

Cite this: *Chem. Sci.*, 2023, 14, 573

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Fatty acid capped, metal oxo clusters as the smallest conceivable nanocrystal prototypes†

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Metal oxo clusters of the type $M_6O_4(OH)_4(OOCR)_{12}$ ($M = Zr$ or Hf) are valuable building blocks for materials science. Here, we synthesize a series of zirconium and hafnium oxo clusters with ligands that are typically used to stabilize oxide nanocrystals (fatty acids with long and/or branched chains). The fatty acid capped oxo clusters have a high solubility but do not crystallize, precluding traditional purification and single-crystal XRD analysis. We thus develop alternative purification strategies and we use X-ray total scattering and Pair Distribution Function (PDF) analysis as our main method to elucidate the structure of the cluster core. We identify the correct structure from a series of possible clusters (**Zr3**, **Zr4**, **Zr6**, **Zr12**, **Zr10**, and **Zr26**). Excellent refinements are only obtained when the ligands are part of the structure model. Further evidence for the cluster composition is provided by nuclear magnetic resonance (NMR), infrared spectroscopy (FTIR), thermogravimetry analysis (TGA), and mass spectrometry (MS). We find that hydrogen bonded carboxylic acid is an intrinsic part of the oxo cluster. Using our analytical tools, we elucidate the conversion from a **Zr6** monomer to a **Zr12** dimer (and *vice versa*), induced by carboxylate ligand exchange. Finally, we compare the catalytic performance of **Zr12**-oleate clusters with oleate capped, 5.5 nm zirconium oxide nanocrystals in the esterification of oleic acid with ethanol. The oxo clusters present a five times higher reaction rate, due to their higher surface area. Since the oxo clusters are the lower limit of downscaling oxide nanocrystals, we present them as appealing catalytic materials, and as atomically precise model systems. In addition, the lessons learned regarding PDF analysis are applicable to other areas of cluster science as well, from semiconductor and metal clusters, to polyoxometalates.

Received 8th September 2022
Accepted 26th November 2022

DOI: 10.1039/d2sc05037d

rsc.li/chemical-science

1 Introduction

Atomically precise clusters are an exciting material class.^{1,2} They span a broad range from oxo clusters, over metal clusters to semiconducting clusters, e.g., $Zr_6O_4(OH)_4(OMe)_{12}$ (**Zr6**, OMe = methacrylate),³ $[Hf_6O_4(OH)_4(OOCMe)_{12}]_2$ (**Hf12**),⁴ $Ti_8O_8(OOCPh)_{16}$,⁵ $Au_{55}(PPh_3)_{12}Cl_6$,⁶ $Cd_{84}Se_{56}(O_2CPh)_{56}(RNH_2)_{56}$,⁷ or $In_{37}P_{20}(O_2CCH_2Ph)_{51}$.⁸ Clusters are used to build stimulative responsive materials,⁹ or hierarchical (porous) structures.^{10,11} Clusters can be precursors or intermediates in nanocrystal synthesis.^{12–15} Composites with improved strength are produced by incorporating zirconium oxo clusters into polymers.^{16,17} Zirconium and hafnium oxo clusters are used as catalysts to make or break amide bonds,^{18,19} and they are suitable as inorganic extreme-ultraviolet photoresists.^{20,21} Subtle differences in cluster chemistry between Zr and Hf are exploited in the

industrial separation of Zr from Hf.²² Titanium oxo clusters exhibit exciting photo-redox properties.²³ Group 4 MOFs are based on group 4 metal oxo clusters that act as secondary building units (or nodes) in the MOF framework.^{5,24–28} Apart from the typical **Zr6**, and **Hf6** oxo clusters used in MOFs, many discrete group 4 oxo clusters exist with different nuclearities.^{2–29}

Discrete group 4 oxo clusters are conceptually close to colloidal oxide nanocrystals and share some features with them.^{2,30,31} For example, both have an inorganic core, usually capped by an organic ligand shell. Discrete oxo clusters are usually synthesized with short and rigid ligands (e.g., acetate or benzoate). The resulting clusters crystallize from the reaction mixture, allowing for convenient purification and structural characterization by single crystal X-ray diffraction (XRD).^{3,32,33} For example, **Zr6**-carboxylate clusters have the general formula: $Zr_6O_4(OH)_4(OOCR)_{12}$. The Zr atoms are arranged as an octahedron and are connected through μ_3 -oxygen bridges, with the largest Zr–Zr distance being 0.5 nm. All zirconium atoms are eight-coordinate and the zirconium–oxygen arrangement is almost identical to bulk, cubic zirconia. Therefore, these clusters can be regarded as the lower limit of downscaling zirconia nanocrystals. In contrast to larger nanocrystals, oxo clusters are

