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**Electronic spectra of positively charged carbon clusters -  $C_{2n}^+$  ( $n=6-14$ )**

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(Dated: 24 October 2021)

Electronic spectra are measured for mass-selected  $C_{2n}^+$  ( $n=6-14$ ) clusters over the visible and near infrared spectral range through resonance enhanced photodissociation of clusters tagged with  $N_2$  molecules in a cryogenic ion trap. The carbon cluster cations are generated through laser ablation of a graphite disk, and can be selected according to their collision cross section with He buffer gas and their mass prior to being trapped and spectroscopically probed. The data suggest that the  $C_{2n}^+$  ( $n=6-14$ ) clusters have monocyclic structures with bicyclic structures becoming more prevalent for  $C_{22}^+$  and larger clusters. The  $C_{2n}^+$  electronic spectra are dominated by an origin transition that shifts linearly to longer wavelength with the number of carbon atoms, and associated progressions involving excitation of ring deformation vibrational modes. Bands for  $C_{12}^+$ ,  $C_{16}^+$ ,  $C_{20}^+$ ,  $C_{24}^+$  and  $C_{28}^+$  are relatively broad, possibly due to rapid non-radiative decay from the excited state, whereas bands for  $C_{14}^+$ ,  $C_{18}^+$ ,  $C_{22}^+$  and  $C_{26}^+$  are narrower, consistent with slower non-radiative deactivation.

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## I. INTRODUCTION

Positively charged carbon clusters were first identified in mass spectrometric studies around 80 years ago,<sup>1</sup> and since then have been subject to extensive experimental and theoretical investigations concerned with their structures, reactivity and spectroscopic properties.<sup>2,3</sup> A significant outcome of these studies is a recognition of the structural diversity of carbon cluster cations.<sup>4</sup> Gas-phase ion mobility measurements have demonstrated that  $C_n^+$  clusters formed through laser ablation of graphite exist as linear isomers for  $n=3-10$ , monocycles for  $n \geq 7$ , bicycles for  $n \geq 21$  and fullerenes for  $n \geq 30$ .<sup>4</sup> The coexistence of linear and ring structures for smaller  $C_n^+$  clusters ( $n \leq 20$ ) was originally deduced on the basis of gas-phase reactivity studies from which it was concluded that linear clusters react rapidly with  $H_2$ ,<sup>5</sup>  $D_2$ ,<sup>6,7</sup>  $O_2$ ,<sup>6</sup>  $CO$ ,  $HCN$ ,  $H_2O$ ,<sup>8</sup> and hydrocarbons,<sup>9</sup> whereas the cyclic clusters react slowly, if at all, with these neutral species. Linear and cyclic isomers also decompose to different sized photofragments when exposed to 266 and 355 nm light.<sup>10,11</sup> The preference of mid-sized  $C_n^+$  ( $n=10-18$ ) clusters for cyclic rather than linear structures is consistent with electronic structure calculations.<sup>12</sup>

Spectroscopic studies of carbon cluster cations have mainly been restricted to species with fewer than ten carbon atoms, including  $C_6^+ - C_9^+$ ,<sup>13,14</sup> and the larger  $C_{60}^+$  and  $C_{70}^+$  fullerenes.<sup>15-17</sup> Significantly, the electronic spectrum of  $C_{60}^+$  exhibits features that match four diffuse interstellar bands (DIBs).<sup>15-18</sup> Although mid-sized carbon cluster cations with 10-30 carbon atoms have been characterized through their abundance, drift mobility, photodissociation behaviour and reactivity, there have been few measurements of their infrared and electronic transitions, which are crucial for identifying the presence of these species in remote environments, including interstellar space. To address this paucity of spectroscopic data, we have recorded electronic spectra of  $C_{2n}^+$  ( $n=6-14$ ) clusters, embracing a size range where the positively charged carbon clusters exist mainly as monocycles with the alternative bicyclic structure becoming progressively more important for clusters larger than  $C_{22}^+$ .<sup>4</sup> The presence of co-existing monocyclic, bicyclic and possibly linear structures for the  $C_{2n}^+$  ( $n=6-14$ ) clusters presents challenges for measuring and interpreting electronic and infrared spectra, and ideally requires mass- and isomer-selective approaches to yield unambiguous spectra. To address these challenges, we have adopted an approach whereby, following their formation, the  $C_n^+$  ions pass through a drift tube ion mobility spectrometer and then a quadrupole mass

filter, affording the capacity to separate and select the clusters on the basis of their collision cross sections with He buffer gas and their  $m/z$ . The gas-phase spectra are obtained by resonance enhanced photodissociation (REPD) of the target  $C_n^+$  clusters tagged with  $N_2$  molecules in a cryogenically cooled quadrupole ion trap.

To our knowledge, the only previous spectroscopic study of mid-sized carbon cluster cations was for  $C_{11}^+$ ,  $C_{12}^+$ ,  $C_{15}^+$ ,  $C_{16}^+$ ,  $C_{18}^+$  and  $C_{21}^+$  clusters that were generated by bombarding  $C_{60}$  with electrons, mass-selected and deposited in a Ne matrix where their electronic and infrared absorption spectra were measured.<sup>19,20</sup> The Ne matrix spectra also contained contributions from neutral, anion and fragment carbon clusters, potentially complicating their interpretation. Bands in the visible spectral region were assigned to linear carbon clusters as accompanying time dependent density functional theory (TD-DFT) calculations predicted that linear species absorb strongly in the visible spectral region, whereas cyclic isomers were calculated to possess strong absorptions below 250 nm.<sup>20</sup>

Experimental spectra of mid-sized carbon cluster cations with a controlled isomeric constitution, as obtained in the current study, should eventually help in elucidating the isomeric form of carbon clusters formed in different environments. For example,  $C_{12}^+$  and  $C_{18}^+$  clusters produced through dissociative ionization of overcrowded perchlorohydrocarbons exhibit the same reactivity and dissociation behaviour as clusters generated from the laser ablation of graphite, evidence that they share a common cyclic structure.<sup>21</sup> Also, it has been found that larger PAH cations such as hexa-peri-hexabenzocoronene ( $C_{42}H_{18}^+$ ) exposed to intense pulses of visible light ( $\lambda=532$  nm) lose peripheral H atoms to form cage structures and also smaller  $C_n^+$  clusters with  $n=16-20$ .<sup>22</sup> Furthermore, exposing the coronene radical cation ( $C_{24}H_{14}^+$ ) to UV radiation leads to complete dehydrogenation to form the  $C_{24}^+$  cluster.<sup>23</sup> Comparison of the electronic spectra of carbon clusters formed through decomposition of PAHs with electronic spectra reported in the current study may help decide whether they retain the skeletal structure of the parent molecule, or isomerize to some other structure. These considerations are relevant in the context of the proposed widespread existence of PAHs in interstellar space where exposure to UV light may cause loss of H atoms and formation of bare carbon clusters.<sup>23</sup>

The current spectroscopic study of the  $C_{2n}^+$  ( $n=6-14$ ) clusters complements extensive spectroscopic investigations of related neutral and charged carbon clusters, which have been motivated by their role in a range of environments, including in comets, carbon stars and

interstellar molecular clouds, and in flames and furnaces.<sup>2,3</sup> For example, electronic spectra have also been measured for smaller  $C_6^+$ - $C_9^+$  clusters deposited in a Ne matrix, with observed transitions assigned to both linear and cyclic isomers,<sup>13,14</sup> and for  $C_6^+$ -He<sub>*n*</sub> clusters in the gas phase.<sup>24</sup> Electronic spectra have been obtained for  $C_{60}^+$  and  $C_{70}^+$  fullerenes in Ne matrices,<sup>25,26</sup> and through REPD of  $C_{60}^+$ -He<sub>*n*</sub> and  $C_{70}^+$ -He<sub>*n*</sub> complexes in the gas-phase.<sup>15-17</sup> Gas-phase electronic spectra of  $C_{70}^{2+}$ -He<sub>*n*</sub> complexes have been obtained by similar means.<sup>27</sup> Electronic spectra have been obtained for linear  $C_{2n}$  ( $n=3-7$ ) and  $C_{2n+1}$  ( $n=2-10$ ) clusters in a Ne matrix,<sup>28-30</sup> and for cyclic  $C_{14}$ ,  $C_{18}$  and  $C_{22}$  molecules in the gas phase using resonance enhanced multiphoton photoionization.<sup>31,32</sup> Infrared spectra have been obtained for linear  $C_n$  clusters up to  $n=13$  in the gas phase<sup>3,33</sup> and in a Ne matrix.<sup>34</sup> Ionization potentials for neutral  $C_n$  ( $n=2-15$ ) clusters have been measured using tunable VUV radiation.<sup>35</sup> Negatively charged carbon clusters have been investigated using photoelectron spectroscopy,<sup>36,37</sup> matrix isolation spectroscopy,<sup>29,38</sup> and resonance enhanced photodetachment spectroscopy.<sup>39</sup>

