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# Multiple ion isolation and accumulation events for selective chemical noise reduction and dynamic range enhancement in MALDI imaging mass spectrometry†

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Abundant chemical noise in MALDI imaging mass spectrometry experiments can impede the detection of less abundant compounds of interest. This chemical noise commonly originates from the MALDI matrix as well as other endogenous compounds present in high concentrations and/or with high ionization efficiencies. MALDI imaging mass spectrometry of biological tissues measures numerous biomolecular compounds that exist in a wide range of concentrations *in vivo*. When ion trapping instruments are used, highly abundant ions can dominate the charge capacity and lead to space charge effects that hinder the dynamic range and detection of lowly abundant compounds of interest. Gas-phase fractionation has been previously utilized in mass spectrometry to isolate and enrich target analytes. Herein, we have characterized the use of multiple continuous accumulations of selected ions (Multi CASI) to reduce the abundance of chemical noise and diminish the effects of space charge in MALDI imaging mass spectrometry experiments. Multi CASI utilizes the mass-resolving capability of a quadrupole mass filter to perform multiple sequential ion isolation events prior to a single mass analysis of the combined ion population. Multi CASI was used to improve metabolite and lipid detection in the MALDI imaging mass spectrometry analysis of rat brain tissue.

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## Introduction

Matrix-assisted laser desorption/ionization (MALDI) imaging mass spectrometry is a valuable tool for biological tissue analysis, enabling the detection of a wide variety of metabolites, lipids, and proteins over a range of concentrations.<sup>1–5</sup> Extending the limit of detection and dynamic range for the spatial mapping of lowly abundant compounds in tissue is a major analytical challenge in imaging mass spectrometry experiments.<sup>6–9</sup> Oftentimes, MALDI analyses of biological tissues produce highly abundant chemical noise, which suppresses the detection of lowly abundant ions of interest. Chemical noise can originate from the MALDI matrix and/or from abundant endogenous compounds in the tissue environment.<sup>10–13</sup> This issue is exacerbated in instrument configurations that utilize ion trapping devices that have finite charge capacities. The presence of highly abundant ions can introduce space charge effects that detrimentally affect the

limits of detection and dynamic range for more lowly abundant compounds.<sup>14,15</sup>

A variety of approaches have been developed to mitigate chemical noise encountered during biological tissue analysis. For example, orthogonal separation using liquid chromatography (LC) prior to mass spectrometry is routinely used to increase the peak capacity of analysis.<sup>16–23</sup> However, LC separation technologies are largely not directly compatible with imaging mass spectrometry due the lengthy timescales needed for each chromatographic separation.<sup>24</sup> Condensed-phase tissue washing and chemical derivatization approaches are performed prior to subjection to *in vacuo* conditions of the mass spectrometer and can be used to selectively remove undesired compounds that may suppress ion signals of interest and selectively improve the ionization efficiency of targeted analytes, respectively. For example, tissue washing using volatile buffers at different pH ranges has been used to enhance the detection of neutral lipids and proteins.<sup>25–27</sup> However, harsh solvent conditions can remove tissue content and further distort the natural morphology of the tissue environment.<sup>27</sup> Chemical derivatization with reagents, such as 1-(4-(aminomethyl)phenyl)pyridin-1-ium chloride (AMPP), can be performed on tissue, but can suffer from incomplete and side reactions that complicate the chemical milieu.<sup>28,29</sup> These con-

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densified-phase approaches also require additional time-consuming steps to sample preparation protocols.

Gas-phase separation techniques offer rapid and flexible means of fractionating ion populations after ionization, but prior to detection, without manipulating the original tissue sample. Ion mobility-mass spectrometry (IM-MS) is widely recognized for its ability to separate ions based on their collisional cross section (CCS) and is routinely applied to imaging mass spectrometry studies.<sup>30–33</sup> This method of gas-phase separation greatly increases the peak capacity of the analysis, but requires the presence of an ion mobility device on the mass spectrometer. Other gas-phase fractionation strategies include the use of selective ion/ion reactions that allow separation of isobaric (*i.e.*, same nominal  $m/z$ ) compounds on the basis of gas-phase chemical reactivity.<sup>34,35</sup> More broadly applicable gas-phase separation strategies involve the use of selective ion isolation using a mass resolving quadrupole on a hybrid mass spectrometer.<sup>36–39</sup> The instrument used herein is a quadrupole-hexapole Fourier transform ion cyclotron resonance (QhFT-ICR) hybrid mass spectrometer, which accumulates ions in the hexapole ion trap prior to transfer to the ICR for mass analysis. RF/DC isolation can be performed as ions pass through the quadrupole mass filter (QMF) to provide for mass-selective ion accumulation in the hexapole ion trap.<sup>40,41</sup> More recently, mass-selective ion accumulation has been termed continuous accumulation of selected ions (CASI) and used to improve limit of detection and dynamic range in imaging mass spectrometry experiments.<sup>36,42–44</sup> Conversely, selected ion ejection (SIE) can be performed by applying a supplemental alternating current (AC) waveform to the QMF to eject a small  $m/z$  interval as the ions transmit through the device.<sup>45</sup> SIE has been employed to selectively exclude highly abundant chemical noise and the subsequent broadband enrichment of ions of interest in imaging mass spectrometry experiments.<sup>36</sup>

