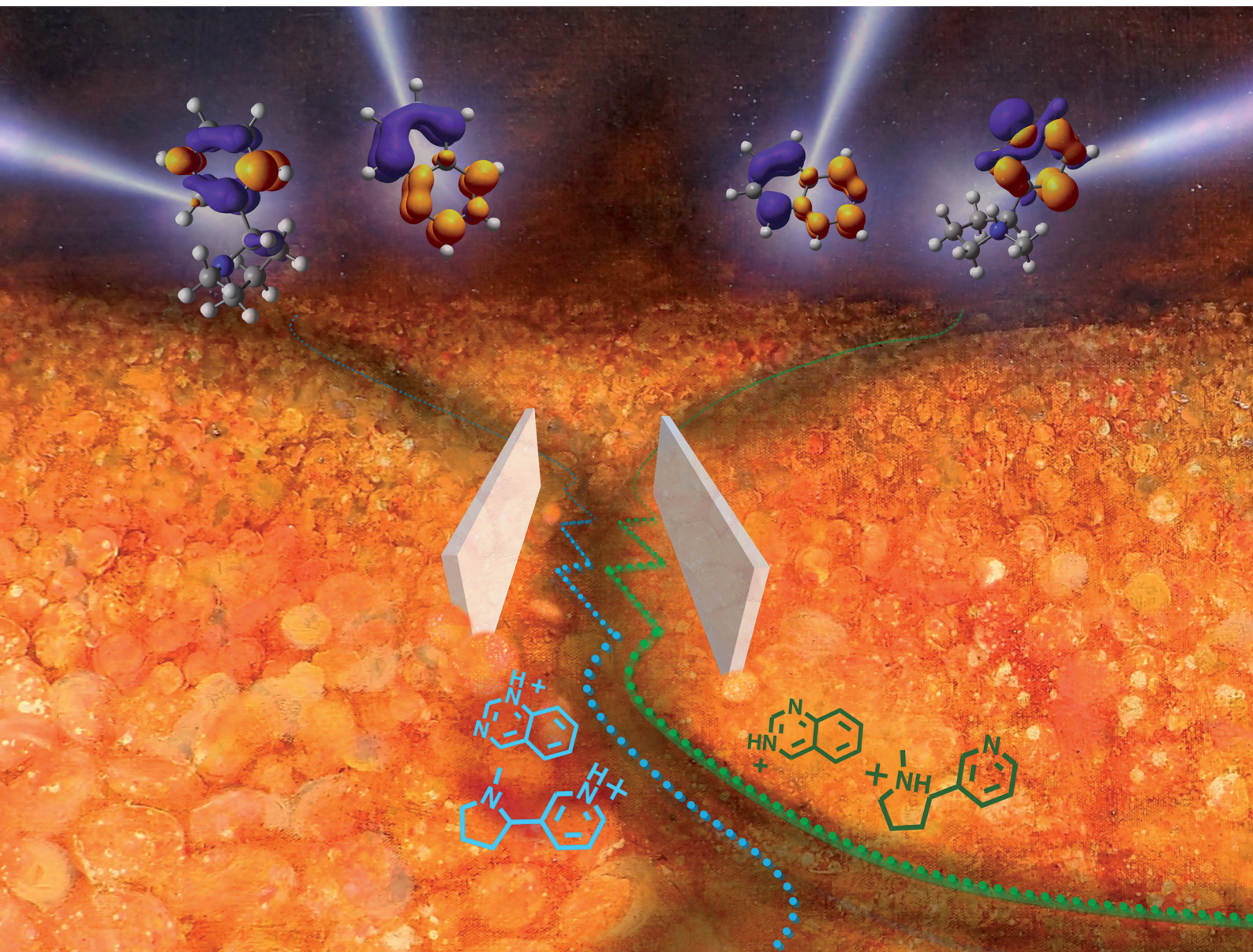


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**FEATURE ARTICLE**

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The combination of laser photodissociation,  
action spectroscopy, and mass spectrometry  
to identify and separate isomers



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# The combination of laser photodissociation, action spectroscopy, and mass spectrometry to identify and separate isomers

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The separation and detection of isomers remains a challenge for many areas of mass spectrometry. This article highlights laser photodissociation and ion mobility strategies that have been recently deployed to meet this challenge with focus on small molecule isomers including protonation isomers, structural isomers, conformation isomers and new studies emerging on chiral isomers. Laser techniques span UV and visible laser photodissociation, time-resolved pump–probe schemes and application of laser hole-burning arrangements to assign isomers within selected ion populations. Also surveyed are applications of ion mobility strategies to separate isomers followed by laser spectroscopic techniques to assign the separated ions. Ultimately, with ongoing refinement in hardware and methods, there are clear pathways forward for laser and mass spectrometry techniques to make decisive breakthroughs in understanding how isomeric details affect biological processes, physiology and disease.

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## Introduction

In mass spectrometry the inability to separate and assign isomers prevents the full extraction of chemical information and this impacts many research areas including metabolomics,<sup>1</sup>

lipidomics,<sup>2</sup> and mass spectrometry imaging.<sup>3</sup> Since the basis of mass spectrometry is the measurement of the mass to charge ratio ( $m/z$ ) of ions, the separation and identification of isomers requires additional strategies that need to be deployed in tandem. Such strategies include ion-mobility,<sup>4</sup> collision-induced dissociation,<sup>5,6</sup> ion-molecule reactions (e.g. ozone-induced dissociation),<sup>7,8</sup> and laser photodissociation techniques.<sup>9,10</sup> Over the last few decades, enormous advances in the conformational biomolecule analysis have occurred leveraging on advances in

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soft-ionisation techniques (e.g. electrospray ionisation (ESI)) together with numerous ion-mobility and computational strategies. Nevertheless, separating and assigning structural isomers for small molecules remains a significant challenge for mass spectrometry and this Feature Article focusses on recent advances in laser photodissociation (PD) mass spectrometry strategies in this context.

In the 1970s, some of the first reports emerged which described UV and visible wavelength photodissociation of mass selected ions as a technique to distinguish isomers. Classic examples include the assignment of various isomers of  $C_7H_8^+$  (radical cations of toluene, cycloheptatriene, and norbornadiene) on the basis of distinct PD spectra.<sup>11</sup> At the time, lasers generally lacked broadband UV-Vis tunability so these pioneering action spectroscopy experiments deployed arc-lamp/monochromator light sources with ion cyclotron resonance ion trap mass spectrometry.<sup>11–13</sup> Studies by the Dunbar and Beauchamp groups are landmark in this area and remain well-cited over 50 years later.<sup>11,12,14</sup>

Since these pioneering experiments, there have been perhaps two major technical advancements that have enhanced the scope and application of what is often referred to as PD action spectroscopy: (i) ESI/MALDI ionisation methods and (ii) tuneable UV-VIS mid-band OPO laser sources. The impact of ESI/MALDI cannot be easily overstated as these techniques have proliferated across the biological, chemical and physical sciences because they permit the study of an enormous variety of gas phase ions. Across all these fields, there are distinct pre and post ESI/MALDI invention epochs.<sup>15–17</sup> The second major advancement is the availability of the solid-state, tuneable, mid-band OPO laser. These are moderately priced (for a tabletop laser system) and are becoming increasingly turn-key and (mostly) hands-free operation. The OPO laser systems used in this context are typically pulsed, with repetition rates commonly (but not limited to) 10–100 Hz and pulse-widths commonly *ca.* 5–10 ns but also picosecond pulse-widths widely available too. The technical advantages of these OPO laser systems are the broadband tunability (spanning hundreds of nanometres) with good pulse energies (mJ), where scanning from the NIR through the visible and into the UV is relatively straightforward. High-energy wavelength cut-offs are typically around 210–225 nm, depending on the system. However, various vendors offer a range of options including deeper UV extensions, higher repetition rates (*ca.* 1 kHz), high pulse energies (*ca.* Joules), IR wavelength extensions, compact footprints, “rugged” housings, and more. There are some limitations and downsides to the common OPO laser systems. For example, OPO laser systems have pulse energies that vary significantly over large scanning ranges and the beam output can be rather divergent with irregular beam spot energy distribution. These factors can be problematic when power normalisation is required for scans over large wavelength ranges and laser beam overlap with ions can vary. Nevertheless, OPO laser systems are robust light sources for acquiring UV-Vis PD action spectra of gas-phase ions.

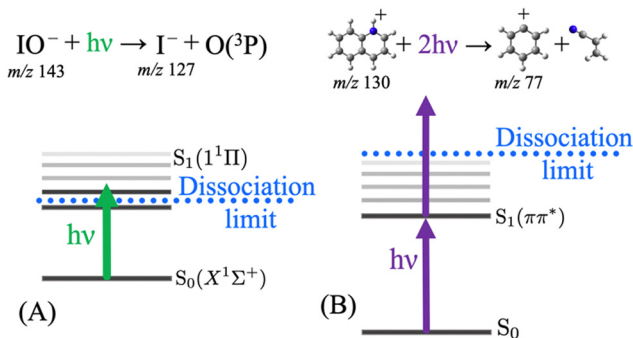
A major portion of the current UV-vis PD action spectroscopy research is conducted with ion traps, both at room temperature

and cryogenically cooled. Ion-mobility strategies such as drift tube ion mobility spectrometry (IMS) and field asymmetric ion mobility spectrometry (FAIMS) are also deployed for pre-filtering of ions. In this Feature Article, the advantages of these techniques are discussed. In defining the scope of this perspective, IR PD techniques—such as infrared multiphoton photodissociation (IRMPD)<sup>18–25</sup> and infrared photodissociation (IRPD)<sup>26–30</sup>—and *m/z* selected photoelectron (PE) spectroscopy<sup>31–34</sup> are not included in detail, with the exception of select examples. The interested reader is directed to these aforementioned references. Also, this Feature Article focusses on studies where ESI is the ionisation source. Other methods that generate ions for action spectroscopy include electron impact ionisation<sup>35–37</sup> and supersonic expansion followed by photoionisation<sup>38</sup> but these areas are not covered. Finally, therefore, this Feature Article is not an extensive review of all PD spectroscopy strategies. For extended resources the reader is directed to the following reviews on ion-mobility,<sup>4,39–41</sup> the history of ion spectroscopy,<sup>11,13,42</sup> advances and developments in ion spectroscopy,<sup>43</sup> UV-vis photodissociation action spectroscopy,<sup>44</sup> lasers and mass spectrometry for structure determination,<sup>45</sup> conformer resolved photoinduced dynamics for neutral and ionic peptides,<sup>46</sup> action spectroscopy of DNA radical cations,<sup>47</sup> gas-phase dynamics including isomer selective spectroscopy,<sup>48</sup> biological relevant ions with room temperature<sup>49–51</sup> and cryogenic ion spectroscopy.<sup>52,53</sup>