Although carbon clusters have mainly been generated and characterized in the gas phase or in Ne matrixes, drawn by their predicted unique electronic and optical properties, synthetic chemists have sought to generate cyclocarbons in condensed phases.<sup>40,41</sup> One recent novel synthesis involved production of  $C_{18}$  on a cooled NaCl surface by elimination of CO molecules from the  $C_{24}O_6$  cyclocarbon oxide molecule.<sup>42</sup> High resolution atomic force microscopy showed that  $C_{18}$  formed in this way has a cyclic polyynic structure, as predicted by electronic structure calculations.<sup>43</sup> Ultimately, gas-phase electronic spectra of cyclocarbon cations reported in this paper should provide useful data for benchmarking theoretical approaches, which, in turn, should help guide synthetic condensed-phase studies.

## II. EXPERIMENTAL METHODS

Spectra of the  $C_n^+$  clusters were obtained by REPD of cryogenically cooled  $C_n^+$ -(N<sub>2</sub>)<sub>*m*</sub> complexes in a three-dimensional quadrupole ion trap (3D QIT). The  $C_n^+$  clusters were generated by laser ablation of a rotating carbon disk using the focused, frequency-doubled output of a pulsed Nd:YAG laser ( $\lambda=532$  nm, 12 mJ/pulse, 20 Hz). The ablation source is situated at the beginning of a drift tube ion mobility spectrometer with which, if required, the clusters can be separated according to their drift time through He buffer gas ( $P\approx 2$  Torr) using a pulsed Bradbury-Nielsen ion gate located at the end of the drift region. Following

passage through the drift region, the charged clusters were collected by an RF ion funnel and passed through a 1 mm orifice into an RF hexapole ion guide (pressure typically  $5 \times 10^{-5}$  Torr), which was used to accumulate ions from 10 ablation laser pulses over 0.5 s. After their exit from the hexapole, the ions travelled through a quadrupole mass filter where they were mass-selected. The ions then passed through an octupole ion guide and into the QIT cooled to  $T \approx 20$  K into which pulses of a He/N<sub>2</sub> 100:1 gas mixture were injected at 2 Hz. The C<sub>*n*</sub><sup>+</sup> clusters were tagged with N<sub>2</sub> molecules in the QIT before being ejected (after 480 ms) into a linear time-of-flight (ToF) mass spectrometer (length 0.9 m). Alternate packets of ions in the QIT were exposed to pulses of unfocused, wavelength-tunable light from an optical parametric oscillator (OPO, EKSPLA NT 342B,  $\leq 1$  mJ/cm<sup>2</sup>/pulse, 6 ns pulse width, bandwidth  $\approx 4$  cm<sup>-1</sup>). The ions were typically exposed to a single pulse of light from the tunable OPO around 400 ms after their introduction into the trap at the end of the trapping cycle. The effect of the light was manifested in the difference between the light-on and light-off ToF mass spectra; when the OPO was tuned to a resonance, the C<sub>*n*</sub><sup>+</sup>-(N<sub>2</sub>)<sub>*m*</sub> signal diminished and the C<sub>*n*</sub><sup>+</sup> photofragment signal grew. As an example, FIG. 1 displays the difference ToF mass spectrum for C<sub>20</sub><sup>+</sup>-(N<sub>2</sub>)<sub>*m*</sub> with the OPO tuned to 897 nm (the origin band for C<sub>20</sub><sup>+</sup>-N<sub>2</sub>). Normally, the light intensity was adjusted so that  $\leq 10\%$  of the N<sub>2</sub> tagged ions were dissociated. Where possible we attempted to obtain spectra with a range of laser powers to test whether power broadening was an issue.

Electronic spectra were obtained by plotting either the integrated C<sub>*n*</sub><sup>+</sup>-(N<sub>2</sub>)<sub>*m*</sub> photodepletion signal or the C<sub>*n*</sub><sup>+</sup> photofragment signal as a function of wavelength with intensities normalised by OPO power. Wavelength calibration was accomplished with a wavemeter (High Finesse, LSA). It is relevant to note that exposure to light over the visible and near-infrared range caused no appreciable photofragmentation of the core C<sub>*n*</sub><sup>+</sup> clusters.

### III. RESULTS AND DISCUSSION

#### A. Ion mobility data

The isomeric composition of carbon clusters produced by the laser ablation source was assessed using the drift tube ion mobility spectrometer in which the ions were separated according to their collision cross section with the He buffer gas; compact ions travel more

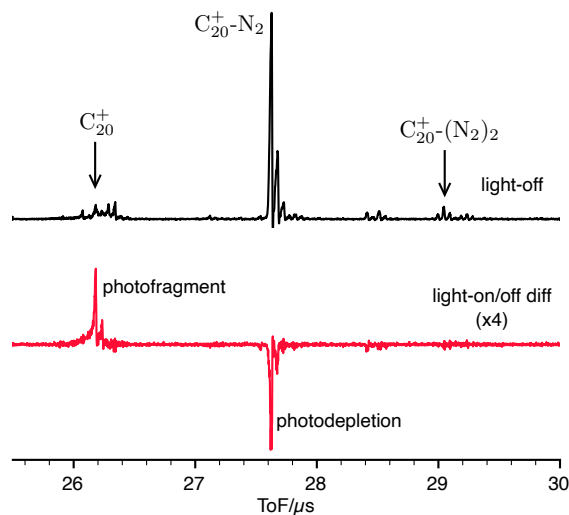


FIG. 1. Light-off and light-on/off difference ToF mass spectra for  $C_{20}^+-(N_2)_m$  ( $m=0-2$ ) clusters with the OPO tuned to 897 nm, the strongest absorption of  $C_{20}^+-N_2$  in the visible/NIR.

quickly through the buffer gas and arrive at the detector earlier than extended ions. FIG. 2 shows a plot of ion current versus arrival time and  $m/z$  for  $C_n^+$  clusters with  $n=10-60$ . The data demonstrate that the  $n=10-28$  clusters are predominantly monocycles, with larger  $n=22-40$  clusters existing as both monocycles and more compact bicycles, and with fullerenes becoming increasingly apparent for  $n \geq 32$ . Families of multiply charged fullerenes ( $C_n^{2+}$ ,  $C_n^{3+}$ ,  $C_n^{4+}$ ) are also evident at shorter arrival times. Relative abundances of the different clusters depend on ion source conditions, particularly the power of the ablation laser. Our ion mobility measurements are consistent with  $C_n^+$  mobility data from von Helden *et al.*<sup>4</sup> For example, drift times for the monocyclic, bicyclic and fullerene isomers of  $C_{36}^+$  have ratios of 1.93:1.76:1, close to drift time ratios calculated using the measured mobilities reported in ref. 4 (1.97:1.80:1).

For the spectroscopic investigations of the  $C_{12}^+$  to  $C_{28}^+$  clusters described below, the apparatus was normally run without mobility selection of the target complex because the monocyclic isomer is present in far greater abundance than the bicyclic isomer and the multiply charged fullerenes that share the same  $m/z$ . In support of this approach we conducted tests for  $C_{28}^+$ , and found that gating the slower  $C_{28}^+$  ions, corresponding to the monocyclic isomer, gave rise to sharp peaks over the 1000-1400 nm wavelength range, whereas the faster  $C_{28}^+$  ions associated with the bicyclic isomer did not absorb appreciably in this region.