Most applications of CASI in imaging mass spectrometry have used a singular mass isolation window for ion accumulation.<sup>36,43</sup> However, multiple ion isolation windows can be used to selectively target multiple analytes simultaneously.<sup>46,47</sup> For example, broadband dipolar AC isolation waveforms that contain multiple frequency notches matched with the secular frequencies of ions of interest can be applied to quadrupole ion traps and quadrupole mass filters to effect multiple mass isolation windows.<sup>48,49</sup> While these approaches allow for the simultaneous isolation of multiple windows, a broadband AC waveform is required. Multiple sequential isolation windows can also be affected by altering the RF/DC settings of the QMF. The resulting ion populations can then be collected in a downstream ion trap and analyzed in a single mass analysis step; a process that has been termed Multi CASI. Multi CASI can be used to selectively exclude one or more interfering ions, while still allowing for broadband enrichment at  $m/z$  values below and above the excluded  $m/z$  window. This process has recently been used to quantify pharmaceuticals in MALDI analysis of biological samples,<sup>12</sup> but has not seen broader adoption in imaging mass spectrometry

and its analytical performance has not been rigorously characterized. Herein, we have used Multi CASI to exclude the signal from the MALDI matrix (*i.e.*, between  $m/z$  150 and 250), while still simultaneously capturing metabolite ion signals below  $m/z$  150 as well as metabolite and lipid ion signals above  $m/z$  250. This reduces chemical noise and increases the limits of detection and dynamic ranges of ions of interest. The figures of merit of full scan, single CASI, and Multi CASI workflows are compared here to assess their analytical utility in imaging mass spectrometry experiments.

## Experimental

### Materials

1,5-Diaminonaphthalene (DAN), hematoxylin, eosin, ammonium acetate, and chloroform were purchased from Sigma-Aldrich Chemical Co. (St Louis, MO, USA). 2-Propanol and methanol were purchased from Fisher Scientific (Pittsburgh, PA, USA). Rat brain (Brown Norway) was purchased from BioIVT (Westbury, NY, USA) and stored at  $-80\text{ }^{\circ}\text{C}$  until analysis.

### Tissue extraction

Microscope slides with uniform metabolite and fatty acid coatings were prepared using an automated robotic sprayer. Metabolites and fatty acids were extracted from rat brain tissue ( $\sim 300\text{ mg}$ ) using the Folch method.<sup>50</sup> Briefly, tissue was homogenized using a FastPrep-24 Classic (MP Biomedicals, LLC, Irvine, CA, USA) with  $50\text{ }\mu\text{L}$  of  $5\text{ mM}$  ammonium acetate per  $10\text{ mg}$  of tissue. The homogenate was centrifuged and  $1\text{ mL}$  of supernatant was collected. To the supernatant,  $4\text{ mL}$  of methanol and  $8\text{ mL}$  of chloroform were added. Aqueous and organic layers were separated and dried with nitrogen gas. The aqueous portion was reconstituted in  $600\text{ }\mu\text{L}$  of methanol and the organic portion was reconstituted in  $600\text{ }\mu\text{L}$  of  $80:20\text{ v/v}$  methanol:2-propanol. These solutions were mixed and  $1\text{ mL}$  of the mixture was applied to a glass slide using an M5 TM Sprayer (flow rate:  $0.15\text{ mL min}^{-1}$ , nitrogen flow:  $10\text{ psi}$ , spray temperature:  $40\text{ }^{\circ}\text{C}$ , four passes with offsets and rotations, spray velocity:  $1300\text{ mm min}^{-1}$ , dry time:  $2\text{ seconds}$ ) (HTX Technologies, LLC, Chapel Hill, NC, USA). A DAN MALDI matrix layer was then applied using a custom-built sublimation apparatus ( $120\text{ }^{\circ}\text{C}$ ,  $8\text{ min}$ ,  $<70\text{ mTorr}$ ).<sup>51,52</sup> Approximately  $2\text{ mg}$  of DAN was sublimated onto each microscope slide. DAN was used as the MALDI matrix for all experiments due to its broad compatibility for many analytes, including the low mass metabolites and fatty acids studied herein. The most abundant of chemical noise from this matrix is in the low  $m/z$  mass range, which typically complicates analysis of low mass metabolites and nicely demonstrates the capabilities of Multi CASI. However, we have also performed Multi CASI using other MALDI matrices (data not shown) and have observed similar improvements in performance to those described herein.