#### How does UV-vis PD action spectroscopy work?

Typically, UV-vis PD action spectroscopy is a method that measures the yield of photoproduct ions as a function of photon energy arising from the irradiation of an *m/z*-selected precursor ion population. Although details can vary, a typical procedure is as follows. A target analyte is introduced from solution *via* ESI, which serves to transfer the analyte from the liquid solution into the gas phase and (if required) to ionise the analyte. Ionisation often arises by protonation (cation) or deprotonation (anion). Charge can also be provided by non-covalent attachment of cations (e.g.  $Li^+$ ,  $Na^+$ ,  $K^+$ , and  $NH_4^+$ ) or anions (e.g.  $Cl^-$ ,  $I^-$ , and  $CH_3COO^-$ ). Alternatively, target ions may be a native cation or anion. At some point following this liberation of the analyte into the gas phase, *m/z* isolation of the target precursor ions is performed by a quadrupole ion guide or within an ion trap. This is a vital step that removes all other ions and provides an essentially background-free baseline in preparation for the detection of photoproduct ions. The target ions are then irradiated with a laser pulse (or many pulses). Following this, a mass spectrum is recorded and the intensity of any product ions arising from the laser pulse is measured against the total ion count. This sequence may be repeated depending on the desired signal-to-noise ratio. The laser wavelength is then altered, at the desired step size, and the process is repeated. At each laser wavelength, the PD yield is recorded as either the sum of all the photoproduct ions or a single *m/z* photoproduct ion spectrum can be plotted as a subset. The success of photodissociation action spectra depends on whether the photon is absorbed (the photon energy corresponds to an electronic transition) and secondly that the

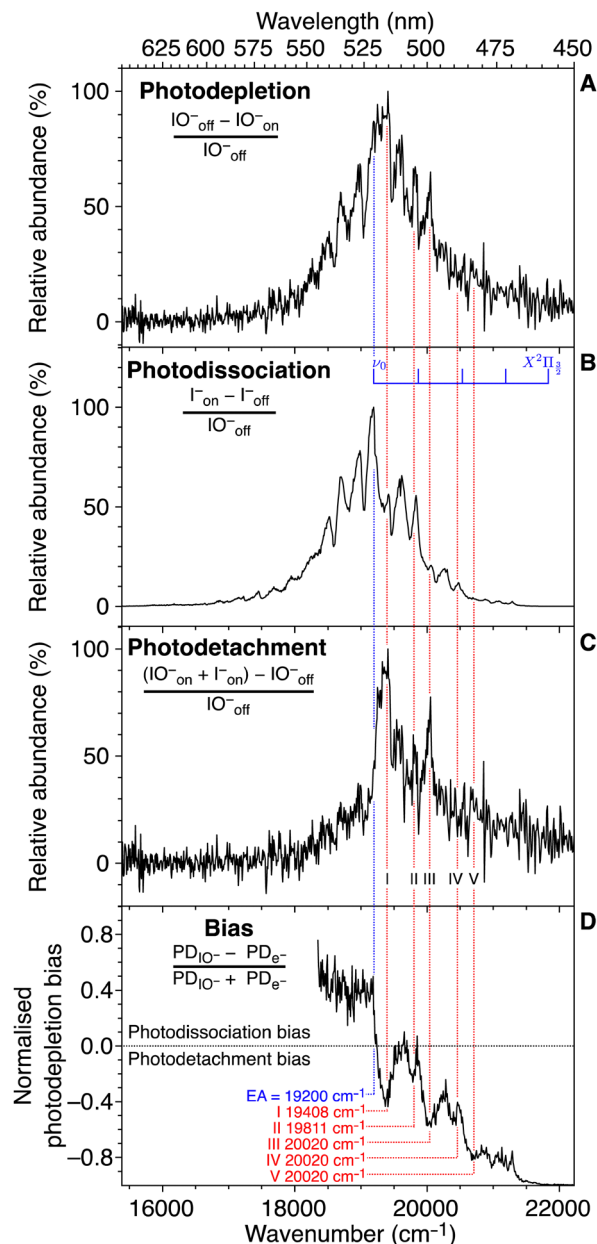




**Fig. 1** Photodissociation action spectra can be recorded resulting from 1-photon dissociation, for example as depicted for (A)  $\text{IO}^-$  excited into the lowest singlet excited state by one photon (which can be tuned to record the action spectrum). Alternatively, resonant 2-photon mediated photodissociation action spectra are possible as is the case for (B) protonated quinoline excited into the lowest singlet excited state by one photon and the subsequent absorption of a second photon is required to overcome the dissociation limit to form products – tuning the photon provides an action spectrum of the  $S_1$  state in this case despite it residing below the photodissociation limit.

photon energy is above the dissociation limit to form product ions (or above any rate limiting barriers that might impede dissociation). Fig. 1 shows two examples where action spectra have been reported using 1-photon mediated photodissociation action spectroscopy and a case where a 2-photon strategy can probe the  $S_1$  state, which is below the photodissociation limit, and a second photon is required to generate photoproducts. In the latter case, the two-photon process occurs within the same nanosecond laser pulse and the multiphoton process is confirmed by tracking the power dependence, as explained in ref. 54. Individual product ions may have different PD action spectra, which might signify different photodissociation processes for a precursor ion or that there are different populations within the precursor ion population (*e.g.* isomers). Interpretation of the PD yield should also take into any account background ions (that are not laser dependent) by background (laser on–laser off) subtraction and also importantly any variations in laser output should be taken into account by laser energy normalisation.

The action spectrum can be of different forms. For example, our recent paper on the action spectroscopy of  $\text{IO}^-$  demonstrated how several processes can be investigated following laser irradiation (Fig. 2).<sup>55</sup> In this example,  $\text{IO}^-$  is  $m/z$  selected and isolated using a room temperature linear ion-trap mass spectrometer and then irradiated. Any loss of  $\text{IO}^-$  parent ions can be recorded with a laser-on vs. laser-off procedure—this is a photodepletion spectrum. Another photoactivated process is bond dissociation, where  $\text{IO}^-$  dissociates into  $\text{I}^- + \text{O}(^3\text{P})$  and thus a photodissociation spectrum is generated by plotting the yield of  $\text{I}^-$  ( $m/z$  127) normalised to the laser-off signal. While no product ions would normally be expected in the laser-off spectrum, the isolation step in an ion trap mass spectrometry can activate fragile ions and give rise to a background of product ions. Since this example is an anion, another laser activated process is photodetachment of electrons, which is



**Fig. 2** Various signal pathways in the action spectroscopy of  $\text{IO}^-$ : photodepletion of the  $\text{IO}^-$  parent ion signal (A); the photodissociation yield of  $\text{I}^-$  (B); the electron photodetachment yield from  $\text{IO}^-$  (C); and the bias between photodissociation and photodetachment (D). The superior S/N of the photodissociation spectrum (B) compared to the other spectra is notable. All channels are monitored concomitantly in the ion trap experiment. Reprinted with permission from B. I. McKinnon, S. J. Marlton, B. Ucur, E. J. Bieske, B. L. Poad, S. J. Blanksby and A. J. Trevitt, *J. Phys. Chem. Lett.*, 2021, **12**, 11939–11944. Copyright 2021 American Chemical Society.<sup>55</sup>

plotted by monitoring the drop in the overall ion count normalised to the laser-off signal. In this example, the competition between photodissociation and photodetachment is plotted out in a bias plot and both are competitive across this photon range. The photodissociation action spectrum, Fig. 2(B), has a superior signal-to-noise ratio, since the  $\text{I}^-$  signal is produced on an essentially zero level of background noise. On comparing



this to the depletion signals, Fig. 2 panels A and C, these are inherently affected by more noise since they monitor the attenuation of the parent ion signal. The final note on this series of spectra is that they are all acquired simultaneously, and the photodepletion spectrum is deconvoluted in the photodissociation and photodetachment spectra.

For the acquisition of PD action spectra, various experimental arrangements have been deployed, including adaptations to commercial ion traps,<sup>56–60</sup> custom built ion-traps,<sup>61</sup> commercial triple-quadrupole MS arrangements,<sup>62</sup> cryogenic ion traps,<sup>53,63–70</sup> and other designs that involve ion mobility selection<sup>71–74</sup> or ion-storage rings.<sup>75,76</sup> All these arrangements allow laser access to selected ions to induce photodissociation. Some of these setups include ion mobility stages, which will be discussed in detail later.

As demonstrated in the above example, PD action spectroscopy is an indirect spectroscopic method. For ions, direct gas-phase absorption measurements are generally challenging since gas-phase ions are typically available in a relatively low number density compared to that of solution-phase or dense gas-phase systems. The number of ions that can be generated by ESI—and subsequently selected in the mass spectrometer—can range from tens of millions down to a few tens and this is much lower than the number density in solution phase absorption, which is normally well above  $10^{12}$  molecules  $\text{cm}^{-3}$  (nM). For a gas-phase ion packet if, say, ten photons are absorbed by the sample from a laser pulse made up of trillions of photons, then the attenuation of photon intensity would be difficult to measure. Alternatively, since PD action spectroscopy relies on the detection of photoproduct ions at different  $m/z$  values from the selected precursor ions it is essentially a background-free measurement and dissociation events are easily recorded even if only a few product ions are generated for each MS cycle.

Another inherent advantage of PD action spectroscopy using mass spectroscopy is that it introduces an element of target selection of ions by their  $m/z$  ratio. As such, background signal and contamination are often eliminated. Of course, this  $m/z$  selection alone does not allow for the selection of isomers or isobaric contaminants. The separation and identification of isomers are an active area of research development and a strategy for addressing isomer separation is using ion-mobility techniques to separate isomers as will be discussed later. Another notable method for separation is chromatography, which has been coupled to IRMPD spectroscopy.<sup>22,77,78</sup> Although ion mobility techniques are a powerful addition for isomer separation and assignment, spectroscopic techniques alone can provide a detailed characterisation of certain isomers.

### Isomer ions

There are many reported studies using UV/vis PD action spectroscopy that interrogate isomeric ions that have been selected before introduction into the mass spectrometer. In these cases, pure samples of structural isomers were available so—provided no isomerisation occurs upon delivery of ions—the spectra from these isomers can be compared. Such studies have targeted substituted pyridines,<sup>79–82</sup> azaindoles,<sup>83,84</sup> diazabenzenes,<sup>85,86</sup> indazole and benzimidazole,<sup>87</sup>

and diazanaphthalenes,<sup>82</sup> as well as anionic systems: deprotonated azaindoles,<sup>84</sup> nitrophenolate ions,<sup>88–90</sup> molecular-iodide clusters,<sup>91</sup> and protonated quinoline and isoquinoline.<sup>54,92</sup>

The case of the structural isomers protonated quinoline and isoquinoline is rather straightforward. The PD action spectra for the  $S_1 \leftarrow S_0$  transitions of quinolineH<sup>+</sup> and isoquinolineH<sup>+</sup> are shown in Fig. 3. These spectra are constructed by tracking the three major product ions, which are the same for both precursor ions.