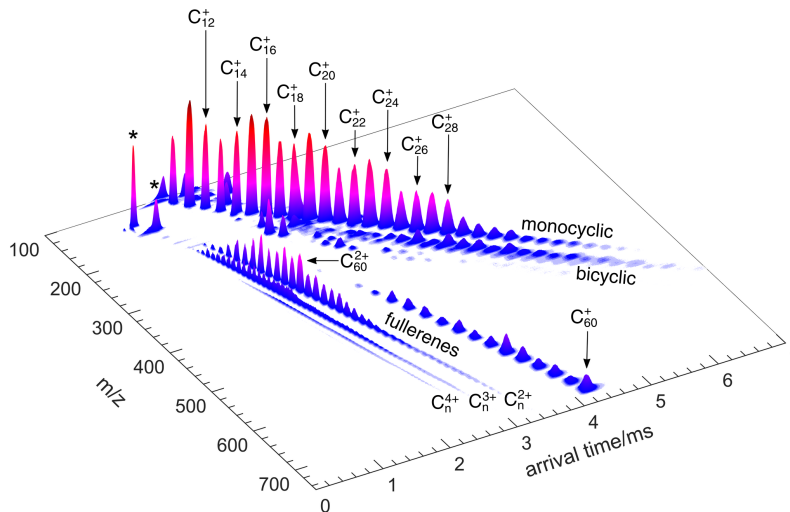


FIG. 2. Ion intensity versus drift time and  $m/z$  for  $C_n^+$  clusters with conditions optimised for generation and detection of smaller clusters. Asterisked peaks with maxima at  $m/z$  138 and 154 are tentatively assigned to impurity  $Ba^+$ ,  $BaO^+$  and  $BaOH^+$  ions.

## B. Electronic spectra

Electronic spectra of  $C_{2n}^+$  ( $n=6-14$ ) carbon clusters tagged with  $N_2$  molecules measured over the visible to NIR range are shown in FIG. 3. Presumably, the observed transitions are from the doublet ground state to the lowest lying doublet excited state of the clusters. Wavenumbers for the peaks are listed in Table I. Each of the  $C_{2n}^+$  ( $n=6-14$ ) spectra is dominated by an origin transition flanked by weaker peaks  $300-600\text{ cm}^{-1}$  to higher energy that presumably correspond to vibronic progressions in ring deformation modes. The most striking feature of the  $C_n^+$  spectra series is that the origin transition shifts linearly to longer wavelength with  $n$  (see FIG. 4). This, along with the ion mobility data shown in FIG. 2 constitutes strong evidence that the  $C_n^+$  clusters responsible for the absorptions have a common structural motif. A similar linear dependence of absorption wavelength on cluster size has been observed for several different homologous linear carbon and species, including  $C_{2n}$ ,  $C_{2n}^-$ ,  $C_{2n}H^-$ ,<sup>30</sup>  $C_{2n+1}$ ,<sup>28</sup> and  $HC_{2n}H^+$ .<sup>44</sup> Perhaps significantly, the wavelength for the  $\tilde{E} \leftarrow \tilde{X}$  origin transition of cyclic  $C_8^+$ , previously measured at 336.5 nm in a Ne matrix, fits with the trend for the  $n=12-28$  series.<sup>14</sup> We have unsuccessfully sought the transition for  $C_{10}^+$ , which, by extrapolation, is predicted to occur in a wavelength region where the output



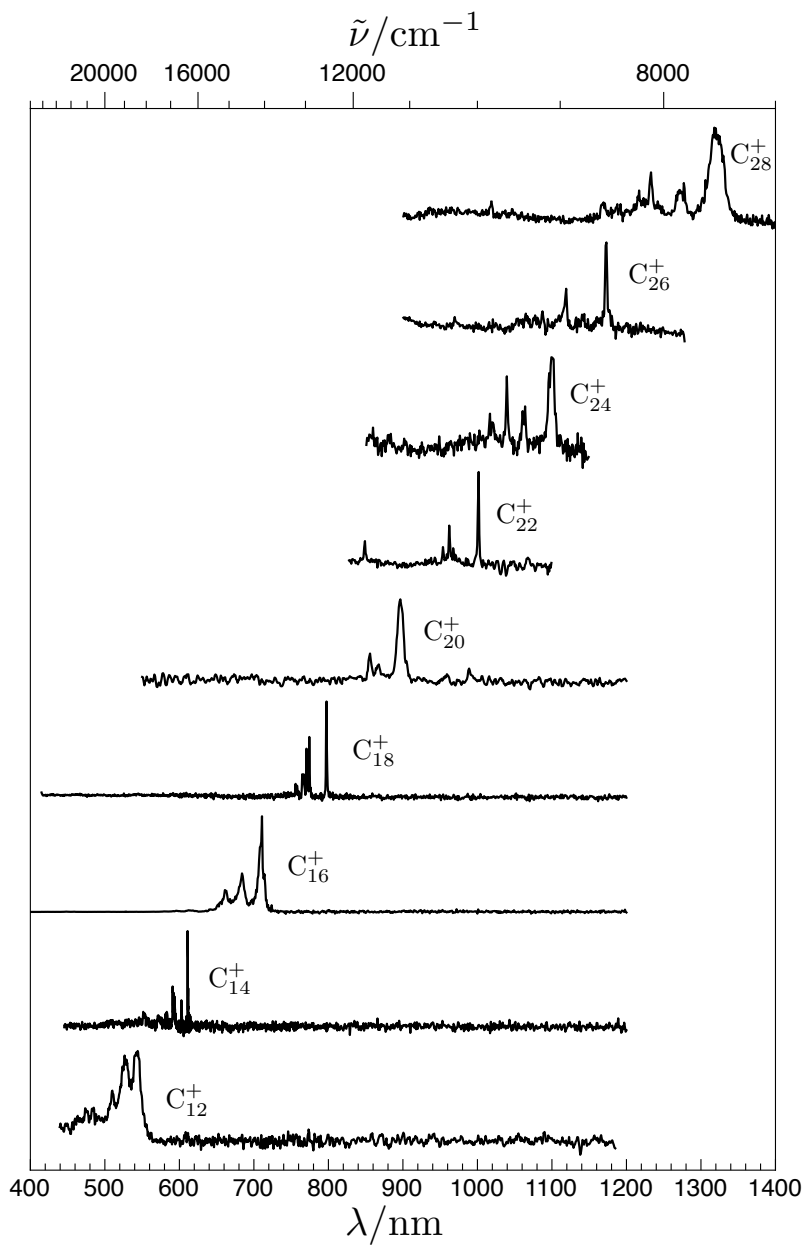


FIG. 3. Electronic spectra for  $C_n^+-(N_2)_n$  clusters obtained by monitoring  $C_n^+$  photofragments following laser excitation. The spectra, which are normalized with respect to OPO power, may be slightly saturated such that the intensities of weaker peaks are exaggerated with respect to the origin transition.

of the OPO is weak ( $\approx 410$  nm).

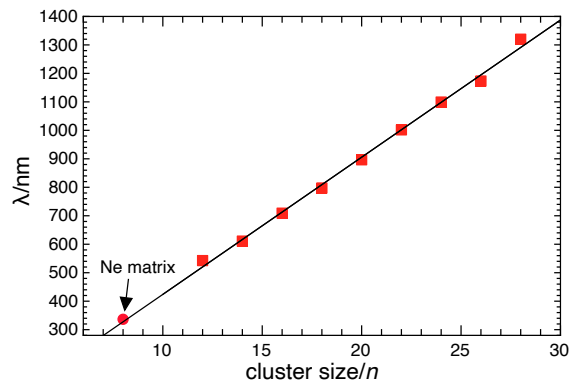


FIG. 4. Plot of the origin band wavelengths versus  $n$  for even  $C_n^+$  clusters. The  $n=8$  point at 336.5 nm corresponds to the  $\tilde{E} - \tilde{X}$  origin transition of cyclic  $C_8^+$  in a Ne matrix (from ref. 14).

TABLE I: Wavenumbers ( $\text{cm}^{-1}$ , vacuum) for transitions of  $\text{C}_n^+$  clusters tagged with  $\text{N}_2$  molecules. Positions are given for spectra obtained by monitoring formation of  $\text{C}_n^+$  photofragments [resulting from photofragmentation of both  $\text{C}_n^+-\text{N}_2$  and  $\text{C}_n^+(\text{N}_2)_2$ ], and from photodepletion spectra of  $\text{C}_n^+-\text{N}_2$  and  $\text{C}_n^+(\text{N}_2)_2$ . Uncertainties in positions depend on cluster size and are estimated as  $\pm 40 \text{ cm}^{-1}$  ( $n=12$ ),  $\pm 5 \text{ cm}^{-1}$  ( $n=14, 18, 22, 26$ ),  $\pm 20 \text{ cm}^{-1}$  ( $n=16, 20, 24, 28$ ).