## Mass spectrometry

All mass spectrometry experiments were performed on a commercial hybrid 7T solariX XR FT-ICR mass spectrometer equipped with an Apollo II dual MALDI/ESI source along with a dynamically harmonized ParaCell (Bruker Daltonics, Billerica, MA).<sup>53</sup> The MALDI ion source utilizes a Smartbeam II Nd:YAG laser system (2 kHz, 355 nm). Mass analysis was performed using a 0.4893-second time-domain transient length, resulting in a mass resolving power of  $\sim 200\,000$  at  $m/z$  130.069. Mass spectra were recorded from  $m/z$  100 to 600 and were visualized in Compass DataAnalysis 5.0 (Bruker Daltonics, Billerica, MA).

Full scan and Multi CASI analyses were compared using the same acquisition conditions, where the only difference between the two experiments was the addition of Multi CASI. During full-scan analysis, no mass isolation is performed. Multi CASI utilizes multiple sequential mass isolation windows to isolate ion populations of disparate  $m/z$  prior to collection of the ion populations in a downstream hexapole ion trap (Fig. 1). For MALDI analysis, the total number of laser shots defined for the acquisition is divided evenly between the sequential mass isolation windows. Following accumulation in the hexapole ion trap, the mass selected ions are jointly transferred to the ICR cell for mass analysis. Full scan and single CASI methods using a defined number of laser shots are compared with Multi CASI methods using double the number of laser shots. This ensures that each isolation window has the same number of laser shots as the full scan and single CASI methods.

## Imaging mass spectrometry

Transverse sections of a control Brown Norway rat brain were collected at 10  $\mu\text{m}$  thickness using a Leica CM 3050S Research Cryostat (Leica Biosystems, Buffalo Grove, IL, USA), thaw mounted onto indium tin oxide (ITO)-coated slides (Delta Technologies, Loveland, CO, USA), and warmed to room temperature in a desiccator for 30 minutes. Following desiccation, a 1,5-diaminonaphthalene (DAN) MALDI matrix layer was applied using a custom-built sublimation apparatus (120  $^{\circ}\text{C}$ , 8 min,  $<70$  mTorr).<sup>51,52</sup> Approximately 2 mg of DAN was sublimated onto each slide.

Metabolite and free fatty acid imaging mass spectrometry of rat brain tissue was performed in negative ion mode. Full scan and Multi CASI experiments were performed to deplete chemical noise and to enrich selected analyte ions. Images were acquired using pixel spacings of 150 or 300  $\mu\text{m}$  in both the  $x$  and  $y$  dimensions (1000 laser shots, 135  $\mu\text{m}$  smart walk raster). Serial tissue sections were stained using hematoxylin and eosin (H&E) and scanned at 10 $\times$  magnification using a Zeiss Axio Imager M2 Microscope (Carl Zeiss Microscopy LLC, White Plains, NY, USA). Data analysis of individual spectra was performed using DataAnalysis (Bruker Daltonics, Billerica, MA, USA). Analyte identifications were assigned using high resolution accurate mass measurements (HRAM) with a tolerance of less than 5 ppm error along and searched against The Human

