

# Formation and Characterisation of Polymetallic Rings in vacuo

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## Abstract

Understanding the (dis)assembly mechanisms of large metallocupramolecules is critical in their design, stability and diverse applications. Yet this task is difficult because of the inherent complexity of the structures, with many potential pathways of combining (or separating) the constituent building blocks. Here, we use collision-induced dissociation mass spectrometry to study the disassembly of heterometallic complexes, which have attracted interest for their magnetic and materials properties. Collisional activation leads to the formation of a series of previously unknown smaller ring products and we characterise their geometry using ion mobility. Specifically, studies of the disassembly of two {Cr<sub>x</sub>Cu<sub>2</sub>} hourglass structures (x = 10, 12) and a {Cr<sub>12</sub>Gd<sub>4</sub>} cluster shows the formation of rare heptametallic rings such as {Cr<sub>6</sub>Cu}, {Cr<sub>5</sub>Cu<sub>2</sub>} and {Cr<sub>5</sub>Gd<sub>2</sub>}, as well as other species like {Cr<sub>10</sub>Cu}, {Cr<sub>12</sub>Cu}, {Cr<sub>10</sub>}, {Cr<sub>12</sub>}, and {Cr<sub>6</sub>Gd<sub>2</sub>}. As these rings are non-trivial to synthesize individually, we propose the presented workflow as a means to identify and characterise feasible target molecules. The collision cross section of cyclic products and precursors has a linear correlation with ion mass, a relationship that does not hold for acyclic systems. Thus, ion mobility mass spectrometry can determine whether a candidate polymetallic complex exists as a closed or open structure.

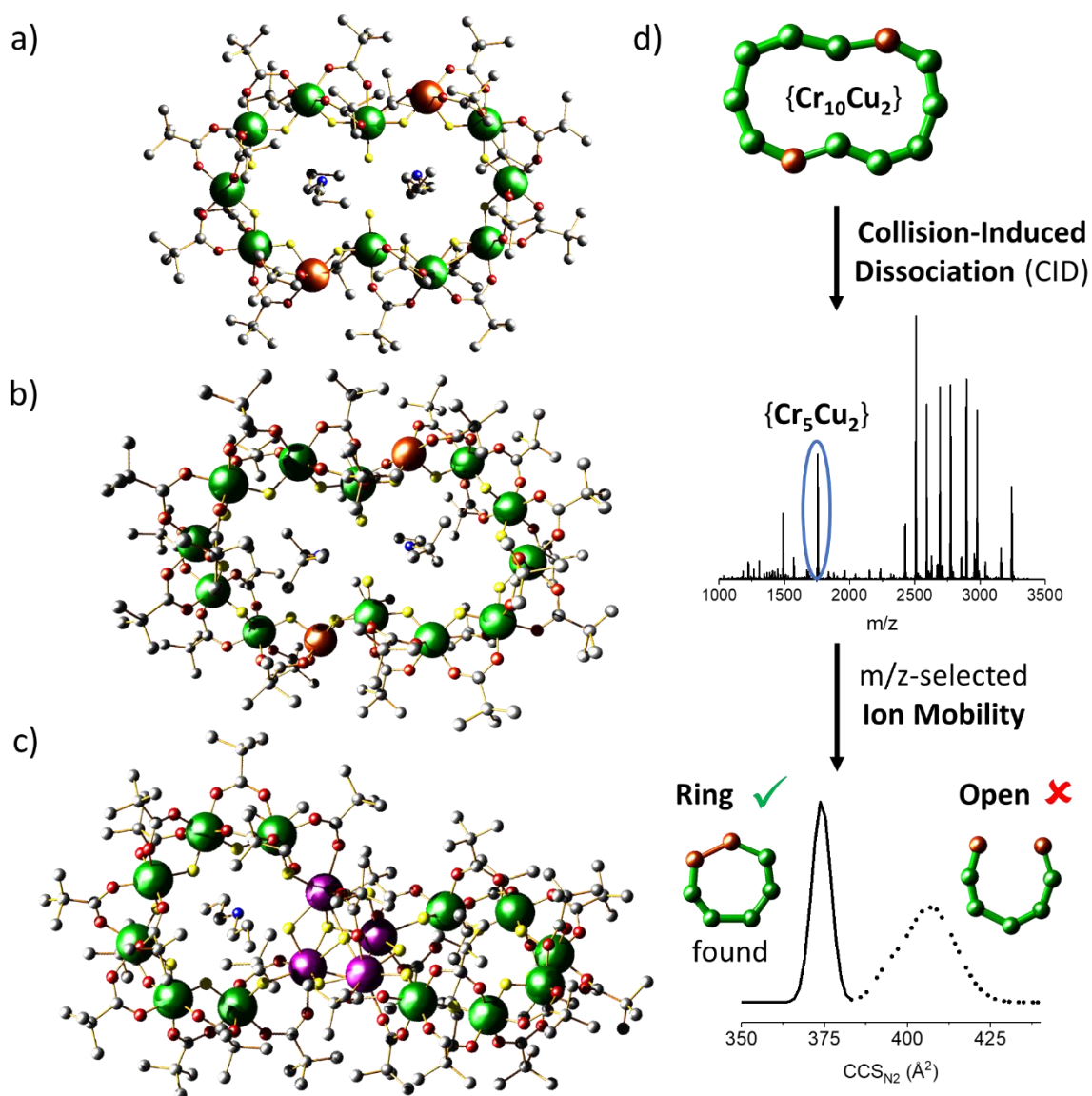
## Introduction

Tandem mass spectrometry (MS<sup>2</sup>) involves isolation of target ions in the gas phase and their subsequent dissociation to smaller ions. The most common fragmentation method is collision-induced dissociation (CID), in which analytes are subjected to collisions with an inert gas at user-defined kinetic energies. The structure of newly formed products as well as non-fragmented precursor ions can be accessed when CID is combined with ion mobility mass spectrometry (IM-MS), which allows the mass as well as size and shape to be measured in the same experiment. Structural information is provided in the form of collision cross sections (CCS), which can be compared to literature data or to theoretical values computed from candidate geometries. Due to the high energies involved in collisional activation, analytes tend to disrupt and undergo major structural change, which becomes visible in the ion mobility spectra of the precursor and/or product ions. Particularly for proteins, the term “collision-induced unfolding” was coined to describe this behaviour similar to denaturation.<sup>1</sup> However, although the disassembly of biomacromolecules is commonly investigated,<sup>2–4</sup> the impact of collisional activation on large synthetic molecules is not well explored.

We have been studying a family of cyclic polymetallic supramolecules that have been proposed as qubits in quantum information processing,<sup>5–8</sup> and more recently have found use as lithographic resists.<sup>9–11</sup> As the interest in these and other metallosupramolecular complexes increases, and similarly the complexity of their structures,<sup>12–14</sup> the synthesis and analysis of their building blocks could aid in rationalising the preference for certain compounds and predict the formation of others.

Perhaps the most prominent example where mass spectrometry was used to find a molecule stable enough to be produced in bulk phase was the discovery of the C<sub>60</sub> buckminsterfullerene by Kroto *et al.* in 1985,<sup>15</sup> which led to the award of the Nobel Prize in Chemistry eleven years later.<sup>16–18</sup> More recently, Cronin and co-workers have used cryospray and ion mobility mass spectrometry to identify polyoxometalate targets for bulk synthesis<sup>19,20</sup> and to unravel their assembly mechanism.<sup>21,22</sup> We have recently applied CID and IM-MS to investigate the disassembly mechanisms and energetics of heterometallic rings and [2]-rotaxanes with the general formula [NH<sub>2</sub>RR']**[Cr<sub>7</sub>MF<sub>8</sub>(O<sub>2</sub>C<sup>t</sup>Bu)<sub>16</sub>]** (M = Mn<sup>II</sup>, Fe<sup>II</sup>, Co<sup>II</sup>, Ni<sup>II</sup>, Cu<sup>II</sup>, Zn<sup>II</sup>, Cd<sup>II</sup>), showing that both the metal M and the R, R' groups can be used to tune the stability and conformational dynamics of these systems.<sup>23</sup>

Here, we investigate the disassembly dynamics of more complex heterometallic systems, two of which have hourglass structures,  $[\text{NH}_2^{\text{n}}\text{Pr}_2]_2[\text{Cr}_{10}\text{Cu}_2\text{F}_{14}(\text{O}_2\text{C}^t\text{Bu})_{22}] = \text{"}\{\text{Cr}_{10}\text{Cu}_2\}\text{"}$  (Figure 1a) and  $[\text{NH}_2^{\text{j}}\text{Pr}_2]_2[\text{Cr}_{12}\text{Cu}_2\text{F}_{16}(\text{O}_2\text{C}^t\text{Bu})_{24}] = \text{"}\{\text{Cr}_{12}\text{Cu}_2\}\text{"}$  (Figure 1b), and a third which involves a lanthanide tetrahedron bound to two  $\{\text{Cr}_6\}$  chains,  $[\text{NH}_2^{\text{n}}\text{Pr}_2]_2[\text{Cr}_{12}\text{Gd}_4\text{F}_{21}(\text{O}_2\text{C}^t\text{Bu})_{29}] = \text{"}\{\text{Cr}_{12}\text{Gd}_4\}\text{"}$  (Figure 1c). We demonstrate how CID, aided by IM-MS, can discriminate different disassembly pathways, which lead to the production of polymetallic rings that have not been made previously *via* solution synthesis (Figure 1d). We suggest that this workflow could therefore inspire new synthetic targets and inform the tools available for the formation of metallosupramolecular compounds. Using the information from the CID-IM-MS measurements, we also propose a workflow for initial topological assignment of potential new polymetallic systems, which applies to structures synthesised in solution as well as those formed in the gas phase.