$n$	assignment	mass channel monitored					
		$\text{C}_n^+$		$\text{C}_n^+-\text{N}_2$		$\text{C}_n^+(\text{N}_2)_2$	
		photofrag.		depletion		depletion	
		$\tilde{\nu}$	$\Delta\tilde{\nu}$	$\tilde{\nu}$	$\Delta\tilde{\nu}$	$\tilde{\nu}$	$\Delta\tilde{\nu}$
12	$0_0^0$	18426	0	18444	0	18481	0
		18934	508	18934	490	18988	507
		19621	1195	19659	1215	19699	1218
				-	-	20848	2367
14	$0_0^0$	16338	0	16338	0	16362	0
		16362	24	-	-	-	
		16585	247	16565	227	16590	228
		16830	492	-	-	16836	474
		16873	535	16849	511	16873	511
		16912	574			16921	559
16	$0_0^0$	14099	0	14090	0	14082	0
		14615	516	14615	525	14615	533
		15101	1002	15112	1022	15112	1030
18	$0_0^0$	12536	0	-	-	12536	0
		12910	374	-	-	12918	382
		12973	437	-	-	12973	437
		13053	517	-	-	13053	517
		13189	653	-	-	13189	653
		13225	689				
20	$0_0^0$	11151	0	11125	0	-	-
		11536	385	11510	385	-	-
		11694	543	11694	569	-	-
22	$0_0^0$	9981	0	9981	0	-	-
		10333	352	10333	352	-	-
		10390	409	10390	409	-	-

*Continued on next page*

TABLE I – *Continued from previous page*

$n$ assignment		mass channel monitored					
		$C_n^+$		$C_n^+-N_2$		$C_n^+-(N_2)_2$	
		photofrag.		depletion		depletion	
		$\tilde{\nu}$	$\Delta\tilde{\nu}$	$\tilde{\nu}$	$\Delta\tilde{\nu}$	$\tilde{\nu}$	$\Delta\tilde{\nu}$
		10482	501	10482	501	-	-
		11775	1794	11775	1794	-	-
24	$0_0^0$	9094	0	9094	0	-	-
		9416	322	9416	322	-	-
		9618	524	9618	524	-	-
		9804	710	9813	719	-	-
26	$0_0^0$	8522	0	8522	0	8522	0
		8933	411	8942	420	8951	429
28	$0_0^0$	7576	0	7570	0	-	-
		7855	279	7837	267	-	-
		8117	541	8110	540	-	-

There are several reasons to believe that the absorptions shown in FIG. 3 are due to monocyclic  $C_n^+$  clusters. First, as noted above, the ion mobility data for the  $C_n^+$  clusters with  $n \geq 10$  are similar to previously reported measurements,<sup>4</sup> and are consistent with monocyclic structures for  $n=10-20$  and the coexistence of monocyclic and bicyclic structures for  $n=22-28$ . Second, for clusters with  $n \geq 10$  the monocyclic clusters are calculated to lie considerably lower in energy than the linear clusters.<sup>12</sup> For example, cyclic  $C_{16}^+$  is calculated to be 3.24 eV more stable than linear  $C_{16}^+$  at the  $\omega$ B97X-D/cc-pVTZ level of density functional theory (DFT). Third, linear clusters, if they do form through isomerization of cyclic clusters following their passage through the ion mobility spectrometer drift region - either when they were stored in the hexapole, passing through the octopole ion guide, or contained in the cryogenic QIT - would be expected to react rapidly with trace  $H_2O$  or  $H_2$  to form linear  $C_nH^+$  or  $HC_nH^+$  molecules.<sup>5-8</sup> We tested this by recording spectra of the  $C_n^+$  clusters with both  $H_2/N_2$  (100:1) and  $He/N_2$  (100:1) buffer gas mixtures in the QIT and found no substantial differences in ToF mass spectra or in the electronic spectra. Last, preliminary experiments with mobility selection of the  $C_n^+$  clusters clearly show that the spectral features apparent in FIG. 3 are associated with the slower, less compact monocyclic clusters rather than with

the faster, more compact bicyclic clusters. Furthermore, when the bicyclic clusters were selected there was no trace of the sharp spectral features indicating that collision-induced bicyclic→monocyclic isomerization does not occur following mobility selection of the clusters in the drift region.

Some of the transitions measured for  $C_n^+$  clusters in the gas phase correspond to bands previously reported for mass-selected  $C_n^+$  clusters in a Ne matrix, where the ions were generated by subjecting  $C_{60}$  vapour to electron impact.<sup>19,20</sup> For example, the strongest bands in the gas-phase  $C_{16}^+$  spectrum at 710.87 and 684.05 nm (FIG. 3) correspond in positions and relative intensities to prominent, unassigned peaks near 710 and 685 nm in the Ne matrix spectrum obtained when  $C_{16}^+$  was mass-selected and deposited (FIG. S12 in the Supporting Information for ref. 20). Furthermore, the strongest bands in the gas-phase spectrum for  $C_{20}^+$  shown in FIG. 3 at 896.5 nm and 854.9 nm correspond in position and relative intensities to peaks at 899 and 857 nm reported in ref. 19 for carbon cluster cations with  $m/z$  240 deposited in a Ne matrix, which were attributed to  $C_{60}^{3+}$ . For the following reasons, the species responsible for the absorptions is almost certainly  $C_{20}^+$  rather than  $C_{60}^{3+}$ . First, the ion mobility data show that for typical operating conditions the laser ablation ion source produces relatively small quantities of  $C_{60}^{3+}$  compared to  $C_{20}^+$  (see FIG. 2). Second, the ToF mass peak for the  $m/z$  240 species tagged with  $N_2$ , which was irradiated to obtain the gas-phase spectrum, appears at  $m/z$  268 (as expected for  $C_{20}^+-N_2$ ) rather than at  $m/z$  249.3 (where it would be for  $C_{60}^{3+}-N_2$ ). Finally, the wavelength of the origin band at 896.5 nm fits the pattern established for the other  $C_n^+$  clusters (see FIG. 4), which are certainly not multiply charged fullerenes.

There are few theoretical predictions for the transition energies of monocyclic  $C_n^+$  clusters that can be compared with the experimental data reported here. Although large-scale multi-reference configuration interaction (MRD-CI) calculations have been carried out for the smaller  $C_6^+$  and  $C_8^+$  clusters and predict electronic transitions with weak to medium intensity occur across the visible and NIR spectral range,<sup>45</sup> calculations of similar quality are unavailable for larger cyclic clusters. TD-DFT calculations reported in ref. 20 predict that cyclic  $C_n^+$  ( $n=10-20$ ) clusters have very strong transitions below 250 nm with oscillator strengths ranging from 1.4 to 7.7. Wavelengths for transitions occurring in the visible region, which were calculated to be relatively weak, were not reported.<sup>20</sup>

### C. Vibronic structure

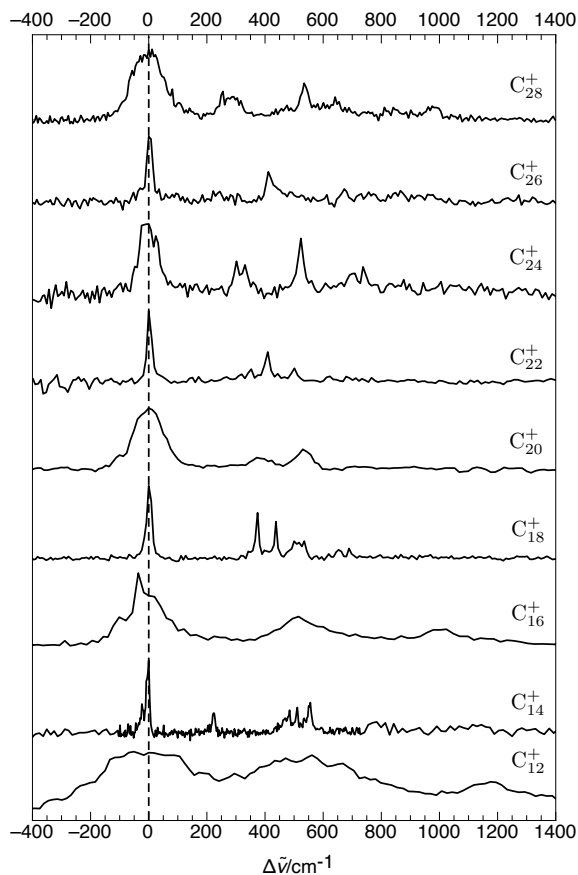


FIG. 5. Expanded view of the spectra for  $C_n^+$  clusters. The dashed vertical line represents the positions of the aligned origin bands for the  $C_n^+$  clusters.