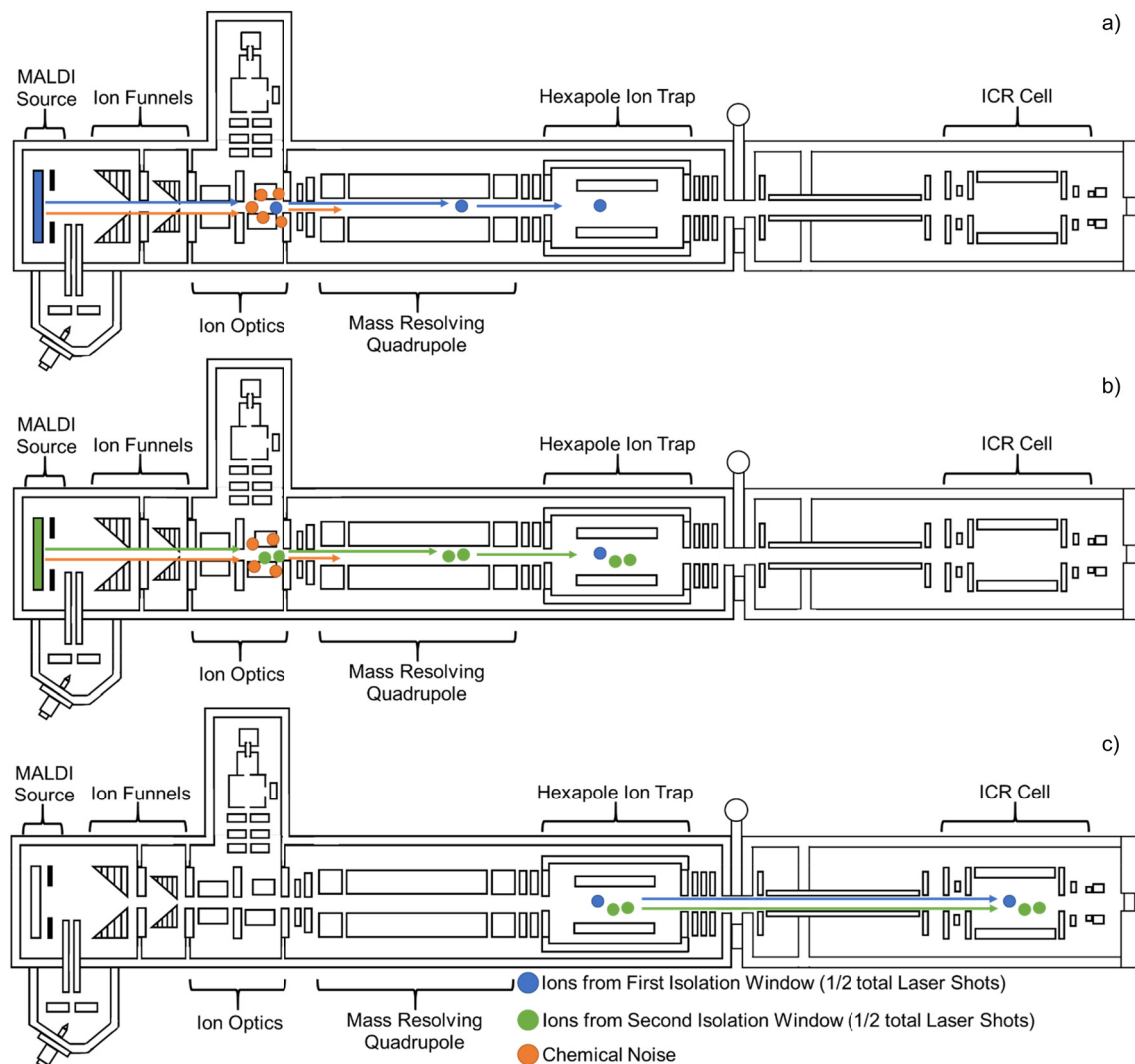
Metabolome Database (<https://hmdb.ca/>). Data analysis of imaging experiments was performed using flexImaging (Bruker Daltonics, Billerica, MA, USA).

## Results

CASI using a single mass isolation window has previously been shown to improve the limit of detection in imaging mass spectrometry experiments.<sup>36</sup> Multiple continuous accumulations of selected ions (Multi CASI) uses multiple sequential RF/DC isolation windows in a QMF to selectively accumulate disparate  $m/z$  ion populations in a downstream hexapole ion trap. The resulting ion population is then transferred to the ICR cell and mass analyzed in a single acquisition (Fig. 1). When using MALDI, the total number of laser shots for Multi CASI acquisition is divided evenly between the successive mass isolation windows. Multi CASI methods generally enrich populations of selected ions, while preventing the accumulation of ions outside of the selected  $m/z$  windows through quadrupole isolation events. This more efficiently fills the charge capacity of the ion trap with analytes of interest, improving the limit of detection and dynamic range. Both the hexapole and ICR cells are ion trap devices with finite charge capacities. When the space charge limit is reached, the confinement of ions can be diminished, and peak abundances may be distorted in the mass spectrum. Space charge artifacts in the ICR cell may manifest *via* peak coalescence and peak shifting.<sup>54–57</sup> The multiple mass isolation windows employed during Multi CASI reduces the number of ions transmitted to the hexapole and ICR cell, which significantly reduces space charge effects and allows for greater accumulation of the ions within the selected  $m/z$  windows.

In MALDI mass spectrometry, Multi CASI can be used to improve the limit of detection and dynamic range for lowly abundant compounds of interest. MALDI mass spectrometry analysis was performed on a complex lipid mixture extracted from rat brain tissue. Full scan and Multi CASI spectra were collected to demonstrate the depletion of chemical noise and enrichment of selected  $m/z$  regions (Fig. 2). The Multi CASI method excluded much of the chemical noise originating from the DAN matrix peak ( $m/z$  157.007, 0.172 ppm) and related isotopic peaks. The depletion of this chemical noise led to enrichment of the ions in the  $m/z$   $131 \pm 5$  Da and  $239 \pm 40$  Da mass regions selected by the Multi CASI windows. These mass windows were selected due to their proximity to the DAN  $[\text{M} - \text{H}]^-$  peak ( $m/z$  157.007). The proximity of these isolation windows to the primary noise peak allows for a rigorous evaluation of chemical noise exclusion. However, Multi CASI is also effective at other mass ranges (*e.g.*,  $m/z$  700–1000 to sample phospholipids, data not shown), depending on the chemical noise to be excluded.