**Figure 1:** Single crystal X-ray structure of a)  $\{Cr_{10}Cu_2\}$ , b)  $\{Cr_{12}Cu_2\}^{24}$  and c)  $\{Cr_{12}Gd_4\}$ ; Cr: dark green, Cu: brown, Gd: purple, F: yellow, O: red, N: blue, C: grey). Solvent molecules and hydrogen atoms have been removed for clarity. d) CID-IM-MS workflow following nano-electrospray ionisation of the precursor  $\{Cr_{10}Cu_2\}$  from solution, and  $m/z$ -selection of  $[\{Cr_{10}Cu_2\} + Na]^+$ . Upon collisional activation of  $[\{Cr_{10}Cu_2\} + Na]^+$ , several fragment ions were observed with different masses, and their  $CCS_{N2}$  distribution can be extracted as shown for an ion of the type  $\{Cr_5Cu_2\}$ . Different conformations of  $\{Cr_5Cu_2\}$  are possible, e.g. a closed ring (low  $CCS_{N2}$ , narrow  $CCS_{N2}$  distribution) or open forms (high  $CCS_{N2}$ , wide  $CCS_{N2}$  distribution), of which only the closed  $\{Cr_5Cu_2\}$  ring was found.

## Results

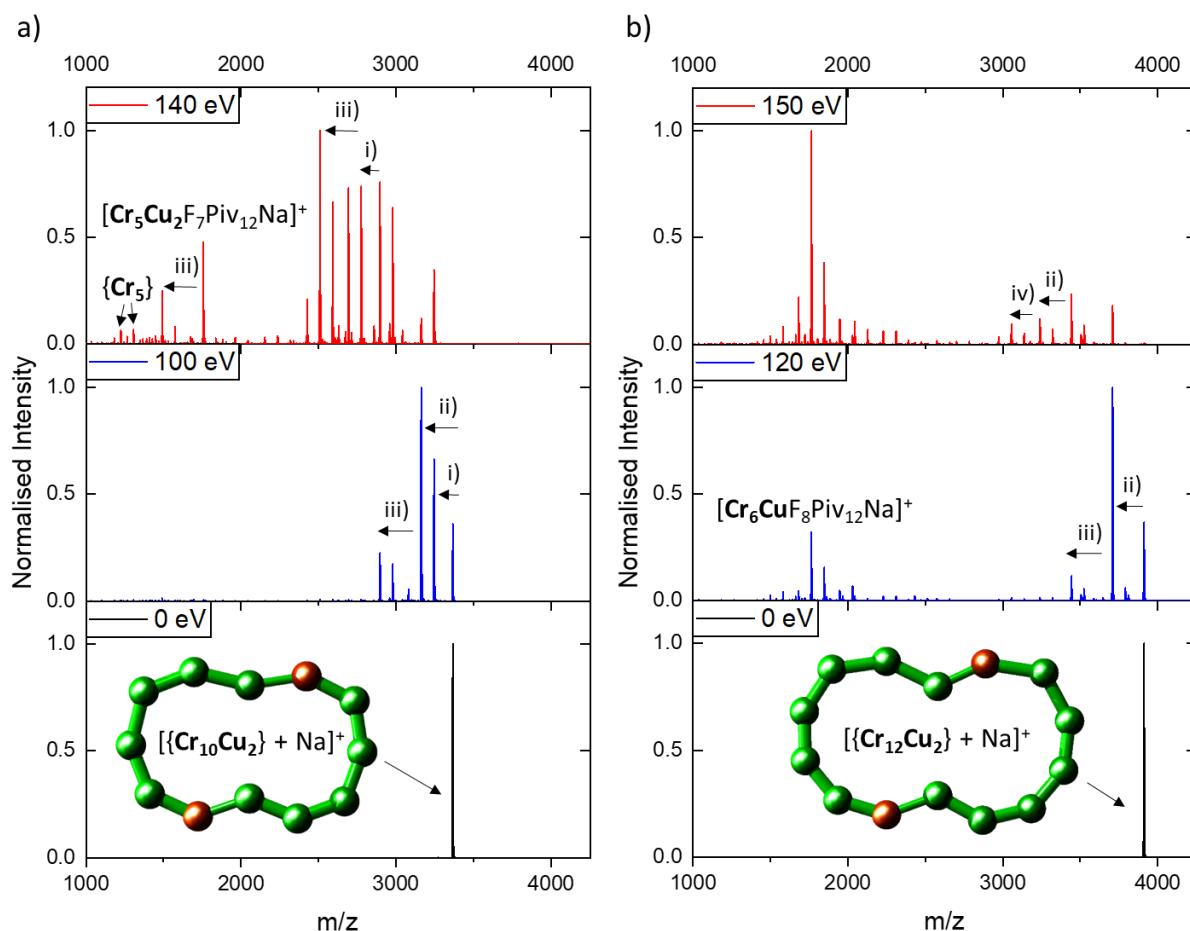
### $\{\text{Cr}_{10}\text{Cu}_2\}$ and $\{\text{Cr}_{12}\text{Cu}_2\}$ Hourglasses

The hourglass structures  $\{\text{Cr}_x\text{Cu}_2\}$  ( $x = 10, 12$ ) consist of two  $\text{Cu}^{\text{II}}$  and  $x$   $\text{Cr}^{\text{III}}$  centres, which are bridged *via* pivalate ligands ( $^-\text{O}_2\text{C}^t\text{Bu} = \text{Piv}^-$ ) on the outside, and fluorides on the inside of the hourglass structure (Figure 1a and b). The coordination environment of the chromium ions involves four pivalates and two *cis*-fluorides with the exception of those located at the hourglass bottleneck, where three pivalates and three *mer*-fluorides are present. These centres are adjacent to the two copper ions, which are penta-coordinated by three pivalates and two fluorides. Additionally, one secondary ammonium cation is located in each half of the hourglass (for  $\{\text{Cr}_{10}\text{Cu}_2\}$ :  $[\text{NH}_2^i\text{Pr}_2]^+$  and for  $\{\text{Cr}_{12}\text{Cu}_2\}$ :  $[\text{NH}_2^i\text{Pr}_2]^+$ ), where it exhibits hydrogen bonds to the fluorides, particularly to the terminal fluorides attached to the two bottleneck chromiums.<sup>24,25</sup> Following optimization of solvent and nano-electrospray ionisation (nESI) source conditions, mass spectra of  $\{\text{Cr}_x\text{Cu}_2\}$  ( $x = 10, 12$ ) were recorded from solutions of sodium iodide in positive mode. Cations of the type  $[\{\text{Cr}_x\text{Cu}_2\} + \text{Na}]^+$  were obtained for both hourglasses (Figures S1, S2).

*Disassembly of  $\{\text{Cr}_{10}\text{Cu}_2\}$  and  $\{\text{Cr}_{12}\text{Cu}_2\}$ .* We isolated the ions  $[\{\text{Cr}_{10}\text{Cu}_2\} + \text{Na}]^+$  and  $[\{\text{Cr}_{12}\text{Cu}_2\} + \text{Na}]^+$  and ramped the energy of collisional activation with nitrogen gas, whilst recording the arrival time distributions (ATD) of both analytes and their products. The ATD of all ions were converted to  $\text{CCS}_{\text{N}_2}$  distributions as a function of collision energy. The tandem mass spectra ( $\text{MS}^2$ ) of both precursor ions show the loss of one secondary ammonium cation along with an anionic ligand, predominantly a pivalate, as the first dissociation step (Figure 2).  $E_{50}$  values, known as a relative measure of ion stability, were determined for both precursor ions and show a slightly higher stability for  $[\{\text{Cr}_{12}\text{Cu}_2\} + \text{Na}]^+$  (Figure S3).

Accurate mass and isotopic distributions were used to assign hourglass fragments in the subsequent dissociation steps at higher collision energies. A variety of dissociation channels occur and ions are observed in two regions of the mass spectrum, one at lower  $m/z$  ( $x = 10$ :  $< 1800$   $m/z$ ;  $x = 12$ :  $< 2200$   $m/z$ ) and one at higher  $m/z$  ( $x = 10$ :  $2500 - 3200$   $m/z$ ;  $x = 12$ :  $2000 - 3700$   $m/z$ ). All of the observed species are singly charged cations, and between these two regions only minor ion populations were found (Figure 2). For  $[\{\text{Cr}_{12}\text{Cu}_2\} + \text{Na}]^+$ , more ions were found in the low mass region compared to  $[\{\text{Cr}_{10}\text{Cu}_2\} + \text{Na}]^+$ , for which the high mass

region clearly dominates. The ratio between the precursor and product ions in both regions depends both on the collision energy (Figure 2) and the instrument used (Figure S4).