In each of the  $C_n^+$  spectra, the strongest band is flanked by weaker peaks lying 300–600  $\text{cm}^{-1}$  to higher energy that are presumably associated with progressions involving the low frequency ring vibrational modes. These spectral features can be appreciated in FIG. 5 where the origin bands for the different  $C_n^+$  clusters are aligned. Calculated vibrational frequencies for  $C_{12}^+$ ,  $C_{14}^+$  and  $C_{16}^+$  in their ground electronic states are given in the Supplemental Material. Assigning these peaks to particular vibrational modes is difficult, given that the geometry and vibrational frequencies for the clusters in their excited electronic states are unknown, and because of the large number of low frequency vibrational modes. For example,  $C_{14}^+$  has 15 in-plane vibrational modes lying below 1000  $\text{cm}^{-1}$  (see Table S7 in the Supplementary Material). One can only say that if the rings have a cumulenic structure with equal C=C bond lengths and C=C=C bond angles in the ground and excited states ( $D_{nh}$  symmetry) then

the band systems should only exhibit progressions in the totally symmetric ring expansion mode. Presence of additional peaks indicates that the molecules have a lower symmetry in one or both of these electronic states.

#### D. Effect of N<sub>2</sub> tag molecules

Because the spectra are obtained through resonance enhanced photodissociation of C<sub>*n*</sub><sup>+</sup>-(N<sub>2</sub>)<sub>*m*</sub> clusters, it is relevant to consider the effect of the attached N<sub>2</sub> molecule(s) on the structures and transition energies of the carbon clusters. DFT calculations conducted as part of this study at the  $\omega$ B97X-D/aug-cc-pVDZ level of theory suggest that for cyclic clusters up to C<sub>18</sub><sup>+</sup>, an attached N<sub>2</sub> molecule prefers to sit above or below the ring (see FIG. 6), but that for C<sub>19</sub><sup>+</sup> and larger clusters, a single N<sub>2</sub> molecule can be accommodated inside the ring. Typically, the N<sub>2</sub> molecules are bound by several hundred cm<sup>-1</sup>. For example, the binding energy of N<sub>2</sub> to C<sub>14</sub><sup>+</sup> is calculated to be 790 cm<sup>-1</sup>, with the second N<sub>2</sub> bound by a similar amount (775 cm<sup>-1</sup>). The dissociation energy of C<sub>20</sub><sup>+</sup>-N<sub>2</sub>, for which the N<sub>2</sub> molecule resides inside the ring, is calculated to be slightly higher (984 cm<sup>-1</sup>). The attached N<sub>2</sub> molecules are predicted to have a minor effect on the vibrational frequencies of the core C<sub>*n*</sub><sup>+</sup> clusters, with shifts of less than 20 cm<sup>-1</sup> (see Table S7 in the Supplemental Information for calculated frequencies of C<sub>14</sub><sup>+</sup>, C<sub>14</sub><sup>+</sup>-N<sub>2</sub> and C<sub>14</sub><sup>+</sup>-(N<sub>2</sub>)<sub>2</sub>). Frequencies of the intermolecular modes involving relative motion of the ring and N<sub>2</sub> molecule have relatively low calculated harmonic frequencies (for example, 13, 49, 67, 78, 97 cm<sup>-1</sup> for C<sub>14</sub><sup>+</sup>-N<sub>2</sub>). The absence of progressions with peak spacings less than 100 cm<sup>-1</sup> in the electronic spectra indicates that the interaction between the C<sub>*n*</sub><sup>+</sup> and N<sub>2</sub> molecule is largely unaffected by electronic excitation.

In some instances we have managed to measure spectra of both C<sub>*n*</sub><sup>+</sup>-N<sub>2</sub> and C<sub>*n*</sub><sup>+</sup>-(N<sub>2</sub>)<sub>2</sub> clusters. For example, FIG. 7 shows photodepletion action spectra for C<sub>14</sub><sup>+</sup>-N<sub>2</sub> and C<sub>14</sub><sup>+</sup>-(N<sub>2</sub>)<sub>2</sub>, which have origin peaks at 611.9 and 611.0 nm, respectively. Assuming that each of the two attached N<sub>2</sub> molecules causes the same spectral shift, as expected for the structures shown in FIG. 6, the origin transition of the bare C<sub>14</sub><sup>+</sup> cluster would lie at 612.8 nm.

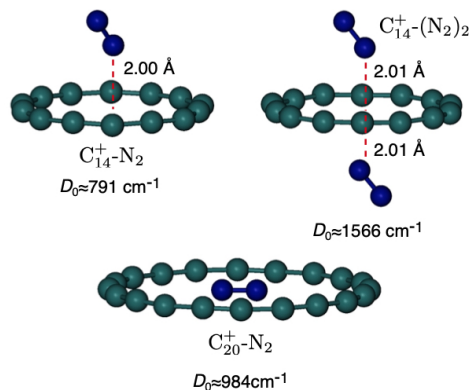


FIG. 6. Minimum energy structures for  $C_{14}^+-N_2$ ,  $C_{14}^+-(N_2)_2$  and  $C_{20}^+-N_2$  showing the  $N_2$  tag molecule sitting above and below the plane of the ring for  $C_{14}^+$  and located in the middle of the ring for  $C_{20}^+$ . Distances are given from the closest N atom to the plane of the ring.

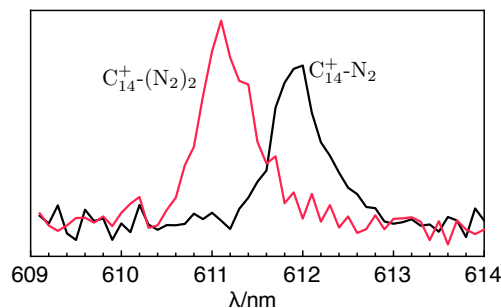


FIG. 7. Origin transitions for photodepletion spectra of  $C_{14}^+-N_2$  and  $C_{14}^+-(N_2)_2$ .

### E. Transition bandwidths

A distinctive feature of the spectral series shown in FIGS. 3 and 5 is that the transitions for the  $C_{4k}^+$  and  $C_{4k+2}^+$  clusters have different widths. Electronic spectra of the  $C_{4k+2}^+$  clusters ( $C_{14}^+$ ,  $C_{18}^+$ ,  $C_{22}^+$ ,  $C_{26}^+$ ) exhibit relatively sharp peaks (FWHM 10-20  $\text{cm}^{-1}$ ), whereas peaks for the  $C_{4k}^+$  clusters ( $C_{12}^+$ ,  $C_{16}^+$ ,  $C_{20}^+$ ,  $C_{24}^+$ ,  $C_{28}^+$ ) are noticeably broader (FWHM > 100  $\text{cm}^{-1}$ ). The peak profiles will be influenced to some extent by the presence of the attached  $N_2$  molecule(s) due to hot band transitions involving low frequency intermolecular modes. However, the broadening from this source should be similar for both series of clusters. A possible explanation for the broader peaks for the  $C_{4k}^+$  clusters involves lifetime broadening associated with rapid nonradiative deactivation through intersystem crossing (ISC). The



lowest quartet state for the  $C_{4k}^+$  clusters is calculated to lie just above the doublet ground state (by 0.4 eV for both  $C_{12}^+$  and  $C_{16}^+$  according to DFT calculations<sup>12</sup>), such that there should be a high density of quartet vibronic states iso-energetic with the excited doublet state, facilitating rapid intersystem crossing. In contrast, the lowest quartet state for the  $C_{4k+2}^+$  clusters is predicted to lie much higher above the ground state (by 3.5 eV for  $C_{10}^+$ , 2.7 eV for  $C_{14}^+$ , 1.5 eV for  $C_{18}^+$ ) so that there should be lower densities of quartet vibrational states iso-energetic with the excited doublet states and consequently lower ISC crossing rates.