The pure electronic transition ( $\nu' = 0 \leftarrow \nu'' = 0$ ), labelled 0–0, is often the key value for assigning experimental spectra because it is a clearly defined value in both calculations and experiment.<sup>93,94</sup> The difference in the 0–0 energy of protonated isoquinolineH<sup>+</sup> and quinolineH<sup>+</sup> is *ca.* 200  $\text{cm}^{-1}$ . Note that a hot-band was incorrectly assigned as the 0–0 for isoquinolineH<sup>+</sup> in the original publication.<sup>54</sup> The 0–0 energies of the two isomers were subsequently measured by Jouvét and co-workers at cryogenic temperatures to be within 200  $\text{cm}^{-1}$ .<sup>92</sup> The difficulty in characterising small differences between 0–0 energies is compounded by the fact that excited state calculations tend to be less accurate and less robust than ground state calculations. Time dependent density functional theory (TD-DFT) calculations have typical absolute uncertainties in the order of 2000  $\text{cm}^{-1}$  and the coupled cluster method CC2, (which is the gold standard method for excited state calculations of medium sized molecules) has typical uncertainties of

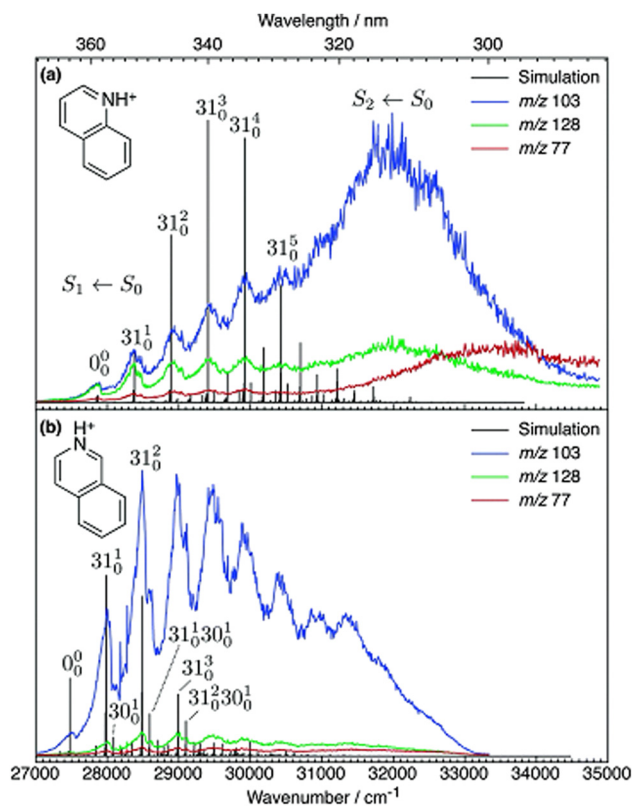


Fig. 3 UV PD action spectra for (a) quinolineH<sup>+</sup> and (b) isoquinolineH<sup>+</sup> ions. Reproduced from ref. 54 with permission from the PCCP Owner Societies.<sup>54</sup> These spectra are acquired in a linear ion trap (LTQ).



*ca.* 1000 cm<sup>-1</sup>. Methods like EOM-CCSDT have typical errors of above 200 cm<sup>-1</sup> and are too expensive to be routinely used for molecules of this size and larger.<sup>95</sup> The fact that EOM-CCSDT and other high level methods—which aspire to reach chemical accuracy (<1 kcal mol<sup>-1</sup> or 350 cm<sup>-1</sup>)—emphasises the importance of benchmark experimental data to validate these claims. In these aforementioned isomers studies, the isomers were generated from chemical standards where the identity of the isomer was known. But what if both isomers were present in one sample? To meet this challenge, Coughlan *et al.* used the quinolineH<sup>+</sup> and isoquinolineH<sup>+</sup> reference spectra to assign isoquinolineH<sup>+</sup> and quinolineH<sup>+</sup> co-sprayed into the instrument and separated within the mass spectrometer using differential ion mobility spectrometry (DMS).<sup>96</sup> Therefore, these isomer resolved spectra can serve as references for future diagnostic applications.

A common structural variable that gives rise to isomeric ions in mass spectrometry is variation in the protonation site. Such isomers are termed protomers. Protomers present in the solution may be transferred to the gas phase following ESI, along with protomers that are favoured in the gas phase.<sup>21,97–100</sup> However, the relative protomer populations generated under ESI are affected by the ESI conditions and do not necessarily reflect the statistically preferred distributions in either the gas phase or the solution phase.<sup>97–100</sup> Because protomer populations are affected by ESI (and because the exact mechanisms by which ions are formed in ESI are not fully understood), it can be difficult to predict which protomer is going to dominate in the experiment. This uncertainty means that the protomers present need to be assigned and characterised.

### Room temperature UVPD

Targeting protomers, Matthews and Dessent reported the room temperature UVPD action spectra of protomer isomers co-populated in an ion trap in their study of para-aminobenzoic acid (PABA).<sup>9</sup> The room temperature action spectra could be deconvoluted to two protomers of PABA, as shown in Fig. 4. Fig. 4a shows the spectrum of the O-protonated form and Fig. 4b shows the spectrum of the N-protonated form. Because there are major differences in the spectra of these protomer ions, and the major product ions are characteristic to the protomer precursor ions, the isomers are clearly distinguished. Since these product ions are—for the most part—unique to the precursor protomer ion, any isomerization between protomers after photoactivation must be minimal. However, depending on the structure of the molecule, the availability of proton scrambling pathways, and the difference in the protonatable functional groups, protomer ions can give rise to the same photoproduct ions. In this case, the protomers of PABA have different photoproduct ions as well as different action spectra.

Photodissociation product ions can provide insight into assignment of the protomer precursor ion. For example, Fig. 4b follows both the generation of the H atom (–1 Da) and the NH<sub>3</sub> fragment (–17 Da), previously assigned to the N protomer.<sup>101,102</sup> Therefore, based on the photoproducts, one can begin to assign the spectra to different protomers before analysing each spectrum.

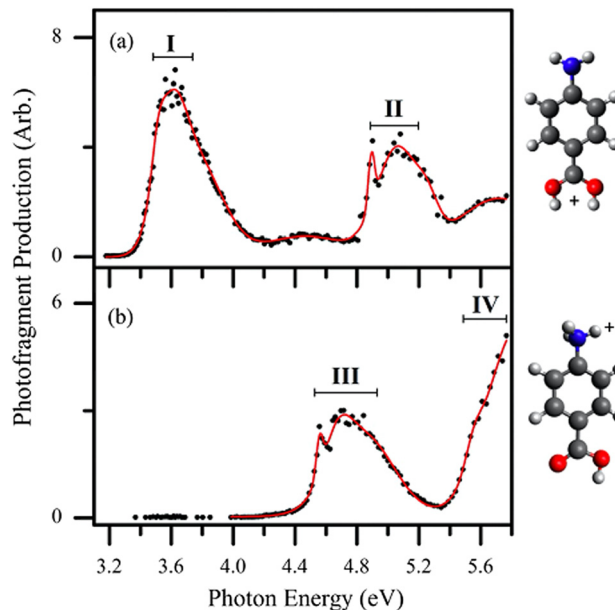


Fig. 4 UV PD action spectra for para-aminobenzoic acid (PABA) ions in different solvents following different PD products. The spectra correspond to two protomers co-populated in the ion trap, (a) the O-protomer and the (b) N-protomer, which are distinguishable by their different action spectra and PD products. Reproduced from ref. 9 with permission from the PCCP Owner Societies.<sup>9</sup>

For direct assignment of isomers, photodissociation action spectroscopy can be compared to calculated spectroscopic transitions. For example, the first electronic transition for the O-protonated form of PABA is calculated to be centred around 3.58 eV (using the high level MRCI method) and the peak in the experimental spectrum (Fig. 4a) is centred around 3.51 eV. Similarly, the calculated first transition of N protonated PABA is calculated to be 4.50 eV, which is close to the experimental transition around 4.56 eV (Fig. 4b). This agreement between calculations and experimental spectra—in addition to other observed states—provides an unequivocal assignment of these spectra to the O and N protomers, which are co-located in the ion-trap.

The Dessent group has also deployed PD action spectroscopy to study protomers of flavin ions<sup>103</sup> and nicotinamide.<sup>104</sup> In these experiments, the various isomers are co-located in the ion trap. Other UV PD studies on nucleobases<sup>105,106</sup> distinguished between multiple protomers. These studies demonstrate how the electronic transitions and non-radiative decay are affected by the protonation site, which is noteworthy considering protonation bears some resemblance to the hydrogen bonding and isomerism that occur for nucleobases in DNA. From the PD action spectra of these molecules, their intrinsic photoproperties can be studied. These properties include absorption transitions and non-radiative decay mechanisms. This is benchmark information to understand the effect of more complex environments, including solvent shifts or other environmental factors. UV PD has also been employed to understand how structural changes affect the photochemistry of sunscreens, for example, by distinguishing and characterising how the binding site of a



cation can inhibit the excited-state proton transfer that affects the photostability of sunscreen agents.<sup>107–109</sup>

Nucleobase radical cation isomers,<sup>47,110–113</sup> as well as several other biologically relevant radical cation isomers, have been analysed by Turaček and co-workers using UV action spectroscopy in a modified commercial ion trap, which takes advantage of the inert ion trap environment. Reactive radical ions, which would otherwise be transient intermediates that quickly react away, can be isolated and stored for the acquisition of PD spectra.

The Turaček group has also compared the results from the two most common commercial mass spectrometers that are modified for PD action spectroscopy (THERMO LTQ and Bruker AmaZon). These modified setups are shown in Fig. 5a and b.<sup>56</sup> The main difference between these arrangements is that for the linear ion trap (THERMO LTQ) optical access is afforded through a window on the rear of the mass spectrometer, whereas optical access is afforded through a window on the top of the mass spectrometer to the 3D ion trap (Bruker AmaZon) with a clear path in and out of the instrument. The spectra in Fig. 2 and 3 were acquired using a THERMO LTQ mass spectrometer, whereas the spectra in Fig. 4, from Dessent and co-workers, were acquired using a Bruker AmaZon mass spectrometer. It is evident that both configurations are suited for PD action spectroscopy; however, these spectra should not be used to directly compare the performance of the two mass spectrometer setups because they also employed different lasers, span different photon energies, and report spectra from different ions. For direct comparisons between equivalent instruments using a THERMO LTQ and a Bruker AmaZon mass spectrometer see ref. 56.