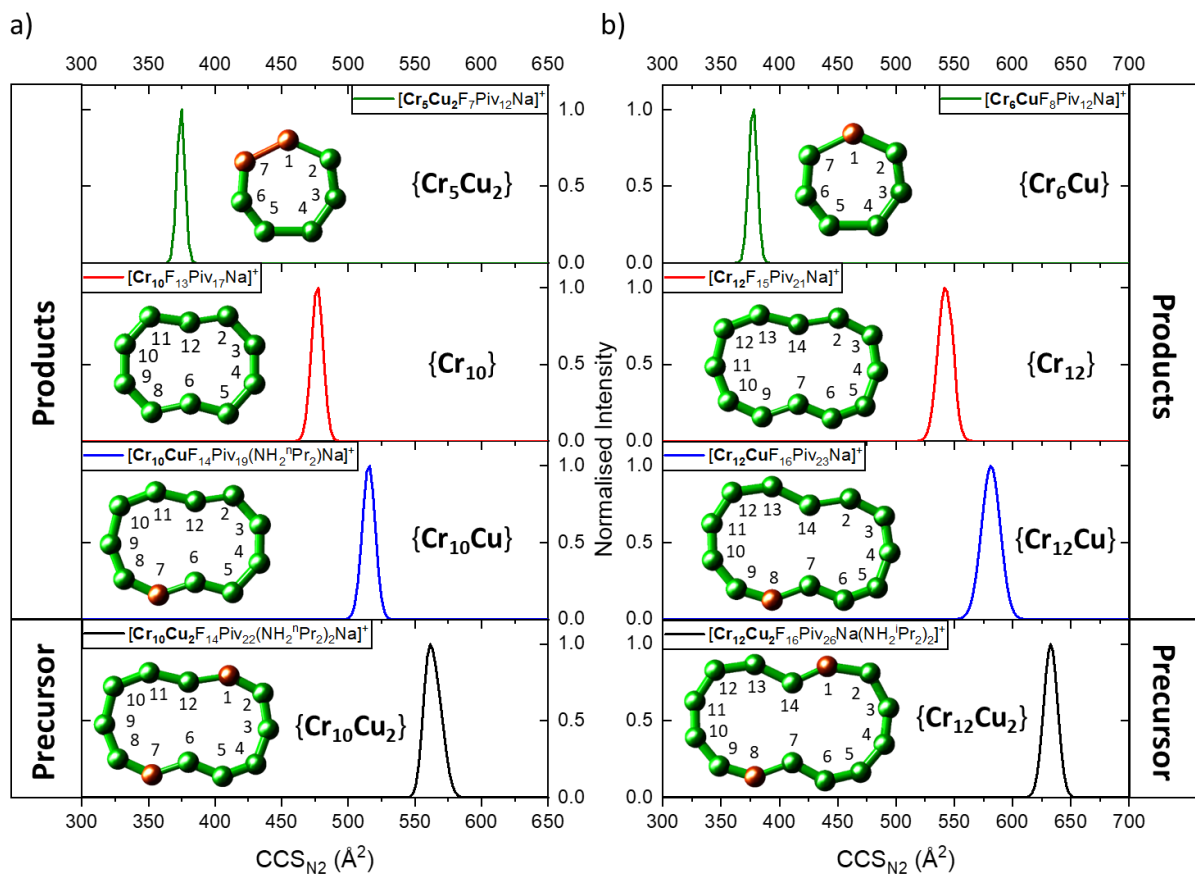


**Figure 2:** Tandem Mass Spectra of a)  $[\{\text{Cr}_{10}\text{Cu}_2\} + \text{Na}]^+$  ( $m/z = 3365$ ) at collision energies of 0 eV, 100 eV and 140 eV and b)  $[\{\text{Cr}_{12}\text{Cu}_2\} + \text{Na}]^+$  ( $m/z = 3910$ ) at collision energies of 0 eV, 120 eV and 140 eV. Occurring fragmentation pathways are labelled: i) –  $[\text{NH}_2^{n/i}\text{Pr}_2]^+$ , –  $\text{F}^-$ ; ii) –  $[\text{NH}_2^{n/i}\text{Pr}_2]^+$ , –  $\text{Piv}^-$ ; iii) –  $\text{Cu}^{\text{II}}$ , –  $2 \text{Piv}^-$  and iv) –  $\text{Cu}^{\text{II}}$ , –  $\text{Piv}^-$ , –  $\text{F}^-$ . The spectra of both ions present two regions, one at lower  $m/z$  and one at higher  $m/z$ . Insets: Schematics of  $\{\text{Cr}_{10}\text{Cu}_2\}$  and  $\{\text{Cr}_{12}\text{Cu}_2\}$  (Cr: green, Cu: brown).

The dominant fragment ions of the low mass region contain seven transition metal ions. From the precursor  $[\{\text{Cr}_{10}\text{Cu}_2\} + \text{Na}]^+$ , it is  $\{\text{Cr}_5\text{Cu}_2\}$  (mainly  $[\text{Cr}_5\text{Cu}_2\text{F}_7\text{Piv}_{12}\text{Na}]^+$  at 1756  $m/z$ ). We also observe in minor amounts  $\{\text{Cr}_5\}$  that likely is the other direct fragment of the  $\{\text{Cr}_{10}\text{Cu}_2\}$  dissociation (Figure 2a top and Supplementary Data Set).  $\{\text{Cr}_5\text{Cu}\}$  was also found in low numbers (mainly  $[\text{Cr}_5\text{CuF}_7\text{Piv}_{10}\text{Na}]^+$  at 1491  $m/z$ ), however this, and other low abundance ions, are presumably secondary fragment from  $\{\text{Cr}_5\text{Cu}_2\}$  as they appear at higher energies than the latter (for  $\{\text{Cr}_5\text{Cu}\}$ : loss of  $\text{Cu}^{\text{II}}$  and two  $\text{Piv}^-$ , Figure 2a). In contrast, the precursor  $[\{\text{Cr}_{12}\text{Cu}_2\} +$

Na)<sup>+</sup> dissociates to {Cr<sub>6</sub>Cu} (mainly [Cr<sub>6</sub>CuF<sub>8</sub>Piv<sub>12</sub>Na]<sup>+</sup> at 1764 *m/z*, Figure 2b). The species in the high mass region follow fragmentation channels where Cu<sup>II</sup> dissociates along with two anionic ligands, before the second ammonium cation and another ligand are lost. This in turn is followed by the dissociation of the second Cu<sup>II</sup> centre, again along with two anionic ligands (Figure 2). Overall, this yields species of the type Cr<sub>x</sub> (x = 10, 12) as the product ions of the hourglass ions [{Cr<sub>x</sub>Cu<sub>2</sub>} + Na]<sup>+</sup>, which was confirmed by their isotopic distributions as illustrated for [Cr<sub>12</sub>F<sub>15</sub>Piv<sub>21</sub>Na]<sup>+</sup> (*m/z* = 3056, Figure S5).

Ion mobility allows us to probe the structure of both precursor and product ions *via* their CCS<sub>N2</sub> distributions (Figure 3). The majority of products exhibit narrow, unimodal conformations, although slightly wider distributions were found for {Cr<sub>12</sub>Cu} and {Cr<sub>12</sub>} fragment ions of [Cr<sub>12</sub>Cu<sub>2</sub> + Na]<sup>+</sup> (Figure 3b) as well as for some {Cr<sub>5</sub>} species formed in the disassembly of [{Cr<sub>10</sub>Cu<sub>2</sub>} + Na]<sup>+</sup> (Supplementary Data Set). The CCS<sub>N2</sub> values of the hourglass precursor ions and the presented products were quantified and compared with data of similar polymetallic rings (Table 1). The data for {Cr<sub>5</sub>Cu<sub>2</sub>} and {Cr<sub>6</sub>Cu} agree well with the CCS<sub>N2</sub> of the seven-membered [Cr<sub>6</sub>MnF<sub>8</sub>Piv<sub>13</sub>]<sup>-</sup> ring, formed from a {Cr<sub>7</sub>Mn} ring *via* CID-MS,<sup>23</sup> and the small difference can be explained with the different number of bulky pivalate ligands ({Cr<sub>5</sub>Cu<sub>2</sub>}, {Cr<sub>6</sub>Cu}: 12 Piv<sup>-</sup>, {Cr<sub>6</sub>Mn}: 13 Piv<sup>-</sup>). This, and their unimodal, narrow distributions clearly suggest a limited conformational flexibility and hence the presence of {Cr<sub>5</sub>Cu<sub>2</sub>} and {Cr<sub>6</sub>Cu} rings, which is distinct from other open {Cr<sub>6</sub>Mn} horseshoe fragments (Table 1).<sup>23</sup> The other products and precursor ions, involving 10 to 14 metal centres, cannot be directly compared to previous data. However, their only slightly increased CCS<sub>N2</sub> values and similarly unimodal and narrow peak shapes indicate cyclic structures as well for these species (Figure 3). For the minor {Cr<sub>5</sub>} fragments, depending on the exact chemical composition of the cations, the conformational landscape is more diverse (Supplementary Data Set).