## F. Comparison with DIBs

Given that the bands of the  $C_{12}^+$ - $C_{28}^+$  clusters span the visible to near infrared region of the spectrum, it is appropriate to consider whether any of the cyclic  $C_n^+$  clusters are carriers of diffuse interstellar bands. Carbon cluster cations can conceivably form in the interstellar medium through a process whereby positively charged polyaromatic hydrocarbons progressively lose H atoms and rearrange following absorption of UV photons,<sup>23</sup> leading to bare dehydrogenated  $C_n^+$  clusters, which, over the  $n=10-28$  range, may have monocyclic structures. Electronic spectra of the neutral carbon rings  $C_{14}$ ,  $C_{18}$  and  $C_{22}$  have previously been recorded, but no convincing match was found between the measured peaks and any DIBs.<sup>31,32</sup> Although the bands of the  $C_{4k}^+$  series are probably too broad to be associated with DIBs, several of the  $C_{4k+2}^+$  clusters exhibit sharper transitions that are close to DIBs. For example, six DIBs have been measured over the 610-613 nm range in the spectrum for HD 183143 (at 611.079, 611.329, 611.684, 611.874, 612.443, 612.820 nm),<sup>46</sup> nearby the strongest peak for  $C_{14}^+$ - $N_2$  at 611.9 nm. However, even if the origin transition of the  $C_{14}^+$  band corresponds to one of these DIBs there are no reported DIBs with appropriate spacings, relative intensities and widths to explain the two shorter wavelength transitions (at 603.5 and 593.3 nm for  $C_{14}^+$ - $N_2$ ). A similar situation pertains for  $C_{18}^+$ . The most intense transition of  $C_{18}^+$ - $(N_2)_2$  occurs at 797.5 nm, nearby DIBs for HD 183143 at 796.809 nm and 797.192 nm.<sup>46</sup> However, there are no appropriate matches for the  $C_{18}^+$ - $(N_2)_2$  transitions near 773.9 and 770.6 nm. Ultimately, spectral shifts associated with the  $N_2$  tag molecules make absolute comparisons with astronomical data impossible, and higher resolution spectra of the bare clusters are required to decide whether any of the  $C_n^+$  clusters are carriers for DIBs.

#### IV. CONCLUSIONS AND OUTLOOK

In summary, we have obtained electronic spectra of the  $C_{2n}^+$  ( $n=6-14$ ) clusters, which the available evidence suggests have monocyclic structures, and which, aside from the fullerenes, include the largest carbon clusters for which electronic spectra have been measured. The most remarkable feature of the series is the linear dependence of the origin transition wavelength on the number of carbon atoms in the cluster. Although the wavelengths for the absorptions of the  $C_n^+-(N_2)_m$  clusters will be shifted slightly from those of the bare  $C_n^+$  clusters, the spectra reported in this paper should provide a foundation for future measurements for  $C_n^+$  clusters tagged by He or tag-free spectra to determine whether the absorptions correspond to DIBs.

The  $C_n^+$  spectra reported in this paper also provide a new route for identifying the isomeric constitution of charged carbon clusters generated through decomposition of larger carbon-bearing molecules and clusters. Carbon clusters in the  $C_{12}^+$  to  $C_{28}^+$  range have been produced through dehydrogenation of PAHs,<sup>23,47</sup> from overcrowded perchloro hydrocarbons,<sup>21</sup> and electron impact of  $C_{60}^+$ .<sup>20</sup> In principle, electronic spectra of  $C_n^+$  clusters generated in these different ways can be compared with spectra presented in this study to see whether the product clusters are monocycles or have some other structure.

There are several avenues to extend the spectroscopic investigations. First, it would be interesting to examine larger clusters for which there is a coexistence of monocyclic, bicyclic and fullerene structures (for example  $C_{36}^+$ ). It remains to be seen whether bicyclic  $C_n^+$  structures give rise to an orderly series of electronic absorptions as found for linear and ring structures. One complication for clusters with 30 or more carbon atoms is the existence of different isomers with similar collision cross sections but which have distinct structures and electronic spectra. For example, bicyclic  $C_{32}^+$  may have an isomer with 16 atoms in each ring, and another isomer with 14 atoms in one ring and 18 atoms in the other. Such isomers should have distinct electronic spectra, but may be difficult to separate using ion mobility techniques.

Development of robust theoretical treatments that properly describe the ground and excited state properties of the  $C_n^+$  clusters is desirable but represents an ongoing challenge. While density functional theory methods to obtain cluster geometries followed by time-dependent frameworks for electronic transition wavelengths and intensities might seem

computationally tractable, recent studies on a range of carbon clusters have highlighted erroneous results when using many density functionals in standard computational software packages, including not correctly describing the change in bond length alternation for either linear or cyclic isomers with cluster size.<sup>43,48,49</sup> Furthermore, accurate calculations are potentially complicated by Jahn-Teller effects<sup>50</sup> and for cations require open-shell methods which are notorious for wavefunction convergence difficulties. Hopefully these difficulties will be resolved so that theory can better guide the interpretation of experimental spectroscopic data.

## **SUPPLEMENTARY MATERIAL**

The supplementary material contains details of the density functional theory calculations and includes calculated equilibrium structures and vibrational frequencies for  $C_{12}^+$ ,  $C_{14}^+$ ,  $C_{16}^+$  and  $C_{20}^+$ .

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## **CONFLICT OF INTEREST**

The authors have no conflicts to disclose.

## **DATA AVAILABILITY STATEMENT**

The data that support the findings of this study are available from the corresponding author upon reasonable request.