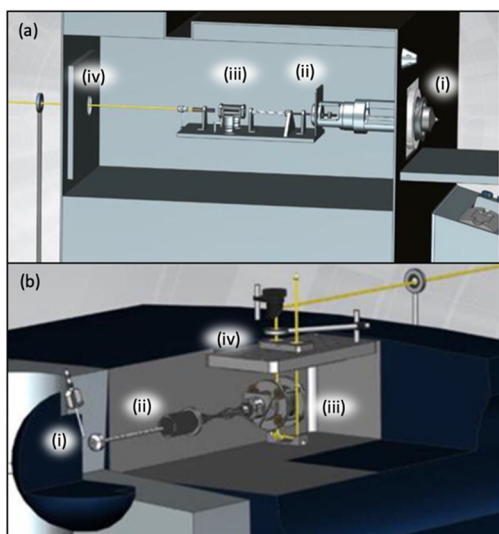


Fig. 5 A diagram of how a THERMO LTQ (a) or a Bruker AMAZON mass spectrometer (b) can be adapted to allow optical access to the ion trap is shown. Figure has been amended to include labels of key components indicating ESI ion source region (i), electrostatic ion guides (ii), ion trap (iii), and laser beam (yellow line) entering the mass spectrometer through a window (iv). Adapted from A. Dang, J. A. Korn, J. Gladden, B. Mozzone and F. Tureček, *Journal of The American Society for Mass Spectrometry*, 2019, **30**, 1558–1564 Copyright 2019 American Chemical Society (ref. 56).

Although there are many cases where room temperature PD action spectroscopy can distinguish isomers, there will be many instances where that the room temperature spectra of isomers are inherently too broad and/or congested. Cooling of ions in cryogenically-cooled ion traps can simplify spectra.

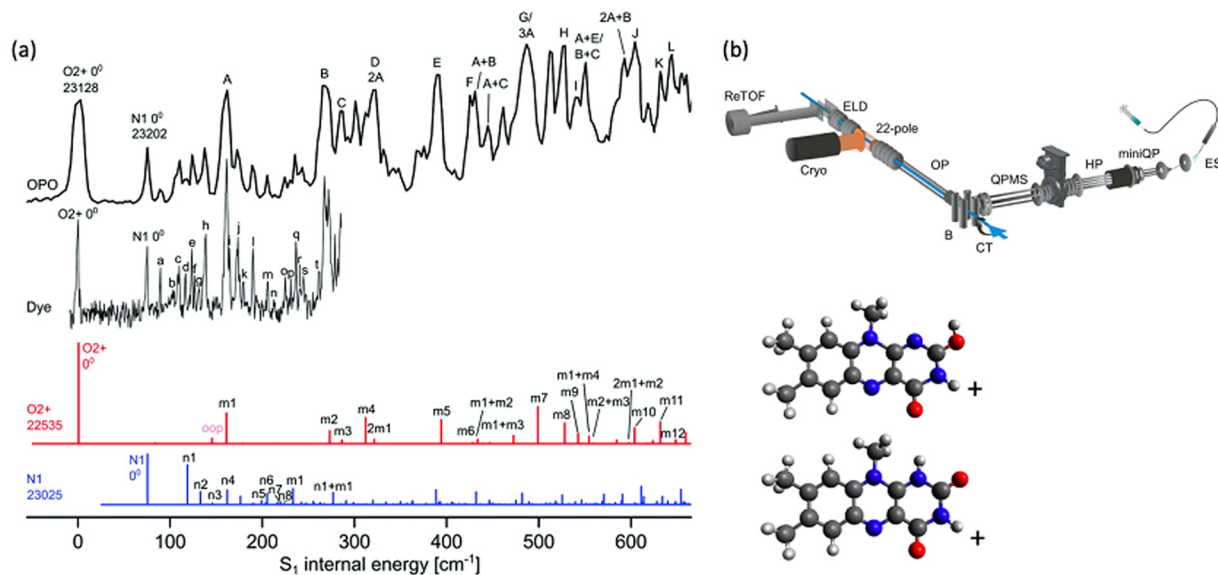
### Cryogenic UVPD

As is well established, vibronic spectra are generally sharper and better resolved when ions are cooled in cryogenic ion traps as this greatly restricts vibrational state populations and narrows rotational state distributions. PD action spectroscopy of ions confined in cryogenically cooled environments is conducted in a similar manner to the previously described room temperature experiments except that the temperature of the ions is cooled to *ca.* 3–50 K using a buffer gas, which is typically helium gas. Fig. 6b shows a schematic of a setup employing a cryogenic ion trap as shown in ref. 63.<sup>63</sup> The ions are generated by ESI and guided into a 22-pole ion trap, which is cooled by a He cryostat. With colder ions, hot band intensities and rotational broadening are significantly reduced or eliminated and the sharpness of the spectra can be vastly improved.

Cryogenic ion-trap PD action spectroscopy has been used to characterise the protonation or metalation-site isomers for ions of flavins,<sup>114–116</sup> nucleobases,<sup>117,118</sup> polycyclic aromatic amines,<sup>119,120</sup> distinguishing between different spin states of high-spin ions,<sup>121,122</sup> and to disentangle *E* and *Z* double-bond isomers of hemithioindigo<sup>123</sup> and the green fluorescent protein chromophore,<sup>124</sup> lipid isomers<sup>125</sup> and carbohydrates.<sup>126</sup> Similar experiments have measured electron photodetachment (rather than PD) as a function of wavelength for deprotonomers of cytosine anions.<sup>31</sup>

The PD action spectra of protonated lumiflavin (LFH<sup>+</sup>) are shown in Fig. 6 (from the study by Dopfer and co-workers).<sup>114</sup> A lower resolution spectrum is recorded using a mid-band OPO laser (linewidth *ca.* 4 cm<sup>-1</sup>) and the high-resolution spectrum is acquired using a dye laser (linewidth *ca.* 0.014 cm<sup>-1</sup>). The first major peak (normalised at 0 cm<sup>-1</sup>) is assigned as the origin of the O protomer. Aided by the simulations, it is apparent that two co-located protomers contribute to the experimental spectrum. The N-protomer 0–0 is assigned at +74 cm<sup>-1</sup> from the O-protomer 0–0. Several low frequency excited state vibrational quanta are assigned within a few 100 cm<sup>-1</sup> above the 0–0 transition due to both protomers. This vibronic structure allows for the assignment of two protomers based on comparison with FC simulations and calculated 0–0 transition energies. The O2<sup>+</sup> protomer (red simulation) is planar in the excited state and is dominated by in-plane vibrations and the geometry of the N1 protomer (blue simulation) is non-planar in the excited state and has a spectrum that has several active low-frequency out of plane vibrations. This illustrates how biological flavins can have altered photoproperties as a consequence of different protonation sites, which then highlights the importance of the flavin environment to biological function. Furthermore, the flavin properties can also be affected by metal cation binding, as other studies using cryogenic PD action spectroscopy have observed that changing the metal binding site can





**Fig. 6** (a) Cryogenic spectra of LFH<sup>+</sup> (black) acquired with a mid-band OPO and a sharper-line-width dye laser. Franck–Condon simulations for two protomers are red and blue. Both protomers are assigned as contributing to the experimental spectrum. Reproduced from ref. 114 with permission from the PCCP Owner Societies.<sup>114</sup> (b) A schematic illustration is included to provide a survey of the cryogenic BerlinTrap apparatus used to acquire these spectra. Ions are generated using ESI, accumulated in a miniature quadrupole (miniQP), transferred into a hexapole (HP), *m/z* selected using a quadrupole mass spectrometer (QPMS), reflected 90° by a quadrupole bend, guided by an octupole (OP) into a cryogenic 22 pole ion trap where they are cooled and can be irradiated by a laser pulse (blue line) after which the ions are detected using a reflectron time of flight mass spectrometer (ReTOF). (b) Reproduced with permission from A. Günther, P. Nieto, D. Müller, A. Sheldrick, D. Gerlich and O. Dopfer, A new cryogenic 22-pole ion trap spectrometer, *J. Mol. Spectrosc.*, 2017, **332**, 8–15. Copyright Elsevier, 2017.<sup>63</sup>

dramatically shift the 0–0 transition energy for the first  $1\pi\pi^*$  state of flavin ions.<sup>114–116</sup>

For protonated lumiflavin, vibronic spectra obtained under cryogenic conditions allow for convincing assignment of protomers. Assignment is achieved by comparison between experimental and calculated results, which typically involves Franck–Condon simulations and calculated 0–0 transition energies.<sup>114,115,119,120,127–130</sup> The interplay between experiment and computational chemistry methods is exemplary in this area of research. Accurate calculations are often needed to assign experimental spectra and, in turn, quantum chemical calculations need experimental data for benchmarking and validation.

Natural linewidth-limited peak shapes recorded using cryogenic ion PD spectroscopy can be used to estimate excited-state lifetimes.<sup>83,85,118</sup> This is possible because rotational broadening of vibronic peaks should be minimised at low temperatures and generally spectra are much less congested. Using cryogenic ion PD spectroscopy, two protomers of protonated uracil were determined to have excited state lifetimes of 40 fs and 2 ns (where the 2 ns lifetime was measured using a pump–probe).<sup>85,118</sup> This difference between lifetimes was attributed to the different energy gaps between the  $\pi\pi^*$  and  $n\pi^*$  states. Thereby, the effect of lone pairs and protonation site on the photostability of DNA nucleobases was spectroscopically observed.