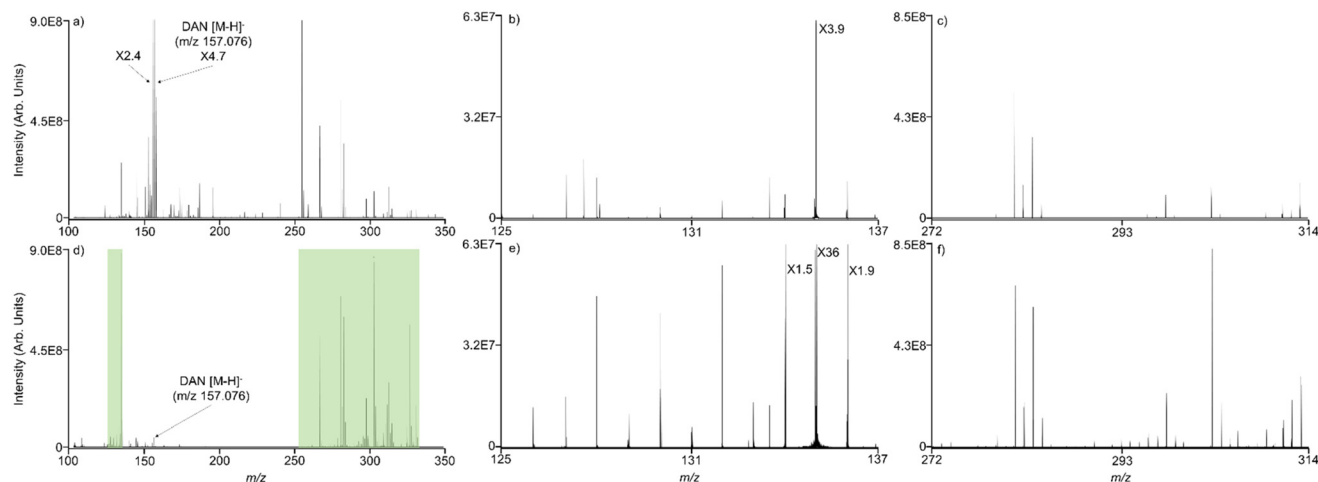
The extent of signal enrichment was monitored as a function of the number of MALDI laser shots accumulated in full scan and Multi CASI methods (Fig. 3). Ion abundance is expected to increase linearly with the number of accumulated



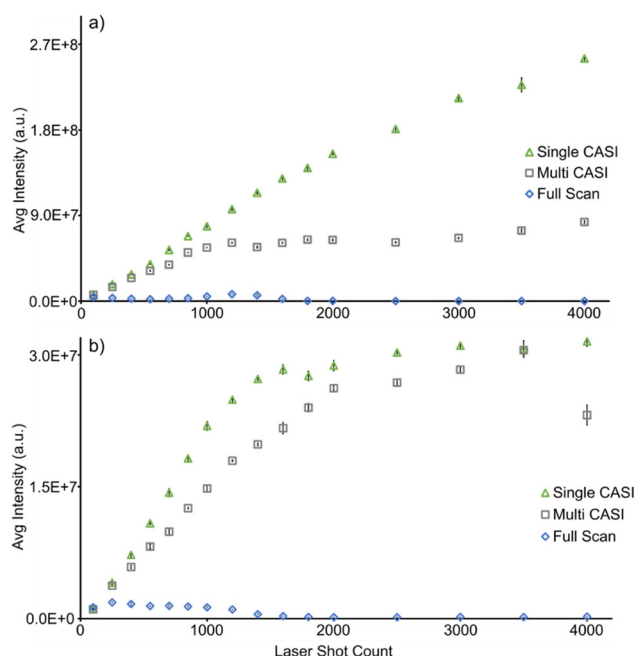
**Fig. 1** A simplified instrument schematic of the 7T solariX FT-ICR platform. (a) During Multi CASI, the first packet of MALDI ions is sequentially isolated in the QMF and stored in the hexapole ion trap. (b) The second packet of MALDI ions is then similarly isolated and mutually stored with the first MALDI event ions in the hexapole ion trap. (c) Both ion populations are sent to the ICR cell for mass analysis.

laser shots, until space charge effects are encountered, which will result in a disruption of the linear trend. Space charge is apparent with as few as 250 laser shots in the full scan experiment, as detected by a decrease in signal intensity for aspartate ( $m/z$  132.030, 0.041 ppm) and eicosadienoate ( $m/z$  307.264, 0.586 ppm) ions of interest (Fig. 3). Conversely, Multi CASI allowed for improvement of over 500-fold in signal intensity using 4000 laser shots for aspartate (present in the first Multi CASI window, Fig. 3a) and over 180-fold improvement in signal intensity using 3500 laser shots for eicosadienoate (present in the second Multi CASI window, Fig. 3b) compared to the full scan measurements. This is due to the reduction in ion population through chemical noise exclusion. The ion population reduction then allows for the traps to be filled with larger amounts of the selected ions. Multi CASI is also able to reduce variance in the data. The average relative standard deviation was 6.35% for full scan, 1.85% for single CASI, and 1.64% for

