Rearrangement of \( \{\alpha-P_2W_{15}\} \) to \( \{PW_6\} \) moieties during the assembly of transition-metal-linked polyoxometalate clusters†

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Polyoxometalates (POMs) are molecular oxide units generally formed by highly oxidised transition metal atoms such as W\(^{6+}\), Mo\(^{5+/6+}\), etc. that yield in most cases highly charged anionic clusters.\(^1\) These clusters have a wide range of different architectures and may possess properties applicable to diverse fields such as catalysis,\(^2\) material science,\(^3\) nanotechnology,\(^4\) magnetism,\(^5\) and medicine.\(^6\) By incorporating other metals to make transition metal substituted polyoxometalates (TMSPs), the structural diversity and functionality of the anions is increased.\(^7\) Many TMSPs are based upon either Keggin\(^8\) or Dawson\(^9\) fragments, this is due to the wide library of lacunary building blocks available\(^10\) for both of these parent architectures. One of the interesting aspects of using lacunary precursors is the transformations that can occur when reacting with transition metals, but this is more commonly seen for Keggin type starting materials.\(^11\)

Herein we report two structurally analogous compounds that show the previously unseen transformation of the Dawson-type \( \{\alpha-P_2W_{15}\} \) precursor to a Keggin-type \( \{PW_6\} \) fragment. Both the cobalt and manganese containing TMSPs are obtained using a synthesis adapted from previously reported methodology.\(^12\) The solution behaviour of the cluster is performed using ESI-MS and structural characterization is performed by XRD crystallography.

Compounds 1 and 2, Na\(_{14}\)Li\(_4\)[Co\(_6\)(PW\(_6\)O\(_{26}\))\(_2\)(H\(_2\)O)\(_2\)]\(_{23}\) \( \times \) 55H\(_2\)O and Na\(_{11}\)Li\(_{12}\)[Mn\(_6\)(PW\(_6\)O\(_{26}\))\(_2\)(H\(_2\)O)\(_2\)]\(_{23}\) \( \times \) 58H\(_2\)O are analogues, possessing a V-shaped sandwich structure (Fig. 1). The general formula for the anion is [M\(_6\)(PW\(_6\)O\(_{26}\))\(_2\)(H\(_2\)O)\(_2\)]\(_{23}\) \( \times \) 55H\(_2\)O for 1 and 141H\(_2\)O for 2. The P–O bond lengths of the central \( \{PW_6\} \) unit are between 1.51–1.55 Å for \( \{\alpha-P_2W_{15}\} \) and the phosphorus of the central \( \{PW_6\} \) unit, is 1.45 Å for 1 and 1.41 Å for 2. The P–O bond lengths of the central \( \{PW_6\} \) unit are between 1.51–1.55 Å for 1a and 1.54–1.58 Å for 2a. The M–O bond lengths are between 2.00–2.21 Å for 1a (M = Co) and 2.10–2.29 Å for 2a (M = Mn). The terminal M–O bond lengths are 2.11–2.13 Å for 1a (M = Co) and 2.17–2.19 Å for 2a (M = Mn) which is indicative that these oxygen atoms are water molecules. This is confirmed by bond valence sum...
Compounds 1 and 2 were synthesised using Na$_{14}$[x-P$_2$W$_{15}$O$_{56}$]·24H$_2$O as the lacunary precursor in the presence of the appropriate transition metal ions, lithium chloride and sodium phosphate as a slurry in a minimal amount of solvent in the pH range 6–7. If the pH value is raised to values higher than pH 7.5 then Na$_{13}$Li$_{14}$H$_2$[M$_6$(H$_2$O)$_6$](x-P$_2$W$_{15}$O$_{56}$)$_3$·65H$_2$O can crystallise. For manganese, this trimer (compound 3, see Fig. 3) was observed only a few times in low yield and full characterisation has not been possible due to lack of material. We can thus see that a small pH change can lead to a different set of structures and we deduce that pH control is the most important factor for the synthesis of these compounds. It is worth noting that in all these reactions, the [M$_6$(P$_2$W$_{15}$)]$_3$ sandwich was also obtained as a by-product, which is expected given that [P$_2$W$_{15}$] is not known to reorganise easily.

The nature of the synthetic strategy used makes it difficult to follow the reaction using analytical methods that would give an insight into the mechanism. This is because the reaction requires such a large excess of Na$_{14}$[x-P$_2$W$_{15}$O$_{56}$]·24H$_2$O that any in situ analysis was unfeasible due to the super-saturation and turbidity of the reaction mixture.