### Pump–probe photodissociation

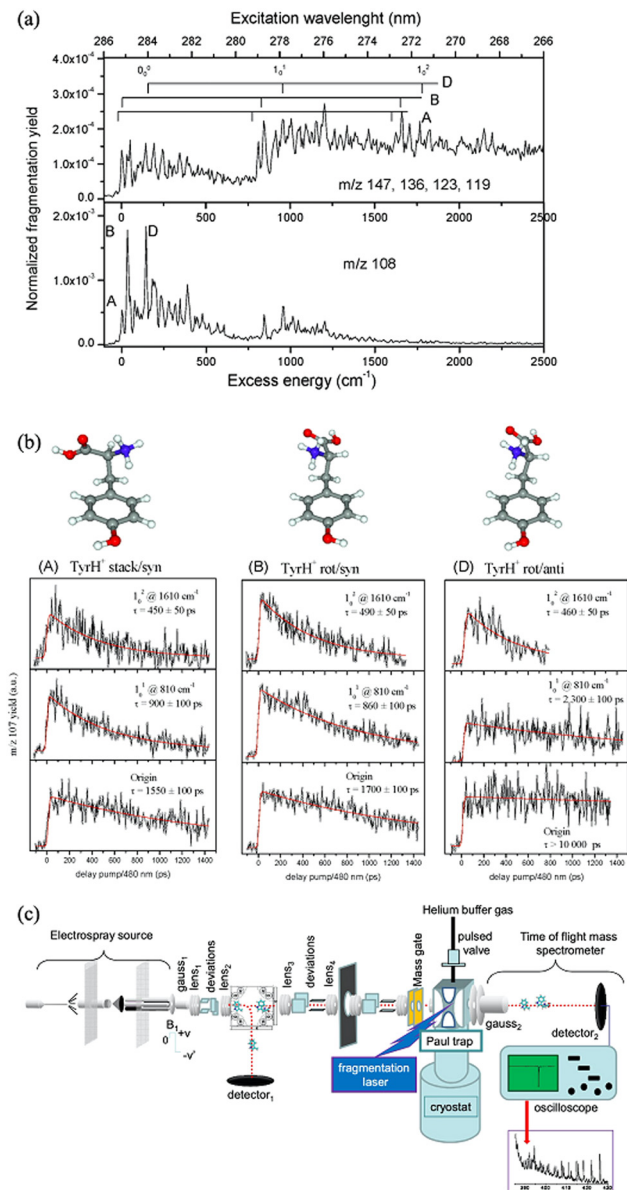
For co-located isomer ions, if vibronic transitions are sufficiently narrow and separated it is possible to tune a laser to selectively excite a vibronic transition of only one isomer

population, leaving the other unaffected. Taking advantage of this, Soorkia *et al.* selectively excited particular conformers of protonated tyrosine ions using resonant transitions. By then introducing a second (probe) laser pulse, they measured pump–probe PD spectra for various conformers (see Fig. 7).<sup>131</sup> A schematic of the experimental setup is also shown in Fig. 7c; all isomers are co-isolated in the ion-trap (Paul trap), which is cooled by a cryostat. Both the pump and probe pulses (see fragmentation laser in Fig. 7c) irradiate the ion cloud in the ion-trap.

This arrangement of the isomer-selective pump–probe experiment was employed to investigate protonated tyrosine isomers<sup>131</sup> and nucleobase isomers.<sup>117,132</sup> Protonated tyrosine conformers are found to have similar excited state lifetimes at 1610 cm<sup>-1</sup> above the 0–0 transition (*ca.* 450 ps). On the other hand, the  $1n\pi^*$  lifetimes of protonated cytosine varied by over an order of magnitude and, as previously mentioned, the excited state lifetimes of uracil conformers varied by almost five orders of magnitude.

Other groups have deployed femtosecond or picosecond pump–probe laser configurations combined with photodissociation mass-spectrometry to interrogate excited-state lifetimes of ions in tandem time-of-flight mass spectrometers,<sup>133–135</sup> triple-quadrupole mass spectrometers,<sup>136</sup> ion storage rings,<sup>137,138</sup> custom built ion traps at room temperature,<sup>139–141</sup> custom built ion traps that are cryogenically cooled,<sup>81,85,132,142</sup> and commercial ion traps.<sup>87,143–146</sup> There are also setups using electronic delays for pump–probe photodissociation on the nanosecond to millisecond timescale.<sup>117,132,146–151</sup> The excited





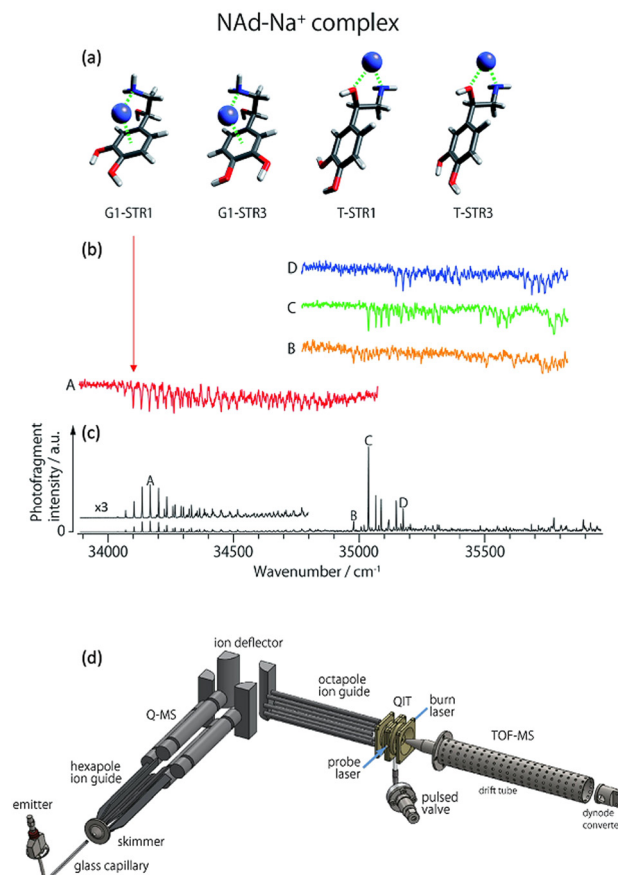
**Fig. 7** (a) Cryogenic UV PD action spectra of tyrosineH<sup>+</sup> ions. (b) pump-probe PD plots following the PD yield of tyrosineH<sup>+</sup> ions as a function of time delay between the pump and probe pulses (right). By setting the pump pulse to excite a vibronic peak for only one isomer, each isomer can be selected and its lifetime can be selectively probed. (a and b) Adapted with permission from S. Soorkia, M. Broquier and G. Grégoire, *J. Phys. Chem. Lett.*, 2014, **5**, 4349–4355. Copyright 2014 American Chemical Society.<sup>131</sup> (c) Schematic of the experimental setup. (c) Adapted with permission from I. Alata, J. Bert, M. Broquier, C. Dedonder, G. Féraud, G. Grégoire, S. Soorkia, E. Marceca and C. Juvet, *J. Phys. Chem. A*, 2013, **117**, 4420–4427. Copyright 2013 American Chemical Society.<sup>64</sup>

state dynamics of few gas-phase ions has been investigated experimentally or computationally, especially relative to solution-phase species.

### Laser hole burning techniques

A particular isomer population that are co-located with other isomers can be selectively excited by tuning a laser to a

characteristic vibronic transition and “burning” out a major fraction of that population. Hole burning spectroscopy takes advantage of this selectivity to obtain a spectrum of only one isomer or conformer. This requires two lasers. First, a single laser photodissociation spectrum would normally be recorded and contain peaks from all isomers/conformers co-located in an ion trap experiment. In the hole-burning experiment, a probe laser is fixed to one peak in the PD spectrum. Then a burn laser is scanned across the spectrum and this is pulsed a short time before (*ca.* a few milliseconds) the probe laser.



**Fig. 8** (a) Some low energy isomers of noradrenaline-Na<sup>+</sup>. (b) Cryogenic UV-PD hole burning spectra of noradrenaline-Na<sup>+</sup> isomers. (c) Cryogenic UV-PD spectrum exhibiting the spectrum of all isomers generated by ESI. Figures (a), (b) and (c) reproduced from ref. 152 with permission from the Royal Society of Chemistry.<sup>152</sup> By setting a “probe” laser pulse to selectively excite one isomer, the PD from that isomer is detected. By introducing a second laser, an isomer selective spectrum can be generated whereby the spectrum dips when the second laser “burns” out the population of the isomer being selected by the “probe”. (d) A schematic illustration is included to provide a survey of the experimental setup. Ions are generated using ESI, guided through a hexapole, *m/z* selected using a quadrupole mass spectrometer (Q-MS), reflected 90° by a quadrupole bender, guided by an octapole into a quadrupole ion trap (QIT) where they are cooled and can be irradiated by one or two laser pulses (blue lines) after which the ions are detected using a reflectron time of flight mass spectrometer (TOF-MS). (d) Reproduced with permission from S.-i. Ishiuchi, H. Wako, D. Kato and M. Fujii, High-cooling-efficiency cryogenic quadrupole ion trap and UV-UV hole burning spectroscopy of protonated tyrosine, *J. Mol. Spectrosc.*, 2017, **332**, 45–51 Copyright Elsevier, 2017.<sup>65</sup>



Variations in the probe laser PD signal are tracked as the burn laser is scanned. When the burn laser is resonant with a transition from an isomer that is also targeted by the probe laser, a depletion in the probe signal is recorded. For other transitions scanned over by the burn laser, the probe laser signal is unaffected. This process is repeated with a probe laser then shifted to a different peak (Fig. 8 includes several hole-burning spectra). Therefore, the hole-burning strategy provides an isomer-selective depletion spectrum.

Examples of hole burning spectra are presented in Fig. 8 where isomer-selective spectra for four noradrenaline- $\text{Na}^+$  conformers (red, orange, green and blue traces) are recorded from an experiment with all isomers co-located in the ion trap. The different spectra are acquired by scanning the burn laser and setting the probe laser to selectively excite one of the peaks A, B, C, or D shown in Fig. 8c. A schematic example of the instrumentation for a hole burning experiment is shown in Fig. 8d (reproduced from ref. 65), which incorporates ESI,  $m/z$  selection with a quadrupole mass spectrometer (labelled Q-MS). Ions are guided into the ion-trap where they are cryogenically cooled by pulsed helium gas and subsequently irradiated by the burn and probe laser pulses. This hole burning experiment was used to assign the vibronic progression labelled A (red trace) to a closed structure where  $\text{Na}^+$  interacts with the aromatic  $\pi$  system. This isomer was not observed for the noradrenaline- $\text{Li}^+$  ion (not shown), which may explain the different effect that  $\text{Li}^+$  and  $\text{Na}^+$  has on the inhibition of noradrenaline molecular recognition.<sup>152</sup>

Biomolecular ions of considerable structural flexibility like polypeptides and proteins have been analysed using combined