Multi CASI (Fig. 3). Space charge effects cause ions to behave unpredictably in ion traps at capacity, leading to detrimental effects such as reduced dynamic range, higher variability, and peak shifting.<sup>14,58,59</sup> Multi CASI is able to exclude chemical noise and therefore reduce the ion population in the instrument. A lower ion population in the ion trap minimizes the repulsion effects of space charge, lowering the overall standard deviation of the data. A reduction in the accumulation of chemical noise also allows for the detection of additional analytes. Multi CASI allowed for the detection of nearly 5-fold more peaks than that in the full scan analysis. While space charge effects are reduced, they are not eliminated. This is evident at 1200 laser shots in the Multi CASI experiment, where space charge effects are observed (Fig. 3a). Space charge in the ICR cell commonly manifests as peak shifting and peak coalescence.<sup>60</sup> As the laser shot count increases, the peaks for aspartate ( $m/z$  132.030) and eicosadienoate ( $m/z$  307.264) shift



**Fig. 2** MALDI mass spectra of rat brain extract using (a, b and c) a full scan acquisition method and (d, e and f) a Multi CASI acquisition method ( $m/z$   $131 \pm 5$  Da and  $293 \pm 40$  Da) designed to eliminate signal from the DAN MALDI matrix. Spectra shown in (b and c) are magnified  $m/z$  regions from the full scan acquisition and spectra shown in (e and f) are magnified  $m/z$  regions from the Multi CASI acquisition. Full scan spectra utilized 1000 laser shots per spectrum and Multi CASI spectra utilized 1000 laser shots per isolation window.



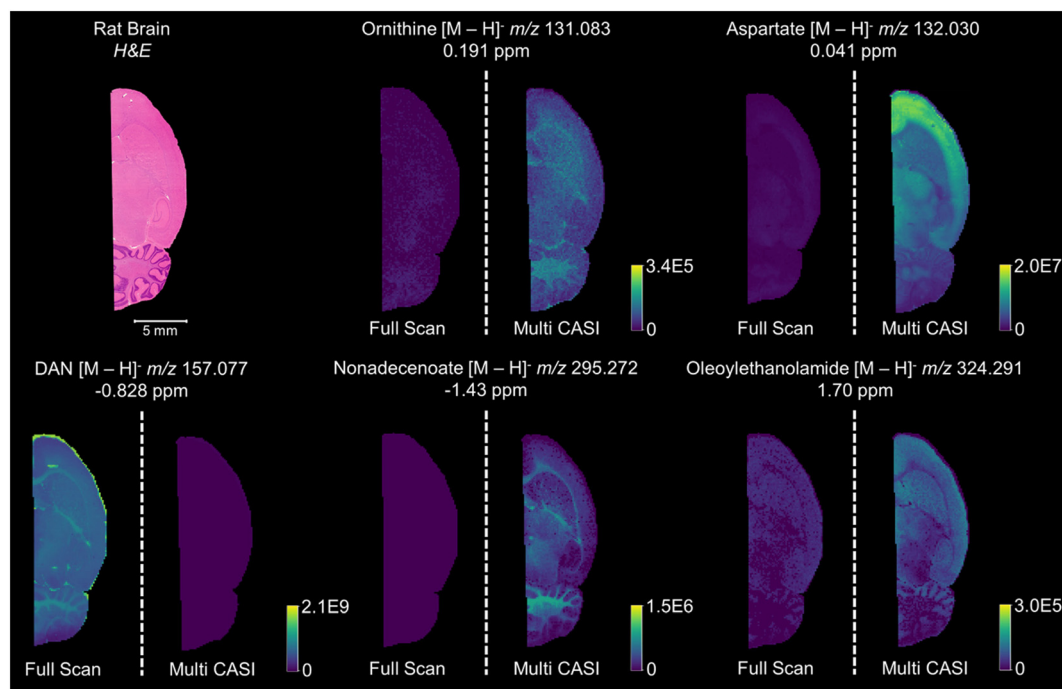
**Fig. 3** Average intensity of (a) aspartate ( $m/z$  132.030, 0.041 ppm) and (b) eicosadienoate ( $m/z$  307.264, 0.586 ppm) as a function of the number of MALDI laser shots per window for a full scan acquisition method (blue diamonds), a single CASI acquisition method [(a)  $m/z$   $131 \pm 5$  Da, (b)  $293 \pm 40$  Da, green triangles], and a Multi CASI acquisition method ( $m/z$   $131 \pm 5$  Da and  $293 \pm 40$  Da, grey squares). Each measurement represents three, five scan average measurements.

to higher  $m/z$  values (ESI Fig. 1†).<sup>60</sup> Ion cyclotron frequency decreases with increasing ion population in the ICR cell, which results in a shift to higher  $m/z$  values.<sup>61,62</sup> Space charge affects ions in the  $m/z$   $131 \pm 5$  Da window at a lower number of laser shots when compared to ions in the  $m/z$   $293 \pm 40$  Da

window (*i.e.*, 1200 versus 2000 laser shots in Fig. 3), and does so independent of the order of the two isolation windows (data not shown). This disparity may be due to a low-mass cutoff (LMCO) or pseudopotential trapping well depth effect in the hexapole ion trap.