There are a few examples in the literature that report banana or V-shaped sandwiches, but all of these architectures are based on Keggin building blocks. All of them have the common [XW$_6$] unit that hinges both parts of the cluster. An iron substituted species that was published by Mialane et al. is a single molecule magnet. Compounds 1 and 2 are special banana TMSPs because there is no known TMSP structure that contains both an [x-P$_2$W$_{15}$] and a {PW$_6$} unit. There are rare examples of mixed Dawson/Keggin species, however we report the first example in which the distinct Keggin and Dawson lacunae have arisen via the transformation of a single lacunary parent. The transformation from [x-P$_2$W$_{15}$] to {PW$_6$} is unexpected as the [x-P$_2$W$_{15}$] Dawson species contains no {PW$_6$} unit of matching connectivity in its structure and it is difficult to postulate how this transformation could occur. A possible mechanism would be the initial decomposition of [x-P$_2$W$_{15}$] to give free tungstate and then it would assemble in two {W$_3$} triads that would react with the {PO$_4$} template generating a [PW$_6$] unit.

In order to investigate the stability of compounds 1 and 2 in solution, electrospray ionisation mass spectrometry (ESI-MS) analysis was conducted. Ionisation for compound 1 was successful and a full spectrum of assignable envelopes was obtained. The ESI-MS spectrum (see Fig. 2, top) shows three envelopes that correspond to the intact cluster. These can be found at the charge values of −6, −7, and −8 at m/z of 1595.5 {Na$_{13}$Li$_{14}$H$_2$[Co$_3$(PW$_6$O$_{26}$)(x-P$_2$W$_{15}$O$_{56}$)]·6H$_2$O}, 1359.8 {Na$_{13}$Li$_{14}$H$_2$[Co$_3$(PW$_6$O$_{26}$)(x-P$_2$W$_{15}$O$_{56}$)]·3H$_2$O} and 1186.8 {Na$_{13}$Li$_{14}$H$_2$[Co$_3$(PW$_6$O$_{26}$)(x-P$_2$W$_{15}$O$_{56}$)]·3H$_2$O}, respectively. A fragment {Co$_3$P$_2$W$_{15}$} species is found at a lower m/z value of 845.4, at a charge of −5 which corresponds to [Na$_{13}$Li$_{14}$H$_2$[Co$_3$P$_2$W$_{15}$O$_{56}$]·11H$_2$O] (see ESI†, Section 5).

Ionisation was also successful for compound 2 and a full spectrum was obtained (see Fig. 2, bottom). In this case there are three envelopes that correspond to the intact cluster. These are found at charge values of −6, −7, and −8 at m/z of 1599.9 [Na$_{13}$Li$_{14}$H$_2$[Mn$_6$(PW$_6$O$_{26}$)(P$_2$W$_{15}$O$_{56}$)]·3(H$_2$O)$_3$·3H$_2$O], 1363.5 [Na$_{12}$Li$_{11}$H$_2$[Mn$_6$(PW$_6$O$_{26}$)(P$_2$W$_{15}$O$_{56}$)]·4H$_2$O] and 1187.2 [Li$_{13}$H$_2$[Mn$_6$(PW$_6$O$_{26}$)(P$_2$W$_{15}$O$_{56}$)]·2(H$_2$O)$_3$·3H$_2$O]. Other peaks at lower m/z values can also be found and they correspond to fragment [Mn$_3$P$_2$W$_{15}$] species. These can be found at the charge values of −5, −6, and −7 at m/z of 817.0 [Li$_{12}$H$_2$[Mn$_3$P$_2$W$_{15}$O$_{56}$]·4H$_2$O], 846.2 [Na$_{13}$Li$_{14}$H$_2$[Mn$_3$P$_2$W$_{15}$O$_{56}$]·12H$_2$O] and 1025.5 [Na$_{13}$Li$_{14}$H$_2$[Mn$_3$P$_2$W$_{15}$O$_{56}$]·4H$_2$O] (see ESI†, Section 5).

Compound 3 is a trimeric structure that consists of a core of nine manganese centres templated by two phosphate units (see Fig. 3). This core is encased by three [x-P$_2$W$_{15}$] units. The full formula of this compound cannot be given due to a lack of reproducibility and very low yields resulting in a full characterisation not being possible. The formula derived solely from crystallographic analysis is
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References


**Fig. 3** Polyhedral and ball-and-stick representation of compound 3 (Mn trimer). Colour scheme: indigo polyhedra: W, yellow spheres: Mn, orange spheres: P and red spheres: O.