**Figure 3:**  $CCS_{N_2}$  Distributions of a)  $[Cr_{10}Cu_2 + Na]^+$  ( $m/z = 3365$ ) including selected product ions of the types  $\{Cr_{10}Cu\}$  ( $m/z = 2896$ ),  $\{Cr_{10}\}$  ( $m/z = 2509$ ),  $\{Cr_5Cu_2\}$  ( $m/z = 1756$ ), and b)  $[Cr_{12}Cu_2 + Na]^+$  ( $m/z = 3910$ ) including selected product ions of the types  $\{Cr_{12}Cu\}$  ( $m/z = 3443$ ),  $\{Cr_{12}\}$  ( $m/z = 3056$ ),  $\{Cr_6Cu\}$  ( $m/z = 1764$ ). Data were recorded at collision energies of 0 eV (both precursor ions) as well as 120 eV (fragments of  $[Cr_{10}Cu_2 + Na]^+$ ) and 160 eV (fragments of  $[Cr_{12}Cu_2 + Na]^+$ ), respectively. Insets: Schematics of  $\{Cr_{10}Cu_2\}$ ,  $\{Cr_{12}Cu_2\}$  and their products (Cr: green, Cu: brown).



**Table 1:**  $^{TW}CCS_{N2}$  values of  $\{Cr_{10}Cu_2\} + Na\}^+$ ,  $\{Cr_{12}Cu_2\} + Na\}^+$  and  $\{Cr_{12}Gd_4\} - Piv\}^+$  including selected product ions.  $^{TW}CCS_{N2}$  values of other structures were added from our previous works for comparisons.<sup>23,26</sup> Data shown represents different collision energies, which can lead to small deviations. For both types of  $\{Cr_6Mn\}$  horseshoes (i.e. open chains), the broad  $^{TW}CCS_{N2}$  distributions were fitted with two Gaussian distributions, and their two maxima are noted in the table.

Source	Cluster Type	Composition	$m/z$	$^{TW}CCS_{N2}$ ( $\text{\AA}^2$ )
Disassembly of $\{Cr_{10}Cu_2\} + Na\}^+$	$\{Cr_{10}Cu_2\}$	$[Cr_{10}Cu_2F_{14}Piv_{22}(NH_2^nPr_2)_2Na]^+$	3365	$566.5 \pm 0.6$
	$\{Cr_{10}Cu\}$	$[Cr_{10}CuF_{14}Piv_{19}(NH_2^nPr_2)Na]^+$	2896	$514.1 \pm 1.3$
	$\{Cr_{10}\}$	$[Cr_{10}F_{13}Piv_{17}Na]^+$	2509	$476.0 \pm 0.8$
	$\{Cr_5Cu_2\}$	$[Cr_5Cu_2F_7Piv_{12}Na]^+$	1756	$374.2 \pm 0.2$
Disassembly of $\{Cr_{12}Cu_2\} + Na\}^+$	$\{Cr_{12}Cu_2\}$	$[Cr_{12}Cu_2F_{16}Piv_{26}(NH_2^iPr_2)_2Na]^+$	3910	$633.2 \pm 0.6$
	$\{Cr_{12}Cu\}$	$[Cr_{12}CuF_{16}Piv_{23}(NH_2^iPr_2)Na]^+$	3443	$580.2 \pm 1.3$
	$\{Cr_{12}\}$	$[Cr_{12}F_{15}Piv_{21}Na]^+$	3056	$541.9 \pm 0.3$
	$\{Cr_6Cu\}$	$[Cr_6CuF_8Piv_{12}Na]^+$	1764	$376.3 \pm 1.0$
Disassembly of $\{Cr_{12}Gd_4\} - Piv\}^+$	$\{Cr_{12}Gd_4\}$	$[Cr_{12}Gd_4F_{21}Piv_{28}(NH_2^nPr_2)_2]^+$	4688	$703.5 \pm 6.3$
	$\{Cr_6Gd_2\}$	$[Cr_6Gd_2F_8Piv_{16}(NH_2^nPr_2)]^+$	2499	$446.1 \pm 0.1$
	$\{Cr_5Gd_2\}$	$[Cr_5Gd_2F_8Piv_{12}]^+$	1940	$377.4 \pm 0.7$
Literature Values	$\{Cr_8\}$	$[Cr_8F_8Piv_{16}Na]^+$	2208	$430.0 \pm 0.8$
	$\{Cr_7Cu\}$	$[Cr_7CuF_8Piv_{16}]^-$	2197	$432.7 \pm 2.0$
	$\{Cr_6Mn\}$	$[Cr_6MnF_7Piv_{14}]^-$	1915	$401.5 \pm 0.1$
	Ring	$[Cr_6MnF_8Piv_{13}]^-$	1833	$394.2 \pm 0.1$
	$\{Cr_6Mn\}$ Open	$[Cr_6MnF_7Piv_{14}]^-$	1915	$429.3 \pm 0.1$ and $444.3 \pm 0.1$
	Horseshoe	$[Cr_6MnF_8Piv_{13}]^-$	1833	$413.4 \pm 0.1$ and $426.9 \pm 0.1$

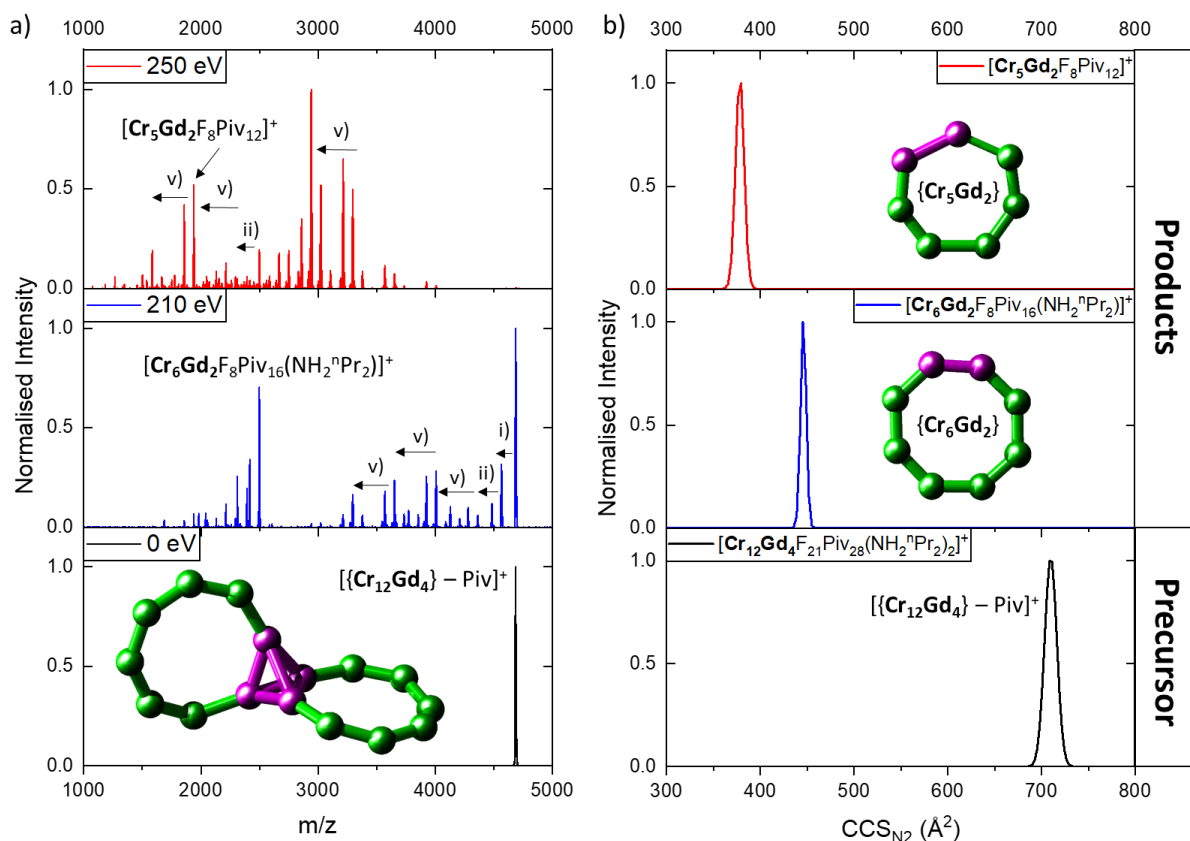
## **{Cr<sub>12</sub>Gd<sub>4</sub>} Cluster**

We further investigated whether collision-induced disassembly would also lead to polymetallic rings when activating a different type of precursor. We chose the lanthanide cluster **{Cr<sub>12</sub>Gd<sub>4</sub>}** (Figure 1c), which consists of a tetrahedral **{Gd<sub>4</sub>F<sub>7</sub>Piv}** cage, in which the fluorides and one pivalate bridge the four **Gd<sup>III</sup>** centres. Each of the octa-coordinated gadolinium ions is attached to a terminus of one of two **{Cr<sub>6</sub>}** chains, and a **[NH<sub>2</sub><sup>n</sup>Pr<sub>2</sub>]<sup>+</sup>** cation is located at each ring centre.<sup>27</sup> The chromium atoms are each connected *via* two pivalate ligands and one fluoride, as in both **{Cr<sub>x</sub>Cu<sub>2</sub>}** hourglasses.