Another method for limit of detection and dynamic range enhancement in MALDI mass spectrometry is single CASI.<sup>19</sup> The signal enrichment capabilities of Single CASI and Multi CASI were compared to quantify the improvements in dynamic range (Fig. 3). The two methods demonstrate comparable improvements in signal enrichment until ICR the charge capacity limit is reached in the Multi CASI experiment around 2000 laser shots (ESI Fig. 2†). Space charge effects will generally appear with fewer numbers of laser shots in the Multi CASI methods, as multiple isolation windows are included in the analysis, though this will depend on the width of the mass windows as well as the number and abundances of peaks in those windows. Increasing the quantity and width of isolation windows allows for more ions to be added to the ion trap and therefore reach the charge capacity limit with a fewer number of laser shots (ESI Fig. 3†). Even with this downside, Multi CASI still is able to detect more analytes as it can cover multiple mass ranges. Multi CASI allowed for the detection of 1.7-fold more peaks than single CASI analysis. Subsequent Multi CASI experiments are all performed using 1000 laser shots per window to minimize the effects of space charge.

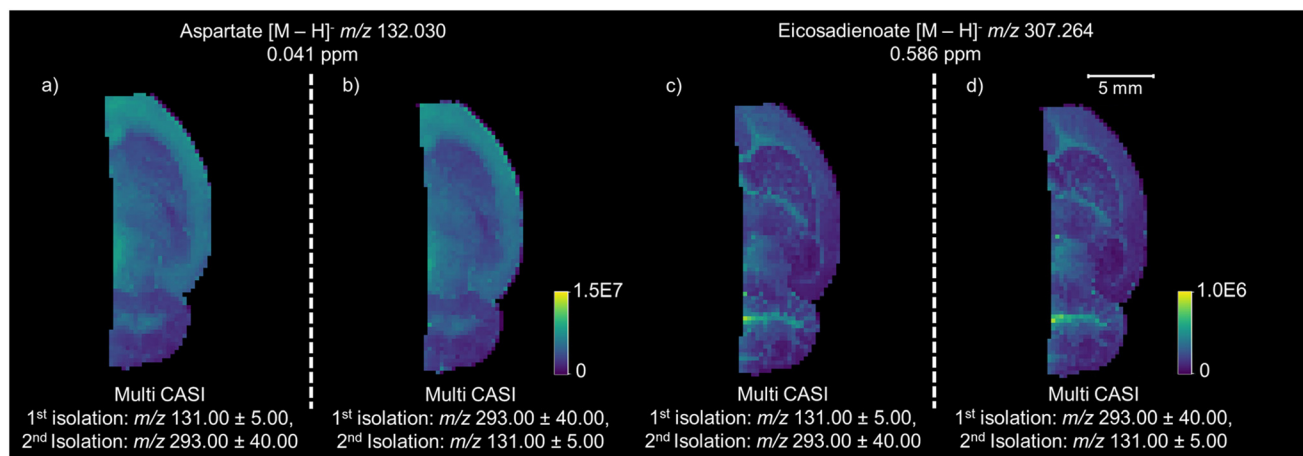
Multi CASI can be used to improve the limits of detection and dynamic ranges for lowly abundant compounds in imaging mass spectrometry experiments. Multi CASI windows centered at  $m/z$   $131 \pm 5$  Da and  $m/z$   $293 \pm 40$  Da are used to acquire an image of rat brain tissue at 150  $\mu$ m spatial resolution (Fig. 4). Full scan and Multi CASI utilized 1000 laser shots per window for both methods for comparison. Many ions that were not detected or detected at low abundance in the full scan experiment are readily detected across the rat brain tissue



**Fig. 4** Imaging mass spectrometry analysis (150  $\mu\text{m}$  spatial resolution, 1000 laser shots per window) of rat brain tissue using (left) a full-scan acquisition and (right) a Multi CASI ( $m/z$  131  $\pm$  5 Da and 293  $\pm$  40 Da) acquisition. Ion images for a range of metabolites and lipids show improved brightness (*i.e.*, limit of detection) and contrast (*i.e.*, dynamic range) with the Multi CASI acquisition. Images are displayed without normalization.