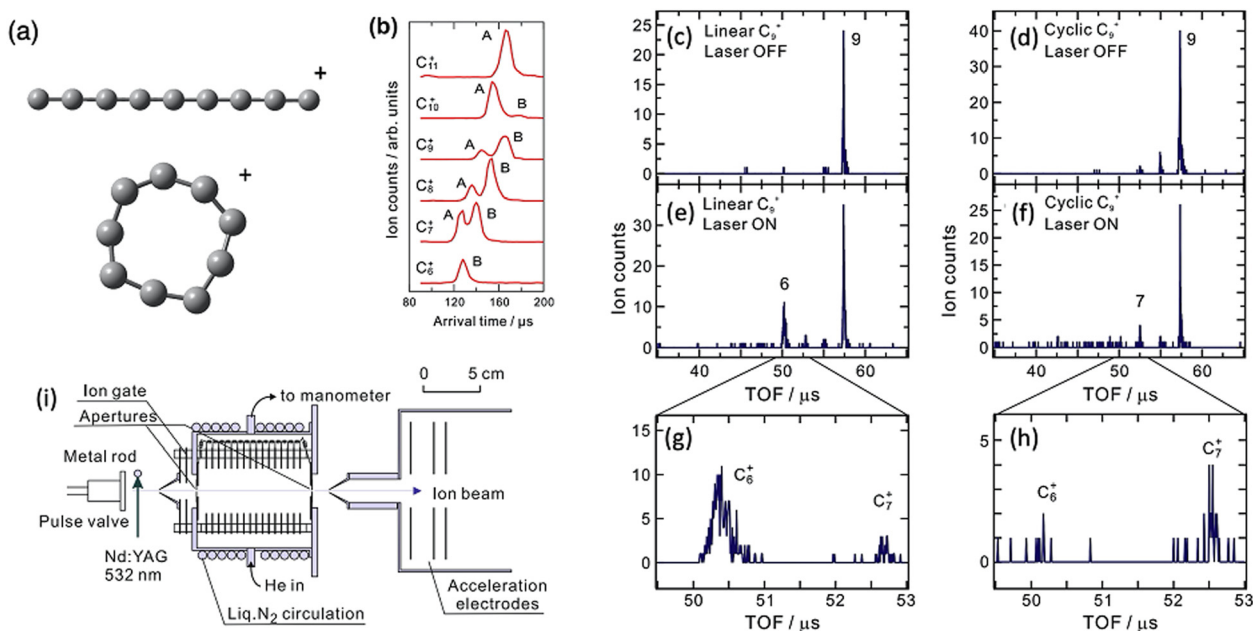
cryogenic and IR and UV hole burning and double resonance experiments.<sup>30,153–155</sup> For more on cold ion spectroscopy, the interested reader is directed to the following reviews for biological ions by Boyarkin and co-workers<sup>52,156</sup> and Soorkia *et al.*<sup>46</sup>

### Ion mobility spectrometry with photodissociation

Ion mobility spectrometry in its various forms is a method to separate ions by physical properties including size, dipole and buffer-gas affinity. By coupling IMS with PD spectroscopy, isomers can first be separated and then spectroscopically investigated without the complication of co-located isomers.

An important example of these techniques used together is from Koyasu *et al.* targeting isomers of  $\text{C}_9^+$  arising from laser vaporisation of graphite. Various  $\text{C}_9^+$  isomers were separated using drift tube ion mobility spectrometry and then irradiated with 355 nm photons to induce photodissociation (see Fig. 9). In IMS, ions drift through a cell that contains a gas, typically  $\text{N}_2$  or He. As the ions drift, their mobility is affected by the ion-molecule collisions as quantified by the collision cross section (CCS). Two types of  $\text{C}_n^+$  isomers are reported in this experiment, linear and cyclic. The linear isomers have a larger CCS and they tumble through the collision cell with longer arrival time delays (peak labelled A in Fig. 9b). The cyclic isomers are more compact with a smaller CCS and, therefore, have the arrival time (peaks labelled B in Fig. 9b).

Because the two  $\text{C}_9^+$  isomers are separated, the ratio between  $\text{C}_6^+$  and  $\text{C}_7^+$  formation is determined, where the linear isomer favours  $\text{C}_6^+$  and the cyclic isomer favours  $\text{C}_7^+$ . Because they are separated, the difference is obvious, which is a clear advantage of the combined ion mobility and PD mass spectrometry



**Fig. 9** Structures of linear and cyclic  $\text{C}_9^+$  (a), Drift IMS arrival times plots of  $\text{C}_n^+$  clusters (b), mass spectra of linear  $\text{C}_9^+$  with laser on (c) and laser off (e and g), mass spectrum of cyclic  $\text{C}_9^+$  with laser on (d) and laser off (f and h). Adapted with permission from K. Koyasu, T. Ohtaki, N. Hori and F. Misaizu, Isomer-resolved dissociation of small carbon cluster cations,  $\text{C}_7^+$ – $\text{C}_{10}^+$ , *Chem. Phys. Lett.*, 2012, **523**, 54–59 Copyright Elsevier 2012.<sup>157</sup> (i) Schematic of experimental setup reproduced from F. Misaizu, N. Hori, H. Tanaka, K. Komatsu, A. Furuya and K. Ohno, Isomer-selected photoreactions of gas-phase cluster ions, *Eur. Phys. J. D*, 2009, **52**, 59–62 with permission of Springer.<sup>74</sup>



arrangement. Action spectra are not acquired in this experiment. A subsequent experimental arrangement that combines IMS and cryogenic PD action spectroscopy has presented the electronic spectra of several cyclic carbon cluster ions.<sup>158,159</sup>

Some other examples of ion mobility coupled with photoelectron spectroscopy or PD using set wavelengths are other carbon cluster ions,<sup>160,161</sup> silicon cluster ions,<sup>161</sup> trisaccharides,<sup>162</sup> DNA anions,<sup>163,164</sup> proteins,<sup>165–169</sup> flavins,<sup>73</sup> and glycans.<sup>170</sup> Also, fixed wavelength arrangements have been used for more exotic experiments such as ion-mobility selected gas-phase fluorescence.<sup>171–173</sup>

### Ion mobility spectrometry coupled with spectroscopy

Ion-mobility and PD action spectroscopy deployed together can reveal considerable detail about ions and isomer populations. For example, protomers of benzocaineH<sup>+</sup> could be separated first using IMS and then assigned using IRMPD.<sup>100</sup> Fig. 10b shows that the mobility selected IRMPD spectrum of isomer II matches the calculated spectrum for the N-protomer of benzocaineH<sup>+</sup>, and the spectrum of I matches the calculated spectrum of the O-protomer. The IRMPD spectra exhibit several diagnostic peaks (like the N-protomer peak at 1250 cm<sup>-1</sup> or the O-protomer peak at 1550 cm<sup>-1</sup>). These peaks in the IRMPD spectra allow for unambiguous assignment of the two ion-mobility separated species. Notably, CID generates the same photoproducts for both protomers and, therefore, does not provide definitive assignment information. However, one could argue that these protomers could be assigned using IRMPD without separation—although the result would be perhaps less definitive without IMS separation. Even though the IMS points to additional isomers (isomers II and II') the IRMPD spectra of these isomers appear almost identical. The populations of these protomers and isomers can be measured by variation of the ESI solvent (Fig. 10a).

Ion mobility combined with room temperature IR PD has been used to distinguish isomers of saccharides,<sup>174</sup> amino acids,<sup>175</sup> amino acid clusters,<sup>176</sup> small molecules,<sup>177</sup> metal-oxide clusters,<sup>178</sup> and  $\beta$ -sheet formation.<sup>179</sup> Combined ion mobility and cryogenic IR PD studies have distinguished isomers of peptides<sup>180–182</sup> and glycans.<sup>183–188</sup> Gas-phase infrared spectroscopy of glycan isomers was also recently reviewed.<sup>189</sup> In the case of many systems mentioned here, especially some glycan isomers, the difference between isomers is very slight and these isomers cannot always be distinguished using one technique alone (*i.e.* just ion mobility or just IR PD). It is only when ion mobility is combined with IR PD that these isomers can be clearly distinguished.

Another ion-mobility separation method is field-asymmetric ion mobility mass spectrometry (FAIMS).<sup>96,98,190</sup> While traditional drift IMS separates ions by arrival time dictated by their cross-sectional mobility in a low-strength electric field, FAIMS filtered ions by mobility in both low and high-field strengths. The principles of FAIMS are described in detail elsewhere.<sup>191,192</sup> Briefly, ions are carried by an inert gas between two electrodes across which a dispersion voltage (DV) is applied, which is an asymmetric voltage that oscillates between high and low voltage of opposite polarity. Ions are separated based on their displacement in the high field and low field regimes and different

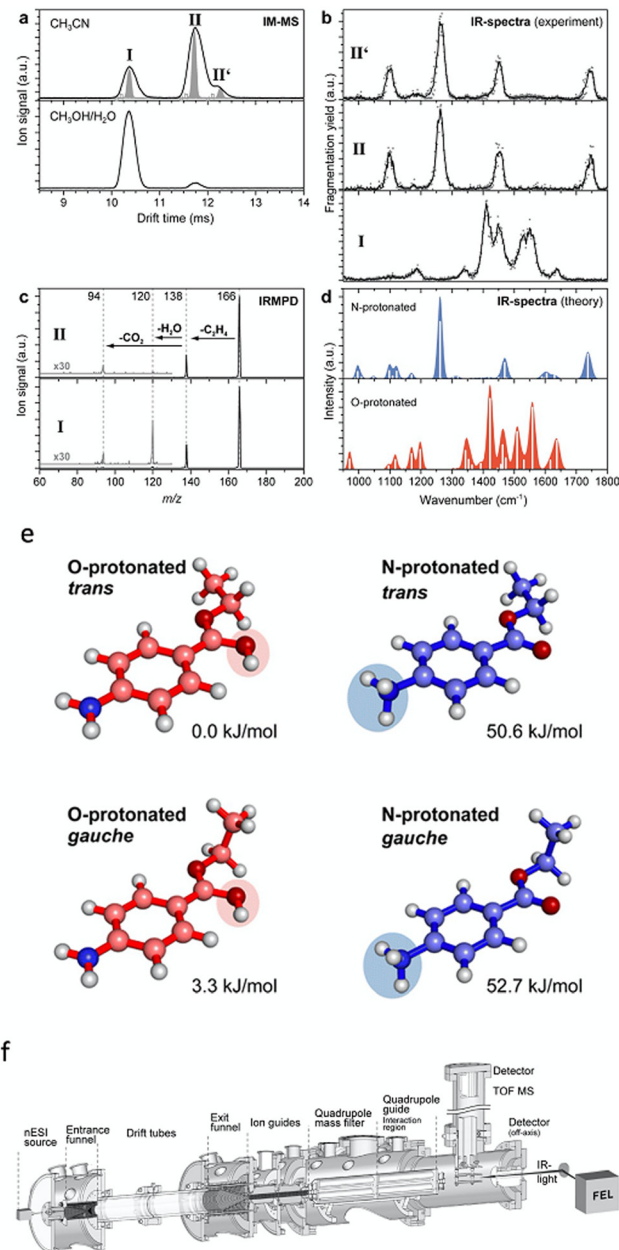
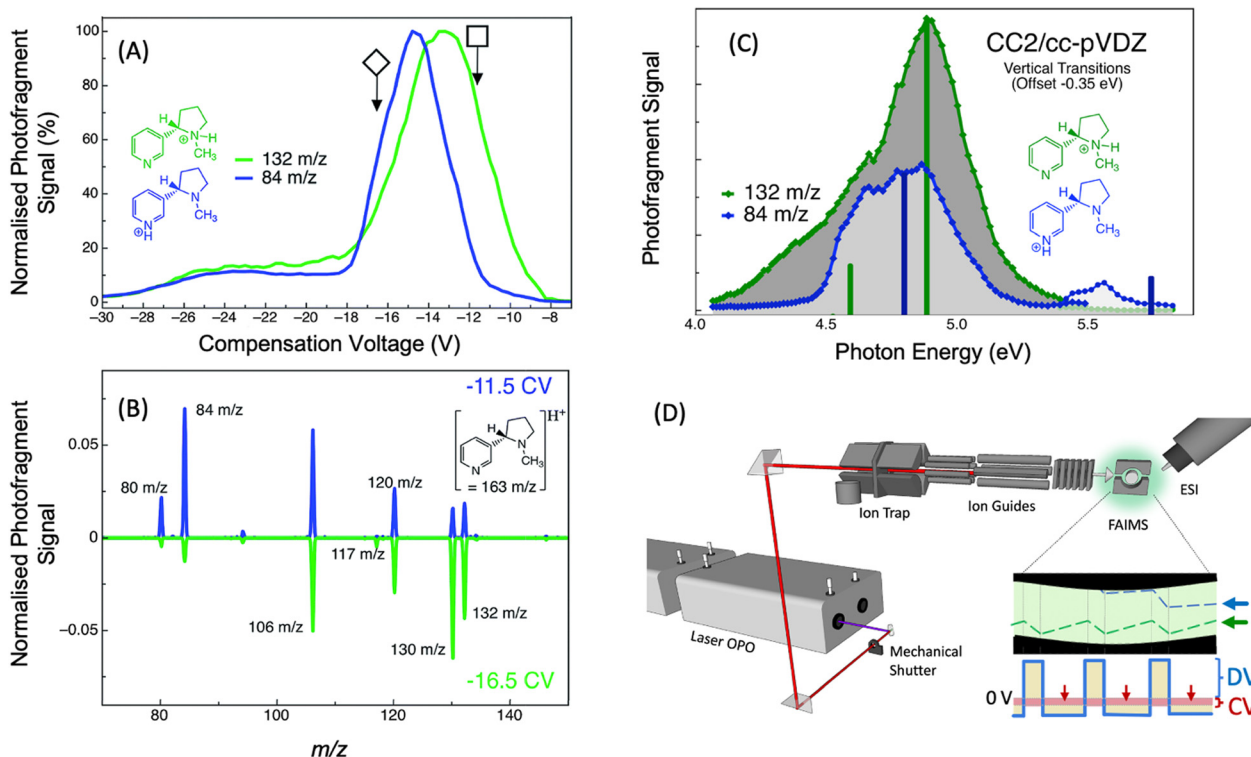