The mass spectrum of **{Cr<sub>12</sub>Gd<sub>4</sub>}**, sprayed from methanol in the presence of sodium iodide using nESI, yielded only a small number of largely intact analyte ions (Figure S6), and we assigned the dominant one to **[{Cr<sub>12</sub>Gd<sub>4</sub>} – Piv]<sup>+</sup>** (*m/z* = 4688, Figure S7 for Isotopic Distribution). Several other polymetallic species were observed in the mass spectrum, including **{Cr<sub>6</sub>Gd}** ions, which are possibly produced by decomposition reactions with the solvent. Using ion mobility, the **{Cr<sub>6</sub>Gd}** clusters were identified as heptametallic rings (CCS<sub>N2</sub> values in Table S1 including discussion of their disassembly; Supplementary Data Set), which are isostructural to a **{Cr<sub>6</sub>Ce}** ring reported previously, but only synthesised in very low yield (see Discussion).<sup>27</sup>

*Disassembly of {Cr<sub>12</sub>Gd<sub>4</sub>}*. We used the tandem mass spectrometry ion mobility workflow (Figure 1d) to study **[{Cr<sub>12</sub>Gd<sub>4</sub>} – Piv]<sup>+</sup>**. Similar to the studies of **[{Cr<sub>x</sub>Cu<sub>2</sub>} + Na]<sup>+</sup>**, the tandem mass spectra show the loss of **[NH<sub>2</sub><sup>n</sup>Pr<sub>2</sub>]<sup>+</sup>** along with one fluoride as the main fragmentation channel, although at higher energies (Figure 4a). The *E*<sub>50</sub> value of **[{Cr<sub>12</sub>Gd<sub>4</sub>} – Piv]<sup>+</sup>** was determined and is more than 50% higher than those of the hourglass ions (Figure S8). Upon increasing the energy further (Figure 4a centre), two main regions were found in the tandem mass spectra: one at 1800 - 2500 *m/z* and the other at 3200 – 4500 *m/z*, however the separation between regions is blurred at higher collision energies (Figure 4a top). All species also occur as singly charged cations. In the lower mass region, the cluster type **{Cr<sub>6</sub>Gd<sub>2</sub>}** was found as the main product (mainly **[Cr<sub>6</sub>Gd<sub>2</sub>F<sub>8</sub>Piv<sub>16</sub>(NH<sub>2</sub><sup>n</sup>Pr<sub>2</sub>)]<sup>+</sup>** at *m/z* = 2499, Figure S9 for Isotopic Distribution). This cluster is one half of the **{Cr<sub>12</sub>Gd<sub>4</sub>}** precursor and we previously reported the solid-state structure of a cyclic **{Cr<sub>6</sub>Y<sub>2</sub>}** complex, albeit in low yield.<sup>14</sup> The **{Cr<sub>6</sub>Gd<sub>2</sub>}** ions undergo further fragmentation at even higher energies, with losses of **[NH<sub>2</sub><sup>n</sup>Pr<sub>2</sub>]<sup>+</sup>** or **Cr<sup>III</sup>**, along with anionic ligands (mainly pivalates, Figure 4a bottom). This for example leads to ions

of the type  $\{\text{Cr}_5\text{Gd}_2\}$  (e.g.  $[\text{Cr}_5\text{Gd}_2\text{F}_8\text{Piv}_{12}]^+$  at  $m/z = 1940$ ). In contrast, the high mass region shows the stepwise disruption of the  $\{\text{Cr}_6\}$  chains with losses of  $[\text{NH}_2^n\text{Pr}_2]^+$  along with  $\text{Piv}^-$ , followed by several dissociations of  $\{\text{Cr}(\text{Piv})_3\}$  units (Figure 4a centre). This infers that the  $\{\text{Gd}_4\}$  tetrahedron remains intact.



**Figure 4:** a) Tandem Mass Spectra of  $[\{\text{Cr}_{12}\text{Gd}_4\} - \text{Piv}]^+$  at collision energies of 0 eV, 210 eV and 250 eV. Occurring fragmentation pathways are labelled: i) –  $[\text{NH}_2^n\text{Pr}_2]^+$ , –  $\text{F}^-$ ; ii) –  $[\text{NH}_2^n\text{Pr}_2]^+$ , –  $\text{Piv}^-$  and v) –  $\text{Cr}^{\text{III}}$ , – 3  $\text{Piv}^-$ . The spectrum at 210 eV shows two regions, one at lower  $m/z$  and one at higher  $m/z$ . Minor contaminant features were observed which were  $m/z$ -selected in the quadrupole along with  $[\{\text{Cr}_{12}\text{Gd}_4\} - \text{Piv}]^+$ . b)  $\text{CCS}_{\text{N}_2}$  Distributions of  $[\{\text{Cr}_{12}\text{Gd}_4\} - \text{Piv}]^+$  ( $m/z = 4688$ ) and the fragment ions  $\{\text{Cr}_6\text{Gd}_2\}$  ( $m/z = 2499$ ) and  $\{\text{Cr}_5\text{Gd}_2\}$  ( $m/z = 1940$ ). Data were recorded at collision energies of 0 eV (precursor), 130 eV for  $\{\text{Cr}_6\text{Gd}_2\}$  and 190 eV for  $\{\text{Cr}_5\text{Gd}_2\}$ , respectively. Insets: Schematics of  $\{\text{Cr}_{12}\text{Gd}_4\}$  and the two products  $\{\text{Cr}_x\text{Gd}_2\}$  ( $x = 5, 6$ ; Cr: green, Gd: purple).

The structure of  $[\{\text{Cr}_{12}\text{Gd}_4\} - \text{Piv}]^+$  and fragment ions was investigated *via* their  $\text{CCS}_{\text{N}_2}$  distributions (Figure 4b for selected ions). The majority of the ions exhibit narrow, unimodal conformations (Supplementary Data Set). We quantified the absolute  $\text{CCS}_{\text{N}_2}$  values of the

precursor  $[\{\text{Cr}_{12}\text{Gd}_4\} - \text{Piv}]^+$  and products of the type  $\{\text{Cr}_6\text{Gd}_2\}$  and  $\{\text{Cr}_5\text{Gd}_2\}$  (Table 1). The  $\text{CCS}_{N_2}$  value of  $\{\text{Cr}_6\text{Gd}_2\}$  agrees well with a related  $\{\text{Cr}_7\text{Cu}\}$  ring,<sup>23</sup> while the value for  $\{\text{Cr}_5\text{Gd}_2\}$  is similar to those of the seven-membered hourglass products  $\{\text{Cr}_5\text{Cu}_2\}$  and  $\{\text{Cr}_6\text{Cu}\}$ , as well as to the  $[\text{Cr}_6\text{MnF}_8\text{Piv}_{13}]^-$  ring.<sup>23</sup> These comparisons suggest cyclic structures for the complexes of the type  $\{\text{Cr}_6\text{Gd}_2\}$  and  $\{\text{Cr}_5\text{Gd}_2\}$ .