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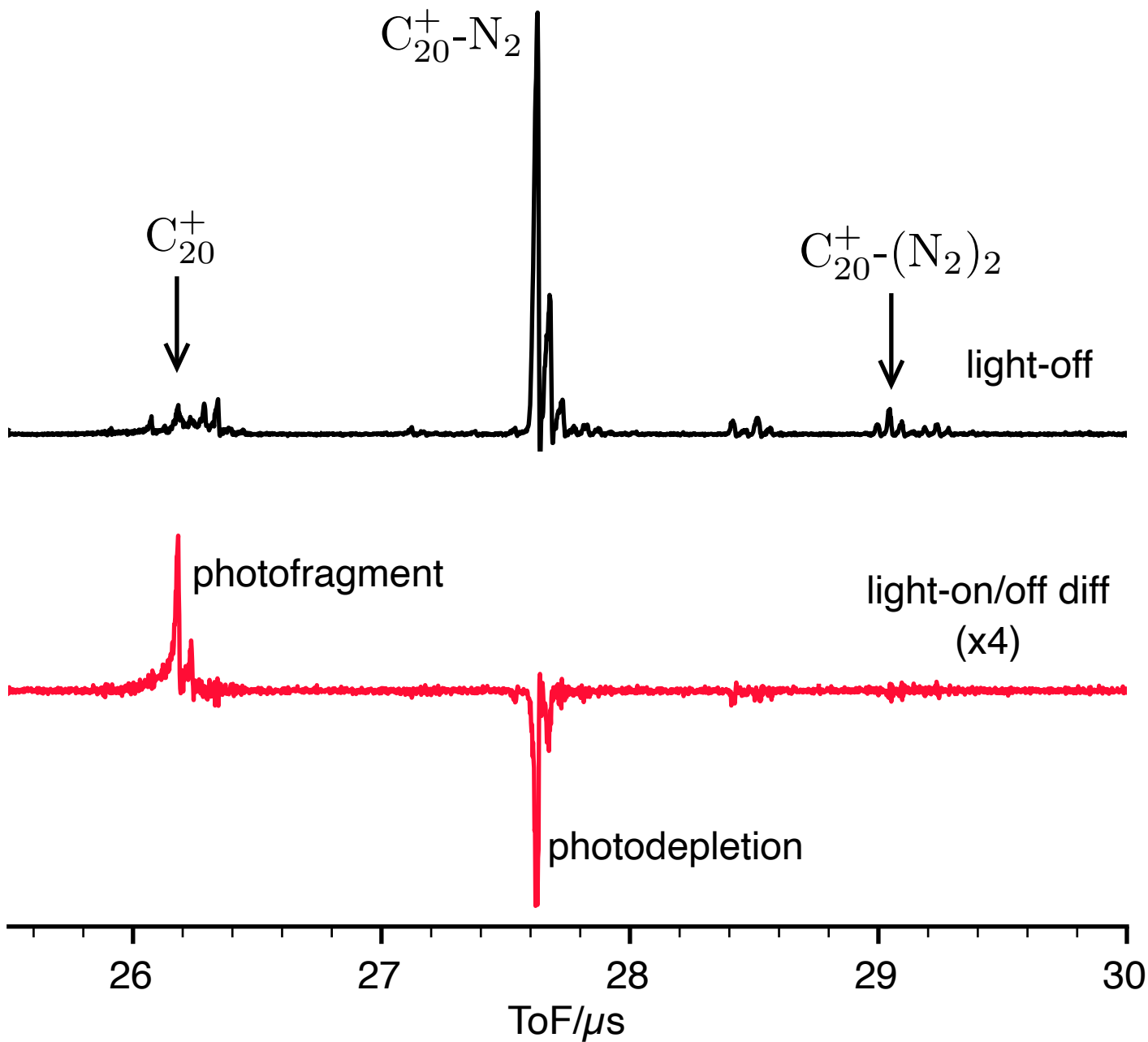
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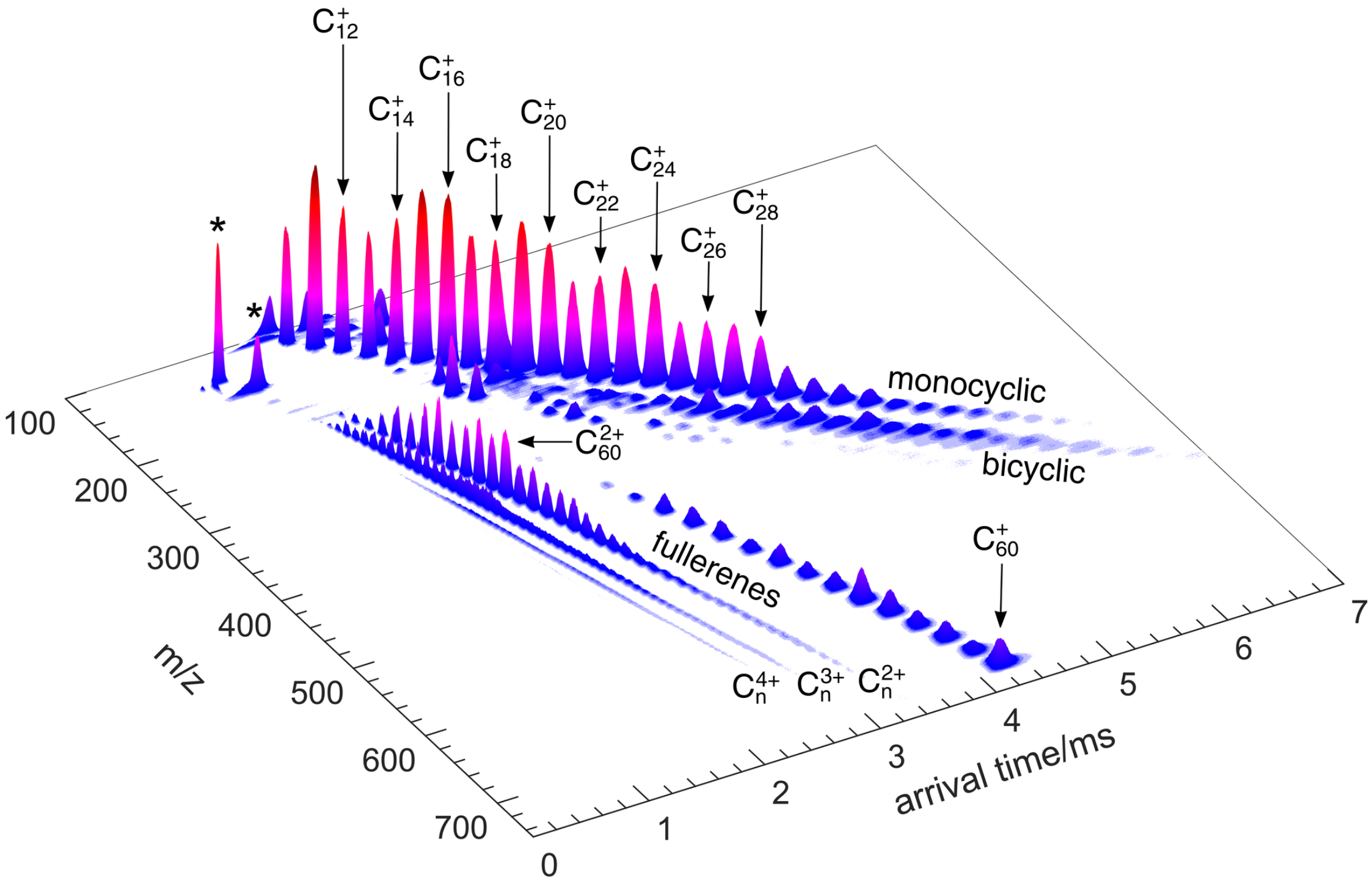
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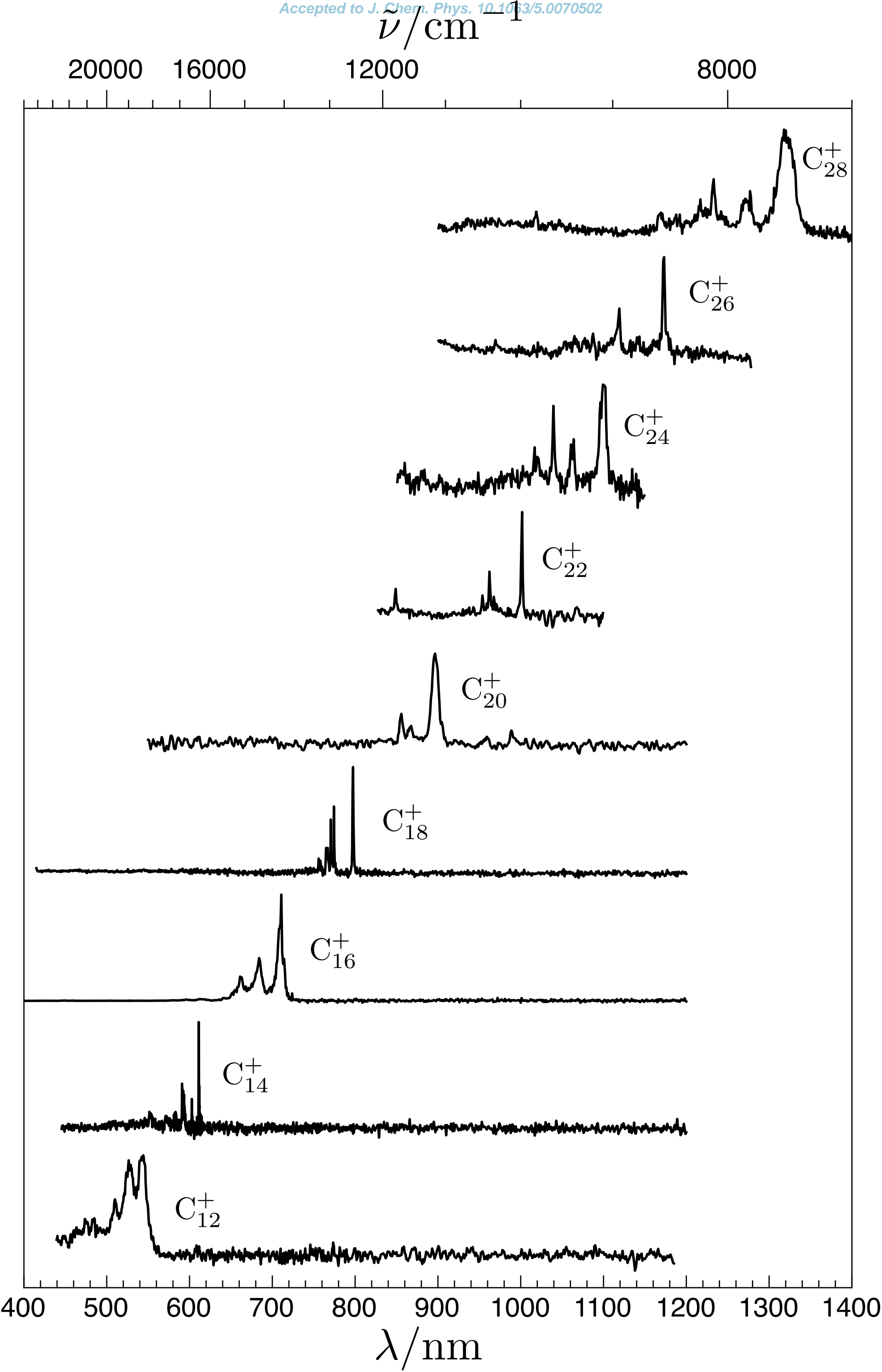
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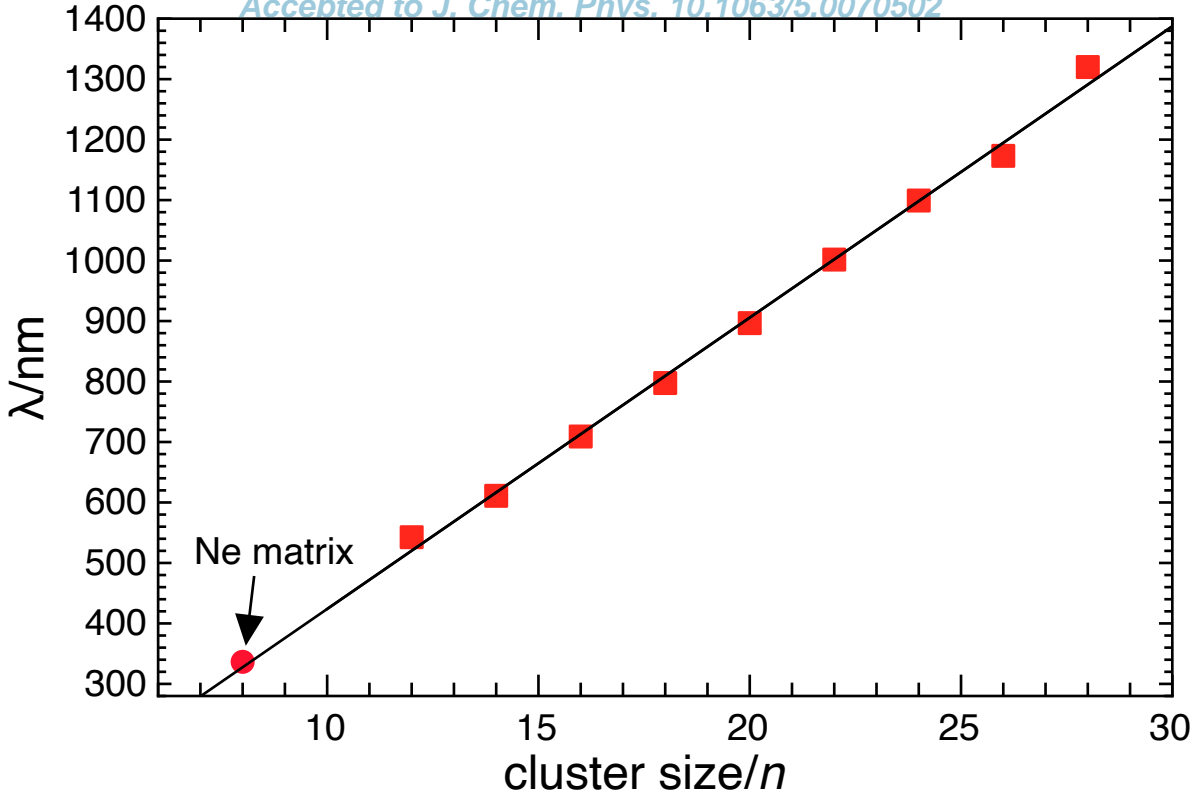
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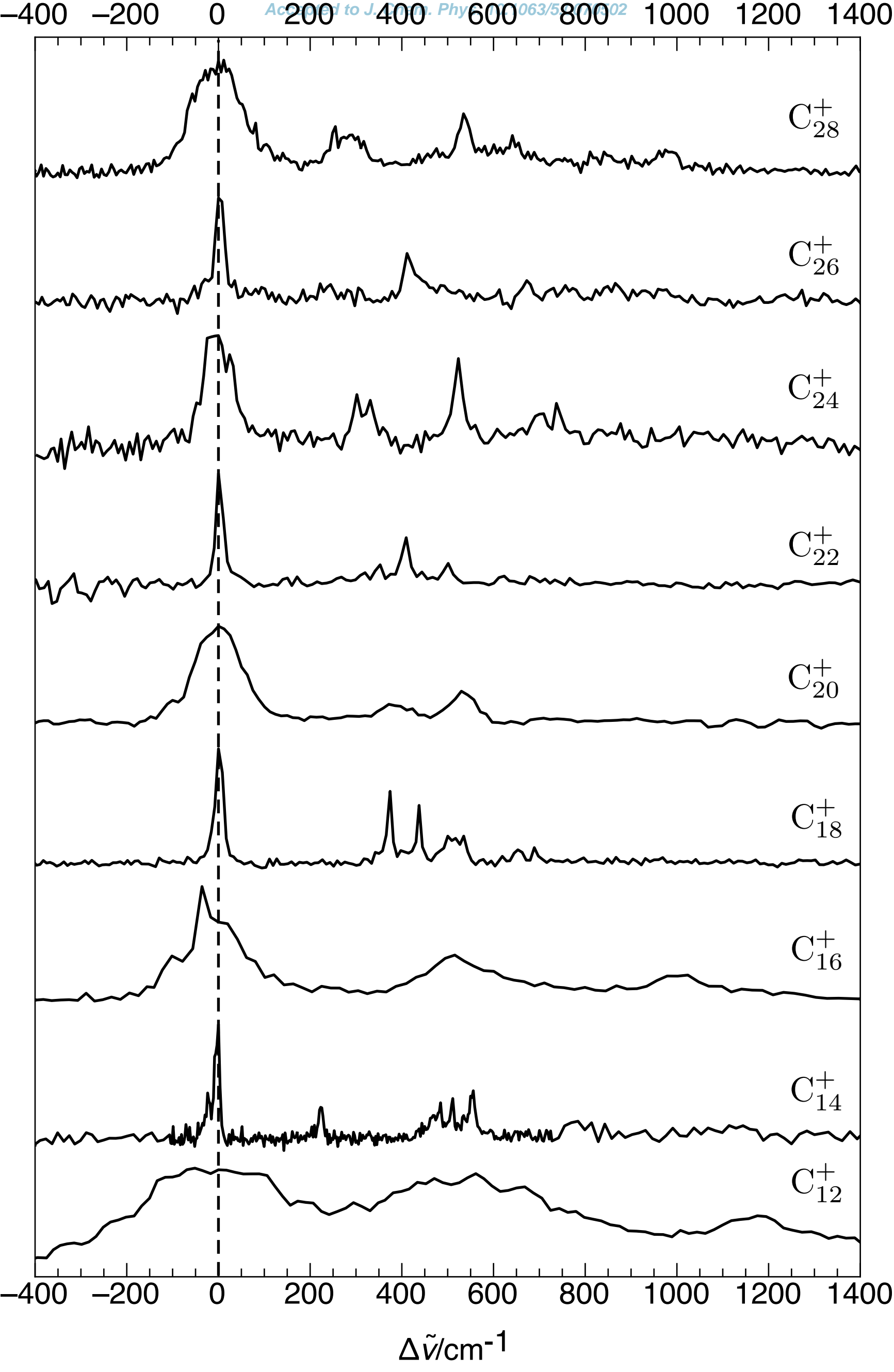


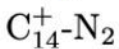
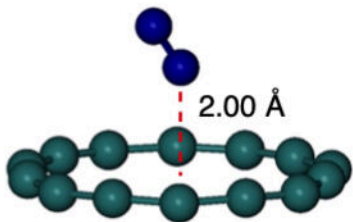




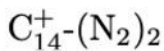
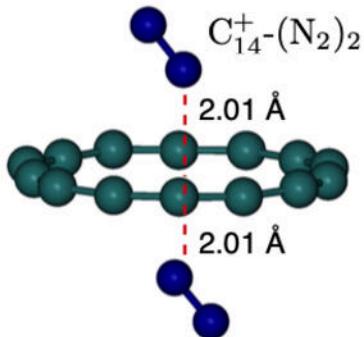








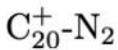
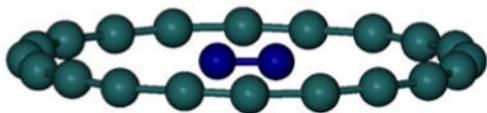
$D_0 \approx 791 \text{ cm}^{-1}$



2.01 Å

2.01 Å

$D_0 \approx 1566 \text{ cm}^{-1}$



$D_0 \approx 984 \text{ cm}^{-1}$