Fig. 10 (a) Drift IMS for benzocaineH<sup>+</sup> generated using acetonitrile or a mixture of methanol and water as the ESI solvent. (b) IRMPD spectra of the ion-mobility selected protomers and isomers of benzocaineH<sup>+</sup>. (c) IRMPD mass spectra. (d) Calculated IR spectra for different benzocaineH<sup>+</sup> protomers. Structures and relative ground state energies for different protomers and isomers of protonated benzocaine (e). Schematic of the experimental setup (f). Adapted from S. Warnke, J. Seo, J. Boschmans, F. Sobott, J. H. Scrivens, C. Bleiholder, M. T. Bowers, S. Gewinner, W. Schöllkopf and K. Pagel, *J. Am. Chem. Soc.*, 2015, **137**, 4236–4242. Original article is in the public domain.<sup>100</sup>

isomers will have different trajectories between the FAIMS electrodes. The trajectory of an isomer can be corrected by applying a compensation voltage (CV) so that only the selected isomer passes between FAIMS electrodes to be detected. The FAIMS voltages and their effect on the trajectory of an ion are shown schematically in Fig. 11D.





**Fig. 11** (A) FAIMS ionogram for nicotineH<sup>+</sup> ( $m/z$  163) following two PD products of nicotineH<sup>+</sup> ( $m/z$  132 and  $m/z$  84), which are indicative of the two nicotine-H<sup>+</sup> protomers. (B) PD mass spectra ( $\lambda = 266$  nm) of nicotineH<sup>+</sup> with FAIMS compensation voltage set to  $-11.5$  V (blue line) or  $-16.5$  V (green line). (C) Photodissociation action spectra of nicotineH<sup>+</sup> following the signal of the  $m/z$  84 photoproduct ion (blue trace) and the  $m/z$  132 photoproduct ion (green trace). Vertical bars show vertical transition energies calculated using CC2 for the pyridine protonated protomer (blue bars) and the pyrrolidine protonated protomer (green bars). Vertical transition energies are offset by  $-0.35$  eV for comparison to experimental spectra. (D) Schematic illustration to provide a survey of the experimental arrangement at University of Wollongong. Ions are generated using ESI, the isomer can then be separated and selected using FAIMS, guided into the linear ion trap (LIT) where they are  $m/z$  selected and stored before being irradiated by a laser pulse (red line) and scanned out of the ion trap to produce a mass spectrum. A schematic showing the DV (solid blue line) and CV (red shaded area) used in FAIMS and the trajectory that two different isomers might take (blue and green dashed lines) between the FAIMS electrodes. Reproduced from ref. 98 with permission from the Royal Society of Chemistry.<sup>98</sup>

Drift IMS and FAIMS each have advantages and disadvantages. While FAIMS can provide separation different from that of drift IMS,<sup>193</sup> the FAIMS separation process is known to thermally activate ions<sup>194</sup> and possibly cause structural rearrangements—this would be of particular concern for native biomolecular ion studies but less of a concern for small molecule investigations targeted here. Drift IMS provides collisional cross-section (CCS) values for separated species. These CCS values provide information on the ion shape and can be compared with other reference CCS values and calculated values. Therefore, even without PD spectroscopy, the CCS values can provide information on structural assignments of different isomers separated using IMS. A method to measure CCS values using FAIMS results is emerging,<sup>195</sup> which is promising but not as established as CCS measurements based on drift IMS.

It is sometimes the case that protomer ions are not possible to disentangle without separation. In particular, broad featureless spectra can be difficult to deconvolve when there are multiple species with similar spectra. For example, in our study of nicotineH<sup>+</sup>, a combination of FAIMS and UV PD action spectroscopy (see Fig. 11D) was able to distinguish the

overlapping spectra of the two nicotineH<sup>+</sup> protomers.<sup>98</sup> The UV PD action spectra of nicotineH<sup>+</sup> (without FAIMS separation) are shown in Fig. 11C following the formation of the  $m/z$  84 photoproduct (blue trace) and the  $m/z$  132 photoproduct (green trace). The spectra obtained by following these two photoproducts are sufficiently different, with the green trace exhibiting a shoulder at a lower energy ( $<4.5$  eV) and the blue trace exhibiting a high energy feature at 5.5 eV.

By comparing these experimental action spectra to vertical transition energies and relative intensities calculated with the CC2 method (vertical bars in Fig. 11C), it is found that there is good agreement between the calculated electronic transitions of the pyrrolidine protonated nicotineH<sup>+</sup> and the spectrum obtained by following formation of the  $m/z$  132 photoproduct and there is good agreement between the calculated electronic transitions of the pyridine protonated nicotineH<sup>+</sup> and the spectrum obtained by following formation of the  $m/z$  84 photoproduct. This is consistent with previous studies of the dissociation of nicotineH<sup>+</sup> assigned to the  $m/z$  132 product ion as forms from the pyrrolidine protomer of nicotineH<sup>+</sup>, while the  $m/z$  84 product ion forms from the pyridine protomer of nicotineH<sup>+</sup>.<sup>196</sup> Based on CC2 calculations, it was determined



that the shoulder at  $<4.5$  eV in the spectrum of the pyrrolidine protomer arises from an  $n\pi^*$  state, which involves the n-orbital from the pyridine-ring nitrogen's lone pair—this state is not present for the pyridine protomer because the pyridine lone pair is involved in the bonding to  $H^+$ . Therefore, when compared to vertical electronic transitions, the UV PD action spectra provide direct spectroscopic evidence for the presence of two protomers of nicotine $H^+$  and direct spectroscopic assignment of these two protomers.

A room temperature IRMPD spectrum of nicotine $H^+$  has been reported,<sup>197</sup> which concluded that just the pyridine-protonated protomer was present. This could signify that only one protomer was generated under those conditions of ESI, or perhaps both protomers were present and the spectrum of one was hidden under the spectrum of the other. Cryogenic infrared photodissociation spectra of  $H_2$  tagged nicotine $H^+$  ions provide additional spectroscopic evidence for the presence of both nicotine $H^+$  protomers.<sup>198</sup>

Another example of FAIMS separation of protomer ions is the case of quinazoline $H^+$  where two protomers present from ESI in comparable yields.<sup>190</sup> With FAIMS separation, the spectra of both protomers were separately recorded, with a vibronic structure (Fig. 12). The presence of this vibronic structure allows for compelling assignment of the isomer structures and this happens with Franck–Condon simulations and comparison to calculated 0–0 transition energies. In this case, the simulations nicely reproduce the experimental vibronic structure and the 0–0 energies for the  $S_1 \leftarrow S_0$  transitions are measured as  $26\,900\text{ cm}^{-1}$

(calculated as  $26\,650\text{ cm}^{-1}$  using MS-CASPT2) for 3-quinazoline $H^+$  and  $27\,600\text{ cm}^{-1}$  (calculated as  $28\,424\text{ cm}^{-1}$  using MS-CASPT2) for 1-quinazoline $H^+$ . However, like the quinoline $H^+$  and isoquinoline $H^+$  spectra shown in Fig. 3, the 0–0 transition energies of quinazoline $H^+$  protomers are close together ( $<1000\text{ cm}^{-1}$ ) and so require the use of an accurate method (in this case MS-CASPT2) to confidently make the assignment.