## Discussion

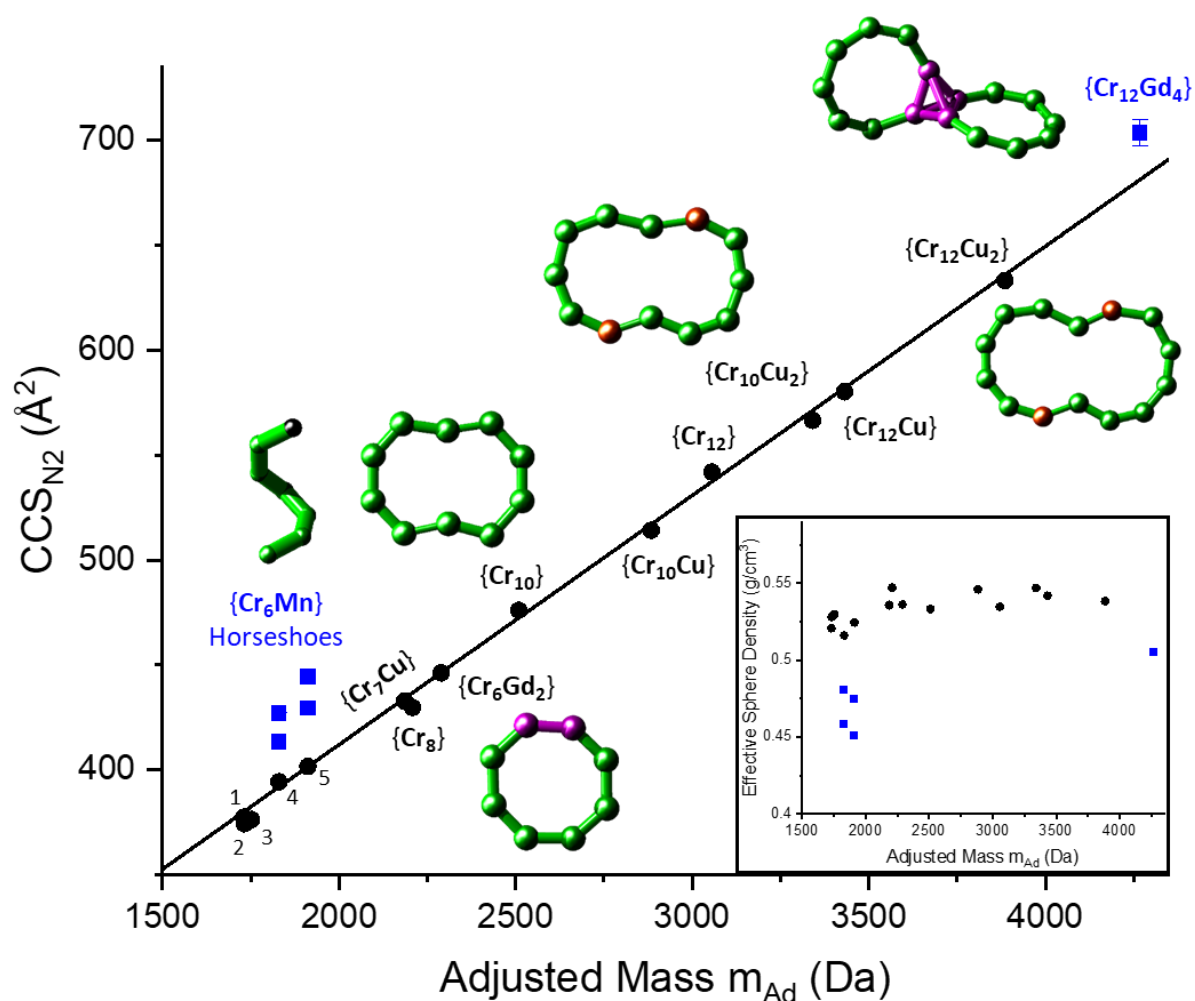
The first major finding from this work is the formation of cyclic products from the collision-induced dissociation of the  $\{\text{Cr}_x\text{Cu}_2\}$  hourglasses ( $x = 10, 12$ ) and from  $\{\text{Cr}_{12}\text{Gd}_4\}$ . The observation that polymetallic rings are formed from all three precursors suggests that this workflow can be used to produce smaller rings of varying sizes, including new types (see below), as long as appropriate larger precursors are available. Remarkable is that these rings form after collisions on a millisecond timescale, which suggests that both rearrangements and new bond formations occur fast. Also noteworthy is the occurrence of gaps in the tandem mass spectra of all three parent ions (Figures 2, 4a), which suggests that potential products in these regions are significantly less stable than those obtained at both lower and higher masses, akin to the magic numbers seen in the spectra for monatomic and molecular cluster ions.<sup>15,28–32</sup> In our case, the primary and presumably the most stable products from the precursor ions are  $\{\text{Cr}_5\text{Cu}_2\}$ ,  $\{\text{Cr}_6\text{Cu}\}$  and  $\{\text{Cr}_6\text{Gd}_2\}$ , whereas others such as  $\{\text{Cr}_5\text{Cu}\}$  and  $\{\text{Cr}_5\text{Gd}_2\}$  are secondary products from these fragments.

Particularly striking is the formation of heptametallic rings  $\{\text{Cr}_5\text{Cu}_2\}$ ,  $\{\text{Cr}_6\text{Cu}\}$  and  $\{\text{Cr}_5\text{Gd}_2\}$ . In the family of polymetallic chromium rings,<sup>33</sup> the only seven-metal ring synthesized to date is  $\{\text{Cr}_6\text{Ce}\}$ , which only forms as a very minor side product,<sup>27</sup> and this is one of three structurally characterised heptametallic rings known.<sup>34,35</sup> Given the sparsity of known compounds it might be argued that these products are unlikely, however in the synthesis of heterometallic chromium rings the difficulty is avoiding the formation of  $[\text{Cr}_8\text{F}_8\text{Piv}_{16}]$ , which is achieved by adding a cationic template that occupies the centre of the cavity and leads to an anionic heterometallic ring.<sup>25</sup> This allows formation of larger rings by choice of larger templates, but for a seven-metal ring there is insufficient space for any organic template. Applying the presented CID-MS approach on these larger rings therefore offers a new route for the formation of heptametallic and smaller rings. We have also previously confirmed the stability

of a heptametallic  $\{\text{Cr}_6\text{Mn}\}$  ring formed by CID-MS, supporting the general presence of heptametallic rings in the gas phase.<sup>23</sup>

The discussed approach can inspire new supramolecular chemistry outside the gas phase. In some cases, rings similar to those produced by CID-MS have been previously synthesised and characterised. For example, several  $\{\text{Cr}_{10}\}$  rings have been reported by our group, however these involved different bridging ligands.<sup>36–38</sup> We have also structurally characterised a  $\{\text{Cr}_6\text{Y}_2\}$  ring of the formula  $[\text{Cr}_6\text{Y}_2\text{F}_8\text{Piv}_{17}(\text{NH}_2\text{Et}_2)(\text{H}_2\text{O})]$ , which is very similar to the ion  $[\text{Cr}_6\text{Gd}_2\text{F}_8\text{Piv}_{16}(\text{NH}_2^{\text{n}}\text{Pr}_2)]^+$  observed upon fragmentation of  $\{\text{Cr}_{12}\text{Gd}_4\}$ . This  $\{\text{Cr}_6\text{Y}_2\}$  was formed only in low yield as the solution synthesis leads to  $\{\text{Cr}_{12}\text{Ln}_4\}$  cages as the major product.<sup>27</sup>

Our second major observation is that ion mobility can be used to identify whether a given polymetallic complex of this family is cyclic or not, and this applies both to structures formed in solution and those *via* CID-MS in the gas phase. The two main advantages here, compared to other bulk phase methods, are small measurement times and low amounts of samples needed. We plotted the  $\text{CCS}_{\text{N}_2}$  of all species in Table 1 against their adjusted masses (Figure 5). This correlation represents the atom density in a given ion,<sup>39–41</sup> which is highest for ions with small CCS and large mass. Previously, Bleiholder *et al.* tracked the self-assembly of peptides and, using an analogous relationship, distinguished between the formation of larger globular peptides and  $\beta$ -sheets relevant for amyloid fibril formation.<sup>40</sup> Later our group showed that the  $\text{CCS}/m$  slope of unfolded and intrinsically disordered proteins is significantly higher (lower density) than for native proteins.<sup>41</sup> In the case of cyclic species from this and previous works, we observe a linear correlation, which does not hold for the acyclic  $\{\text{Cr}_{12}\text{Gd}_4\}$  cluster and the extended  $\{\text{Cr}_6\text{Mn}\}$  horseshoes.<sup>23</sup> These are located above the line as their estimated spherical densities (ESD) are lower than for the polymetallic cyclic structures (Figure 5 Inset). ESD were derived from  $\text{CCS}_{\text{N}_2}$  values, following previously published calculations based on the assumption of a spherical ion,<sup>39</sup> and cannot be directly compared to the macromolecular density in the crystal structures (where available). The latter is typically higher ( $1.2 - 1.4 \text{ g/cm}^3$ ) and represents the packing density of the molecules in the crystal lattice (together with solvent molecules), whereas the ESD discussed here inform on how dense the atoms are in the gas phase ion.



**Figure 5:** Correlation between  $CCS_{N_2}$  and adjusted mass  $m_{ad}$  for all species in Table 1. The cyclic complexes show a linear correlation, which does not hold for the acyclic ions ( $\{Cr_{12}Gd_4\}$  and  $\{Cr_6Mn\}$  horseshoes). 1:  $\{Cr_5Gd_2\}$ ; 2:  $\{Cr_5Cu_2\}$ ; 3:  $\{Cr_6Cu\}$ ; 4, 5:  $\{Cr_6Mn\}$  rings. The mass was adjusted so that every metal centre is accounted for with the mass of chromium (52 Da). This model is reasonable as the nature of the metal in the polymetallic complex has a low overall impact on the  $CCS_{N_2}$ , although some have high masses e.g. in the case of gadolinium. Inset: Schematics of selected cyclic and acyclic complexes (Cr: green, Cu: brown, Gd: purple, Mn: black). Effective sphere density (ESD) derived from  $CCS_{N_2}$  values vs adjusted mass (black circles: cyclic, blue squares: acyclic), showing that the acyclic  $\{Cr_6Mn\}$  horseshoes and  $\{Cr_{12}Gd_4\}$  have a lower ESD than the cyclic species.