in the Multi CASI experiment. An increase in LOD is observed as an increase in the brightness of the image and an increase in dynamic range is observed as an increase in the contrast of the image. For example, a 6.5-fold increase in aspartate ion intensity was observed and 4.5-fold more peaks were detected in Multi CASI analysis when compared to full scan analysis, and DAN matrix intensity decreased by over 50-fold in Multi CASI analysis. Multi CASI is performed by ablating the same

pixel in multiple (*i.e.*, two) discrete MALDI ablation steps. Each MALDI ablation step then uses separate quadrupole mass filter settings to accumulate ions from disparate mass isolation windows. As such, MALDI laser power and shot count need to be optimized to ensure sufficient material is sampled in each Multi CASI ablation event and that signal is not diminished in the subsequent isolation windows. Laser power was optimized for each experiment to yield maximum signal in the



**Fig. 5** Imaging mass spectrometry analysis (300  $\mu\text{m}$  spatial resolution, 1000 laser shots per window) of rat brain tissue using two Multi CASI acquisition methods for detection of (a and b) aspartate using different sequences of mass isolation windows  $m/z$  131  $\pm$  5 Da and 239  $\pm$  40 Da and (c and d) eicosadienoate using different sequences of mass isolation windows  $m/z$  131  $\pm$  5 Da and 239  $\pm$  40 Da. Images are displayed without normalization.

first ablation event, while not diminishing analyte signal in the second ablation event. Two Multi CASI images were acquired using different sequential orders of the two isolation windows to verify that the sampling and accumulation conditions used here had no effect on the imaging mass spectrometry results (Fig. 5). For example, the tissue pixel is ablated and this first packet of ions is isolated with a mass isolation window of  $m/z$   $131 \pm 5$  Da. The same pixel is then ablated a second time and this packet of ions is isolated with a mass isolation window of  $m/z$   $293 \pm 40$  Da. Both ion populations are accumulated in the hexapole ion trap and then mass analyzed together in the ICR cell (Fig. 5a and c). In a second image, the mass isolation windows are then switched so that the first packet of ions is isolated using a mass window of  $m/z$   $293 \pm 40$  Da, followed by a second mass window of  $m/z$   $131 \pm 5$  Da (Fig. 5b and d). The distributions and intensity for aspartate and eicosadienoate are consistent across the tissue when comparing the two methods, indicating that the order of the ablation events does not have an impact on the quality of the imaging data using the settings describe here.

## Conclusions

Herein, we have described a method for gas-phase ion enrichment in imaging mass spectrometry experiments. Multiple continuous accumulations of selected ions (Multi CASI) utilizes multiple sequential RF/DC mass isolation windows in a quadrupole mass filter (QMF) to selectively accumulate ions in a downstream hexapole ion trap. In this study, Multi CASI can provide for nearly 500-fold enrichment in targeted ion signals, resulting in substantial improvements in the limits of detection and dynamic ranges of selected ions within the mass isolation windows. The ion enrichment provided by Multi CASI improves the MALDI analyses of biological tissue. Signal for lowly abundant compounds, such as the amino acids and lipids observed in our analyses, is greatly enriched. Multi CASI methods applied to several other workflows in our lab have also resulted in similar improvements in performance. Multi CASI improves upon typical CASI methods by allowing for the selection of multiple mass windows for analysis, enabling a broader view of compounds in a single image. This has a multiplexing effect compared to other CASI and selected ion monitoring (SIM) configurations. Multi CASI is also enabled without the addition of a supplemental waveform to the quadrupole, as has been reported in workflows that leverage multiple frequency notches. Accurate use of Multi CASI requires sufficient sample at each pixel for multiple laser ablation steps (*i.e.*, for multiple sequential mass isolation windows) on the same spot. Space charge effects can persist at higher laser shot counts due to abundant endogenous molecules, though these effects are observable at far fewer number of laser shots in full scan experiments. Herein, Multi CASI allowed for the detection of 4.6-fold more peaks in imaging mass spectrometry analysis of rat brain when compared to full scan analysis, and 1.7-fold more peaks than single CASI analysis. This leads to an increase

in throughput, reducing the number of targeted windows needed to enrich lowly abundant analytes in imaging mass spectrometry experiments.

## Abbreviations

|            |                                                    |
|------------|----------------------------------------------------|
| MALDI      | Matrix-assisted laser desorption ionization        |
| Multi CASI | Multiple continuous accumulations of selected ions |
| LC-MS      | Liquid chromatography-mass spectrometry            |
| DAN        | 1,5-Diaminonaphthalene                             |

## Author contributions

Troy R. Scoggins IV – data collection, data analysis, experimental design, and manuscript authorship. Jonathan T. Specker – training of primary author, experimental design, and manuscript review. Boone M. Prentice – data analysis, experimental design, manuscript review, and funding acquisition.

## Conflicts of interest

The authors declare no competing financial interest.

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