Other examples of experiments that combine ion mobility with PD action spectroscopy are FAIMS separation with cryogenic UV/vis PD spectroscopy of peptides from Rizzo's group,<sup>180,199,200</sup> FAIMS separation with room temperature UV/vis PD spectroscopy from the Hopkins group,<sup>96,201–204</sup> drift mobility separation with cryogenic spectroscopy from Bieske's group,<sup>158,159</sup> and drift mobility separation with room temperature photodissociation and photoisomerization from Bieske's group<sup>72,205–218</sup> and Dugourd's group.<sup>219,220</sup>

### Tandem IMS and photoactivation

Tandem IMS has been combined with photoactivation to observe photoisomerism. This is possible because photoisomerism can alter the ion CCS. The Bieske group<sup>72,205–218</sup> and the Dugourd group<sup>219,220</sup> have both deployed this strategy. The tandem ion mobility setup uses an IMS-IMS-QMF configuration with two drift ion mobility regions (IMS1 and IMS2). A schematic of this experimental setup is shown in Fig. 13, along with an ion mobility spectrum and photoisomerization action spectrum of protonated *trans* azobenzene reproduced from ref. 207. The *trans* isomer of protonated azobenzene is

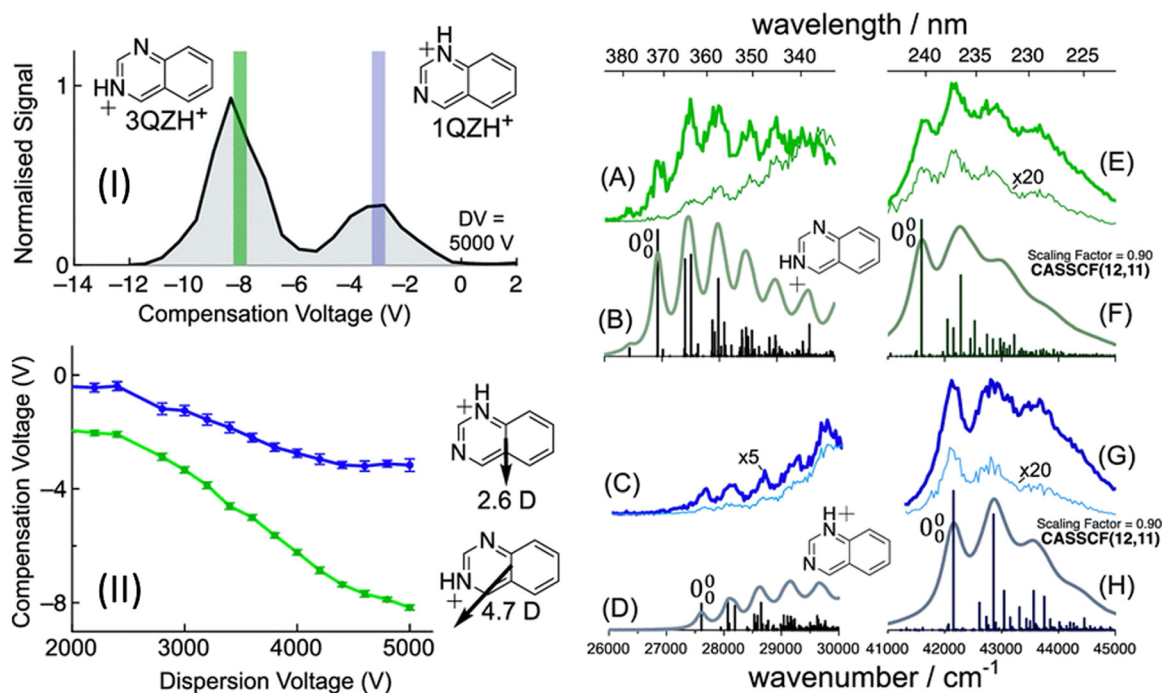


Fig. 12 (I) FAIMS ionogram following quinazoline- $H^+$  ( $m/z$  131). Action spectra of quinazoline- $H^+$  using compensation voltages of  $-8$  V (A and E) or  $-3$  V (C and G). Franck–Condon simulations for the  $S_1$  ( $L_a$ ) state of 3-quinazoline- $H^+$  (B) and 1-quinazoline- $H^+$  (D). Franck–Condon simulations for the  $S_5$  ( $B_b$ ) state of 3-quinazoline- $H^+$  (F) and 1-quinazoline- $H^+$  (H). S. J. Marlton, B. I. McKinnon, B. Ucur, J. P. Bezzina, S. J. Blanksby and A. J. Trevitt, *J. Phys. Chem. Lett.*, 2020, **11**, 4226–4231. Copyright 2020 American Chemical Society.<sup>190</sup>



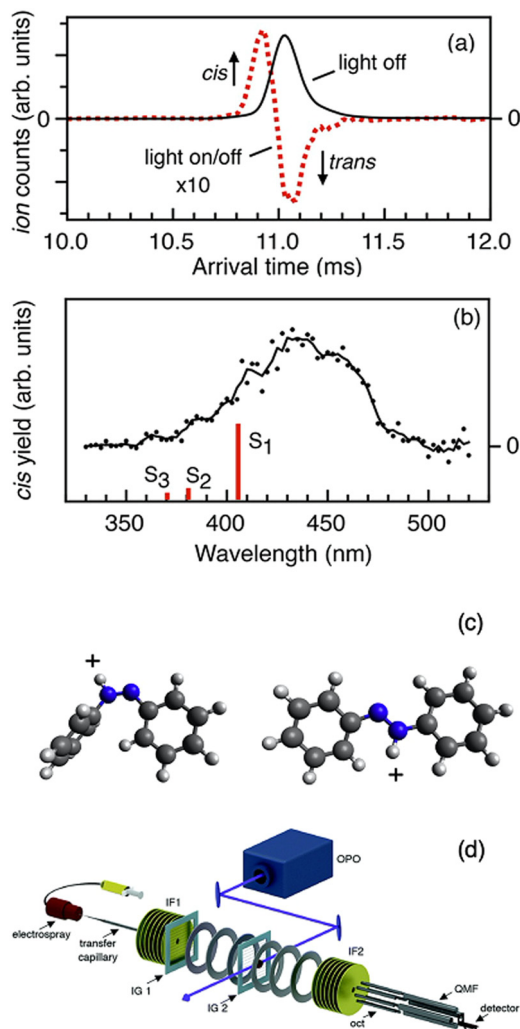


Fig. 13 (a) Arrival time distributions measured by the IMS2 drift region for *trans* azobenzene (that was selected by its mobility in the IMS1 region). The traces correspond to light off (black trace) and 440 nm photoaction (red dashes). (b) Action spectrum following the photoisomerization to the *cis* isomer as a function of wavelength. (c) Structures of protonated *cis* azobenzene and *trans* azobenzene. (d) Schematic of the tandem ion-mobility experimental setup. (a), (b) and (d) Adapted with permission from M. S. Scholz, J. N. Bull, N. J. Coughlan, E. Carrascosa, B. D. Adamson and E. J. Bieske, *J. Phys. Chem. A*, 2017, **121**, 6413–6419.<sup>207</sup>

selected in the first ion mobility step (IMS1) so only this isomer is present. This is confirmed by the IMS2 arrival time distribution, which displays only one peak (black trace in Fig. 13a). If the mobility selected ions are irradiated by a laser between IMS1 and IMS2, photoisomerism can be induced as revealed by the signal from the *cis* isomer and the depletion of the *trans* (dotted line in Fig. 13b). The *cis* isomer peak is assigned by its arrival time from the IMS2 spectrum.

Tandem ion mobility makes up a major proportion of the studies using combined ion-mobility and UV/vis action spectroscopy due to studies by the Bieske group<sup>72,205–218</sup> as well as the Dugourd group.<sup>219,220</sup> The example of protonated azobenzene highlights several key points about this technique. Ion mobility selection can allow for isomer specific investigation, tandem

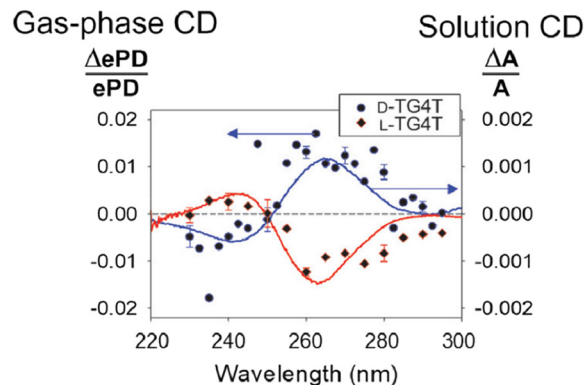


Fig. 14 Gas-phase circular dichroism spectrum of *m/z*-selected right-handed G-quadruplex (blue data points) and left-handed chiral isomer (red data points) compared to conventional solution phase circular dichroism spectrum (solid lines). Adapted with permission from V. Gabelica, *Accounts of Chemical Research*, 2021, **54**, 3691–3699.<sup>228</sup>

ion mobility can be used to study photoisomerism and the connection between conformational isomers.

### Chirality

Chirality affects the biomolecule function including physiological and toxicological behavior.<sup>221</sup> Conventional circular dichroism (CD) spectroscopy takes advantage of the fact that stereoisomers have different absorption values for left and right circularly polarised light. The development of gas phase PD techniques for distinguishing stereoisomers has emerged in recent years. The inherent advantages of coupling CD analysis with mass spectrometry include less sample quantity, lower sample concentration, and reduced purity concerns (the target can be *m/z*-isolated within the mass spectrometer) compared to conventional liquid-phase CD analysis.

Recently, experimental gas phase PD arrangements have successfully distinguished between stereoisomers of small chiral ions with linearly polarised light.<sup>222,223</sup> Experimental setups using circular polarised light to generate gas-phase CD action spectra have also been developed recently. Gas-phase CD action spectra have been reported for small ions in a cryogenic ion trap<sup>224,225</sup> and large DNA polyanions at room temperature.<sup>226</sup> Photoelectron circular dichroism of amino acid anions was also recently reported.<sup>227</sup>

A recent report, shown in Fig. 14, by Gabelica and co-workers described gas-phase chiral analysis where the signal is the normalised difference in electron photodetachment (ePD) yield arising from measurements alternating between left and right circularly polarised light. This is compared with the conventional solution phase CD spectrum (solid lines in Fig. 14). This approach demonstrates how gas phase laser and mass spectrometry techniques can be used to measure CD action spectra of selected ions.<sup>226</sup> It is early days for this application but the incorporation of CD analysis coupled embedded in an 'omics mass spectrometry is an alluring prospect.

### Final thoughts

Since the presence of isomers poses an outstanding challenge for many areas of mass spectrometry, it remains that more



technical advances are required before isomer separation and disentanglement become routine. In this Feature Article, we focused on the techniques for distinguishing isomers that involve UV and visible photodissociation with mass spectrometry. These strategies are also applicable to the study of metal-cluster topological isomers like Au<sub>25</sub>(SR)<sub>18</sub> (thiol-capped gold clusters),<sup>229</sup> which can be liberated into the gas phase with ESI and separated with ion-mobility.<sup>230–232</sup> Similar clusters have also been investigated with gas-phase techniques including photoelectron spectroscopy<sup>233</sup> and photodissociation<sup>234</sup> (but without isomer selection). Many of these experimental strategies are exciting prospects for further deployment and refinement. Incorporation of these techniques into routine mass spectrometry workflows remains an exciting avenue for exploitation. Furthermore, the application of mass spectrometry to areas including disease diagnostics, pathology and tissue imaging, and the numerous omics applications only further highlights the rich chemical information that will be available when isomer differentiation becomes commonplace. It was our intention to show here that photodissociation action spectroscopy strategies can provide definitive insights for a significant number of isomer classes. With suitable technical advancements, and reductions in size and cost of laser and optical hardware, it is easy to envisage a future where commercial mass spectrometers are equipped with tunable lasers affording visible, UV and VUV photoactivation. With these capabilities, PD action spectroscopy will be a vital tool to characterise the role of isomers in many areas including biological processes, physiology and the characterisation of diagnosis of disease.

## Conflicts of interest

There are no conflicts to declare.

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