*Disassembly of  $\{Cr_{10}Cu_2\}$  and  $\{Cr_{12}Cu_2\}$ .* The first dissociation step of both hourglass ions  $[\{Cr_xCu_2\} + Na]^+$  ( $x = 10, 12$ ) is the loss of the secondary ammonium cation, along with an anionic ligand, predominantly a pivalate. This is probably because the cation is the only non-covalently bound species in the system. As the primary fragmentation pathway is the same

for both  $\{Cr_xCu_2\} + Na\}^+$  ions, their  $E_{50}$  values are also similar. The same dissociation channel was observed for the heterometallic rotaxanes  $[NH_2RR']\{Cr_7MF_8Piv_{16}\}$ , where R and R' include bulky phenyl and *tert*-butyl groups.<sup>23</sup> The secondary ammonium cations are smaller here, so the relatively ease of dissociation from the hourglass ions is not surprising.

The further disassembly can be rationalised by considering the sites in the hourglass structure that are most prone to collision-induced dissociation. These are the two  $Cu^{II}$  centres and their ligands, because of the lower oxidation state ( $Cu^{II}$  vs.  $Cr^{III}$ ), and hence a weaker electrostatic attraction between ligand and metal, as well as the coordination number ( $Cu^{II}$ : 5,  $Cr^{III}$ : 6, Figure 1a, b). It seems likely that both the initially lost ammonium cation and pivalate are adjacent to the same  $Cu^{II}$  centre. This would weaken the hourglass scaffold further in this region, which is the reason for its disruption in the second dissociation step (rather than the loss of the second ammonium cation). As discussed above, the hourglass disruption occurs *via* one of two pathways, with one of them yielding ions in the low  $m/z$ -region and the other one at higher  $m/z$ .

Fragmentation associated with higher mass loss leads to rings of the type  $\{Cr_5Cu_2\}$  (from  $\{Cr_{10}Cu_2\} + Na\}^+$ ) and  $\{Cr_6Cu\}$  (from  $\{Cr_{12}Cu_2\} + Na\}^+$ , Figures 2, 3 and Table 1). In the disassembly of  $\{Cr_{12}Cu_2\}$ , the dissociation occurs at the hourglass bottleneck, between **Cu-1** and **Cr-14** as well as **Cr-7** and **Cu-8** (Schematics in Figure 3b). This is likely the favoured site as  $Cu^{II}$  is far more reactive than  $Cr^{III}$ . As a result, the hourglass predominantly dissociates symmetrically to  $\{Cr_6Cu\}$ , which rearranges to a ring structure likely *via* a new connection between one **Cu-1** and **Cr-7**. For the fragmentation of  $\{Cr_{10}Cu_2\}$ , this same mechanism would lead to a six-membered  $\{Cr_5Cu\}$  ring (connection between **Cu-1** and **Cr-6**, Schematics in Figure 3a). Interestingly this ring, characterised by ion mobility, is only seen at higher energies, suggesting that it is a secondary fragment (Figure 2a and Supplementary Data Set). By contrast, we predominantly observe  $\{Cr_5Cu_2\}$  products due to an asymmetric hourglass disruption (presumably between **Cu-7** and the **Cr-8**). Here three bonds need to be broken instead of two and this preference may be due to the stability given by the formation of a seven-membered ring, which implies it is more stable than the alternative six-membered  $\{Cr_5Cu\}$  ring.

No ammonium cation is present in the dominant  $\{\text{Cr}_5\text{Cu}_2\}$  or  $\{\text{Cr}_6\text{Cu}\}$  ring products, which indicates that these are formed from the part of the hourglass where the ammonium cations and  $\text{Piv}^-$  are lost in the primary fragmentation step. This is possibly due to insufficient space in the central cavity of a seven-metal ring to bind an ammonium cation. Instead,  $\text{Na}^+$  was found in all the heptametallic rings observed, where it is likely located in the centre. We have previously found that  $\text{Na}^+$  is too small for a good fit for the cavity of an octametallic ring, both in bulk and gas phase,<sup>26,42</sup> and might hence be better suited for seven-metal rings. The formation of  $\{\text{Cr}_5\text{Cu}_2\}$  from  $\{\text{Cr}_{10}\text{Cu}_2\}$ , likely occurring in a concerted step, implies that  $\{\text{Cr}_5\}$  is also formed. As discussed above, we have a low number of  $\{\text{Cr}_5\}$  ions (Figure 2a top), which is likely due to the relatively higher stability of heptametallic rings and the preference of the charge-carrier  $\text{Na}^+$  to remain with  $\{\text{Cr}_5\text{Cu}_2\}$ , which complicates the detection of  $\{\text{Cr}_5\}$  cations. Our data suggests that  $\{\text{Cr}_5\}$  can occur in different conformations and/or topologies, potentially either as chains and/or rings (Supplementary Data Set).

The mechanism associated with smaller mass losses shows the stepwise disruption of the hourglasses, starting with the loss of one  $\text{Cu}^{\text{II}}$  centre along with two ligands. The formed  $\{\text{Cr}_{10}\text{Cu}\}$  and  $\{\text{Cr}_{12}\text{Cu}\}$  products are likely also rings due to their narrow and unimodal  $\text{CCS}_{\text{N}_2}$  distributions (Figure 3, Table 1), and require new connections, likely between **Cr-2** and **Cr-12** for  $\{\text{Cr}_{10}\text{Cu}\}$  and **Cr-2** and **Cr-14** for  $\{\text{Cr}_{12}\text{Cu}\}$ , respectively. The  $\{\text{Cr}_{12}\text{Cu}\}$  distribution is significantly wider than for  $\{\text{Cr}_{10}\text{Cu}\}$ , indicating a higher conformational flexibility. The same qualitative trend was observed in the next fragmentation step, where the second secondary ammonium cation is lost, predominantly along with a pivalate. After that, the second  $\text{Cu}^{\text{II}}$  centre dissociates along with two anionic ligands, leading to ions of the type  $\{\text{Cr}_{10}\}$  and  $\{\text{Cr}_{12}\}$ . Here both ions exhibit narrow  $\text{CCS}_{\text{N}_2}$  distributions, once more suggesting cyclic structures, presumably due to newly formed connections between **Cr-6** and **Cr-8** for  $\{\text{Cr}_{10}\}$  and **Cr-7** and **Cr-9** for  $\{\text{Cr}_{12}\}$ .

*Disassembly of  $\{\text{Cr}_{12}\text{Gd}_4\}$ .* If similar disassembly mechanisms take place for  $[\{\text{Cr}_{12}\text{Gd}_4\} - \text{Piv}]^+$  as for the hourglass ions, we would also expect smaller ring structures, of the type  $\{\text{Cr}_6\text{Gd}\}$  (as discussed above and in the SI) or  $\{\text{Cr}_6\text{Gd}_2\}$ . For the second route involving low mass loss, we would expect the stepwise disruption of the  $\{\text{Cr}_6\}$  chains, again followed by rearrangements to cyclic structures, as these are more accessible for gas collisions and less strongly bound than the  $\{\text{Gd}_4\}$  unit.



The fragmentation of the ion  $[\{\text{Cr}_{12}\text{Gd}_4\} - \text{Piv}]^+$  follows similar routes as  $[\{\text{Cr}_x\text{Cu}_2\} + \text{Na}]^+$  ( $x = 10, 12$ ) and confirms our hypothesis that the mechanisms observed for the hourglass ions can be transferred to other polymetallic complexes. Here, after the loss of one secondary ammonium cation along with a fluoride, the disassembly associated with higher mass loss leads to  $\{\text{Cr}_6\text{Gd}_2\}$  and subsequently to  $\{\text{Cr}_5\text{Gd}_2\}$  complexes, for which the comparisons with previously reported  $\text{CCS}_{\text{N}_2}$  values clearly indicate ring structures.

