear ion trap has been increased to a maximum of 20,000 through the use of the faster eQ™ electronics. This enabling fast MRM³ scan cycles at very high sensitivity.

### Selectivity

Because of the configuration of the QTRAP® systems, the MRM³ quantification workflow is highly flexible and leverages the sensitivity and selectivity of the system. For the MRM³ workflow, the analyte ion of interest is first isolated in the Q1 quadrupole with a user-selected resolution, usually unit resolution (0.7 Th FWHH). It is then fragmented in the Q2 collision cell, providing a broad range of product ions to be selected in the ion trap.

The in-trap fragmentation is achieved through the application of a single wavelength / narrow band excitation. As shown in Figure 3, this allows very selective fragmentation. The C12 isotope of a product ion can be specifically excited and fragmented to completion with minimal fragmentation of the C13 isotope. This provides further selectivity advantages in the removal of interfering background.

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**Figure 2. Pulsed Gas Valve.** Gas is introduced into the linear ion trap using a high speed pulsed gas valve which rapidly increases LIT pressure and reduces required excitation time for ion trap fragmentation.

**Figure 3. Single Frequency Excitation for High Selectivity.** Narrow band excitation is used to specifically excite and fragment just the C12 isotope of the ion isolated in the LIT (left). This isotope can be fragmented to completion with no impact on the nearby C13 isotopes (right).
Sensitivity

Linear Accelerator™ Trap technology has resulted in ground breaking improvement in the handling of ions inside the linear ion trap of the QTRAP® 4500, 5500, 6500 and 6500+ systems, resulting in up to 100x more sensitivity. Trapped ions are manipulated within the linear ion trap through the use of auxiliary DC fields provided by the addition of small electrodes (Figure 4, top). Ions are gently moved toward the extraction region of the linear ion trap during the cooling period by a voltage applied to the trap collar. A potential barrier is created by increasing the potential on the auxiliary electrodes just before the mass scan to complete the ion concentration process. The application of this axial field has a significant effect on sensitivity (Figure 4, bottom left).

In addition, a radio frequency is applied to the exit lens of the Linear Accelerator Trap resulting in further sensitivity gains (Figure 4, bottom right). These two innovations enable better than unit resolution to be obtained in the trap scan modes at these very high scan speeds.

Removal of Tough Interferences

Innovations in scanning speed, selectivity and sensitivity on the QTRAP systems enable successful implementation of the MRM³ workflow for a wide range of analytes. Sometimes, background noise or interferences can limit the detection of an analyte. Shown in Figure 5 is an example of an interference that has the same MRM transition as Clenbuterol and elutes at the same retention time. Use of MRM³ can completely remove this interference and enable a much lower detection of this analyte.

Figure 4. Linear Accelerator™ Trap Innovations for Sensitivity. Addition of electrodes in this new trap design (top) significantly improves the sensitivity of trap scanning by moving the trapped ions into the extraction region before axial ejection from the trap (bottom left). Further sensitivity gains are achieved by the addition of RF to the exit barrier of the trap (bottom right).
Conclusions

- MRM³ is an effective strategy for quantitation of analytes when high background or interferences make standard MRM quantitation difficult (Figure 5).
- MRM³ can be used to achieve similar LOQ’s with less sample preparation and simplified or faster chromatography.
- MRM³ has been successfully applied to the detection and quantitation of small molecules, peptides, and protein biomarkers.
- MRM³ is significantly improved on the QTRAP® systems, making them useful tools for quantitation in tough matrices.
- The QTRAP® 4500, 5500, 6500 and 6500+ LC-MS/MS systems are high performance triple quadrupole and linear ion trap systems providing users with many powerful quantitative and qualitative tools.

References