The second disassembly mechanism of the  $\{\text{Cr}_{12}\text{Gd}_4\}$  cluster shows the loss of the second  $[\text{NH}_2^+\text{Pr}_2]^+$  and a pivalate, followed by multiple dissociation steps in which  $\{\text{CrPiv}_3\}$  is lost. Here the loss of five  $\{\text{CrPiv}_3\}$  units, occurring as the most abundant pathways, exceeds the number of pivalates present in a  $\{\text{Cr}_6\}$  chain and suggests either major rearrangements, or the disruption of the second  $\{\text{Cr}_6\}$  unit before the complete dissociation of the first. The  $\text{CCS}_{\text{N}_2}$  distributions of the occurring products indicate more flexible structures (larger peak widths) than the precursor  $[\{\text{Cr}_{12}\text{Gd}_4\} - \text{Piv}]^+$ , however most of the products still appear with unimodal conformations (similarly to the fragments of  $[\{\text{Cr}_{12}\text{Cu}_2\} + \text{Na}]^+$ ). This suggests that no major ring perturbation takes place and both horseshoe units are possibly disrupted simultaneously. In either case, the  $\{\text{CrPiv}_3\}$  leaving group seems to be the driving force of this disassembly route, which we also observed in our previous works.<sup>23,26</sup>

## Conclusions

For the three studied compounds  $\{\text{Cr}_x\text{Cu}_2\}$  ( $x = 10, 12$ ) and  $\{\text{Cr}_{12}\text{Gd}_4\}$ , the formation of smaller rings *via* rearrangements and newly established connectivities suggests a “self-healing” capability upon collision-induced dissociation, driven by the stability of polymetallic rings. Several of the latter were formed including  $\{\text{Cr}_6\text{Cu}\}$ ,  $\{\text{Cr}_5\text{Cu}_2\}$ ,  $\{\text{Cr}_{10}\text{Cu}\}$ ,  $\{\text{Cr}_{10}\}$ ,  $\{\text{Cr}_{12}\text{Cu}\}$ ,  $\{\text{Cr}_{12}\}$ ,  $\{\text{Cr}_6\text{Gd}_2\}$  and  $\{\text{Cr}_5\text{Gd}_2\}$ , which we identified by ion mobility. This workflow can potentially inspire new applications in materials science, if these or similar complexes can be made in isolable quantities, and in general aid the design process of polymetallic complexes by identifying feasible targets. We also propose a simple method to assign whether a polymetallic complex exists as a closed or open structure and this approach can in the future be adapted for other compound families. We believe that the presented workflow, with some modifications, will be particularly useful for the structural characterisation of larger metallocupramolecular compounds, where X-Ray crystallography is often not feasible.<sup>43</sup>

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## Methods

**Synthesis.** {Cr<sub>12</sub>Cu<sub>2</sub>} was synthesised according to our previous work.<sup>24</sup> {Cr<sub>10</sub>Cu<sub>2</sub>} was made similarly to [NH<sub>2</sub>Et<sub>2</sub>]<sub>2</sub>[Cr<sub>10</sub>Cu<sub>2</sub>F<sub>14</sub>(O<sub>2</sub>C<sup>t</sup>Bu)<sub>22</sub>],<sup>25</sup> but the precursor diethylamine was replaced with dipropylamine. Elemental analysis (%) calcd. for {Cr<sub>10</sub>Cu<sub>2</sub>}: C 43.84, H 6.94, N 0.84, Cr 15.56, Cu 3.80; found: C 44.13, H 6.99, N 0.89, Cr 15.42, Cu 3.86. {Cr<sub>12</sub>Gd<sub>4</sub>} was synthesized similarly to [NH<sub>2</sub>Et<sub>2</sub>]<sub>2</sub>[Cr<sub>12</sub>Gd<sub>4</sub>F<sub>21</sub>(O<sub>2</sub>C<sup>t</sup>Bu)<sub>29</sub>],<sup>27</sup> by using the previously reported<sup>44</sup> [(NH<sub>2</sub><sup>n</sup>Pr<sub>2</sub>)<sub>3</sub>Cr<sub>6</sub>F<sub>11</sub>Piv<sub>10</sub>]<sub>2</sub> as the reactant instead of [(NH<sub>2</sub>Et<sub>2</sub>)<sub>3</sub>Cr<sub>6</sub>F<sub>11</sub>Piv<sub>10</sub>]<sub>2</sub>. Elemental analysis (%) calcd. for {Cr<sub>12</sub>Gd<sub>4</sub>}: C 39.38, H 6.17, N 0.58, Cr 13.03, Gd 13.13; found: C 39.54, H 6.21, N 0.63, Cr 12.85, Gd 13.45. All reagents and solvents were purchased from Alfa, Fisher Scientific, Fluorochem or Sigma-Aldrich and used without further purification.

**Sample Preparation.** Samples were prepared in 4:1 toluene/methanol with 500 μM NaI. Analyte concentrations of 200 – 500 μM were typically used for IM-MS and MS measurements.

**nano-Electrospray Ionisation (nESI).** Samples were ionised and transferred to the gas phase with a nESI source and were sprayed from borosilicate glass capillaries (World Precision Instruments, Stevenage, UK). The latter were pulled on the Flaming/Brown P-2000 laser puller (Sutter Instrument Company, Novato, CA, US). The capillary voltage (typically 1.0 - 1.8 kV) was

applied through a platinum wire (Diameter 0.125 mm, Goodfellow, Huntingdon, UK) inserted into the nESI capillaries. Source temperatures of 23 °C (Cyclic) or 30 °C (Q Exactive UHMR) were applied.

*Tandem Mass Spectrometry (MS<sup>2</sup>).* The Q Exactive Ultra-High-Mass-Range (UHMR) Hybrid Quadrupole-Orbitrap Mass Spectrometer (Thermo Fisher) was used for the derivation of the  $E_{50}$  values *via* MS<sup>2</sup> experiments, and more precisely collision-induced dissociation (CID).<sup>45</sup> Target ions were  $m/z$ -isolated in a quadrupole filter, accelerated to a user-defined kinetic energy ( $E_{lab}$ : 0 - 300 eV) and injected into the higher-energy C-trap dissociation (HCD) cell, which contained nitrogen gas (trapping gas pressure parameter: 2.0). Non-fragmented precursor ions and fragment ions were transferred to the Orbitrap mass analyser (AGC target: 3E6 ions, maximum inject time: 100 ms, resolution: 25000).

*Ion mobility mass spectrometry (IM-MS)* experiments were performed on a Select Series Cyclic IMS (Waters).<sup>46</sup> Following ionization (Cone Voltage: 20 - 60 V, Source Offset: 10 - 20 V, Purge Gas: 0 – 300 L/h), ions were transferred to the cyclic ion mobility drift ring (Stepwave Ion Guide RF: 300 – 700 V). Ions were separated by using a non-uniform electric field under a constant nitrogen pressure with travelling waves (TW, Height: 20 - 22 V), which push the ions through the drift ring. Ions travelled one pass in the cyclic drift ring (“single path”, separation time: 2 – 60 ms) and were then transferred (Transfer Energy: 4 – 15 V) to a time-of-flight mass analyser.

*Data Processing.*  $E_{50}$  values were obtained a method described in our previous works.<sup>23,26</sup> Mass spectra were recorded at different collision energies and the share of the precursor ion count, relative to the total ion count (“survival yield”), was plotted vs. the collisional energy in the centre-of-mass frame ( $E_{com}$ , Figures S3, S8). Survival yield plots were fitted with a sigmoidal Hill function (Hill1 function in OriginPro 2020b), yielding the point ( $E_{50}$ ) at which the survival yield reaches 0.5 or 50%. This  $E_{50}$  value is known as a relative measure of precursor ion stability.

Experimentally obtained arrival time distributions (ATD) were converted to collisional cross section distributions  $^{TW}CCS_{N_2}$  (TW = “Travelling Waves”) *via* published calibration procedures.<sup>47</sup> The Agilent tune mix was used as a calibrant.<sup>48</sup>

*Crystallographic Data.* Single crystals of {Cr<sub>10</sub>Cu<sub>2</sub>} and {Cr<sub>12</sub>Gd<sub>4</sub>} were grown by slow diffusion of acetonitrile into toluene solutions of the respective complexes. ORTEP Structures of {Cr<sub>10</sub>Cu<sub>2</sub>} (Figure S10) and {Cr<sub>12</sub>Gd<sub>4</sub>} (Figure S11) can be found in the Supporting Information along with crystallographic refinement details. Crystal structures are also presented in Figure 1a and 1c, respectively.

X-ray diffraction data were collected using a dual wavelength Rigaku FR-X rotating anode diffractometer using MoK $\alpha$  ( $\lambda = 0.71073 \text{ \AA}$ ) radiation, equipped with an AFC-11 4-circle goniometer, VariMAX<sup>TM</sup> microfocus optics, a Hypix-6000HE detector and an Oxford Cryosystems 800 plus nitrogen flow gas system, at a temperature of 100K. Data were collected and reduced using Rigaku CrysAlisPro v42<sup>49</sup> and absorption correction was performed using empirical methods (SCALE3 ABSPACK) based upon symmetry-equivalent reflections combined with measurements at different azimuthal angles. The phase problem was solved using SHELXT and the structural model refined against all  $F^2$  values using SHELXL,<sup>50,51</sup> implemented through Olex2 v1.5.<sup>52</sup> All non-solvent atoms were refined anisotropically. Hydrogen atoms were placed in calculated positions and refined using idealised geometries and assigned fixed isotropic displacement parameters.

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## **Conflict of Interests**

There are no conflicts to declare.

## **Supplementary Data**

The supporting data referred to in this manuscript is contained within a supplementary information document and in a supplementary data set available on Figshare [10.6084/m9.figshare.21751442](https://figshare.com/10.6084/m9.figshare.21751442). The latter includes the raw data of ion mobility mass spectrometry and mass spectrometry measurements.