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† Electronic supplementary information (ESI) available. See DOI: <https://doi.org/10.1039/d2sc05037d>



atomically precise (polydispersity = 0),³⁴ and their ligand shell is less densely packed (ligand density = 2.4 ligands per nm² compared to 3–4 ligands per nm² for nanocrystals).^{35,36} When capped by ligands with low steric hindrance, the **Zr6** cluster dimerizes to form a **Zr12** cluster that has two **Zr6** subunits, connected by four inter-cluster bridging carboxylate ligands,² see Fig. 1. Apart from seven regular bridging ligands, there are also three chelating ligands per **Zr6** subunit. There exist two types of **Zr6** monomer structures, either with all bridging ligands or with three chelating and nine bridging ligands, see Fig. 1. Due to their rigid ligands, typical oxo clusters have low solubility, rendering them poorly suitable for subsequent processing.³⁷ Ligands typically used for providing colloidal solubility to nanocrystals (e.g., oleic acid),^{38,39} have not been explored on group 4 oxo clusters. In general, the surface chemistry of group 4 oxo clusters is less studied than that of group 4 oxide nanocrystals.^{35,40–44} Only carboxylate for carboxylate ligand exchange is explored.^{4,45} At room temperature, the **Zr6** and **Zr12** clusters do not interconvert and the bridging ligands appear inaccessible for exchange.

Drawing inspiration from the nanocrystal field, we envision a library of flexible carboxylate ligands to increase the solubility and hence the possible applications of these clusters. Since longer carboxylate ligands do not allow for crystallization, the question becomes how to purify and structurally characterize such objects. Recently, a purification method based on resins with amine groups was developed for oxo clusters with mixed

(but still short) ligand shells.²⁰ About 20% of the acid persists as free acid in the final product. Regarding characterization, there are a few examples where oxo clusters in MOFs have been studied by X-ray total scattering and Pair Distribution Function analysis (PDF).^{46–49} Structural changes upon heating have been observed by a model-free analysis. Recently, oxo clusters were modelled *via* PDF analysis without organic ligands in MOF reaction mixtures.^{47,49} On the other hand, X-ray PDF has been often used to study the structure of nanocrystals. In this case, the organic ligands have usually been neglected in the structural models because the organic ligands scatter very little compared to the inorganic core.^{50–56} In the case of larger clusters (35–144 metal atoms) PDF could confirm the structure of the core,^{7,57–59} again by largely neglecting the organic ligands. However, to our knowledge, PDF has not yet been applied to small discrete oxo clusters capped with organic ligands, as discussed here. One can expect this to be challenging since the core is much smaller.

Here, we establish standardized conditions for the synthesis of a range of carboxylate capped zirconium oxo clusters. We use PDF analysis to structurally characterize the oxo clusters. We find that excellent refinements are only obtained when the structural models include the contribution of the organic ligands. PDF analysis can clearly distinguish a **Zr6** cluster from a **Zr12** cluster and dismiss other possible cluster structures. The organic ligand shell is then comprehensively characterized by FTIR, NMR and TGA. The molecular formula is finally confirmed by electrospray ionization mass spectrometry (ESI-MS). Using our characterization tools, we evaluate carboxylate for carboxylate ligand exchange and we find that the ligand is structure-directing, determining whether a **Zr6** or a **Zr12** cluster is formed. These results are further generalized to hafnium oxo clusters. Finally, we test the notion that the fatty acid capped metal oxo clusters are the smallest conceivable nanocrystal prototypes. We compare the catalytic activity of **Zr12**-oleate clusters with oleic acid capped zirconium oxide nanocrystals, and find a superior activity for the oxo clusters.

2 Results and discussion

2.1 Synthesis of oxo clusters

Zirconium propoxide (70 w% in propanol) or zirconium butoxide (80 w% in butanol) reacts with an excess of acetic acid (8 equivalents) to form a **Zr12**-acetate cluster, which is a dimer of **Zr6**.⁴ The reaction is performed in dichloromethane (DCM) solvent to obtain higher quality crystals.⁴ Both acetic acid and DCM are co-crystallized with the cluster.⁴ The balanced chemical equation is shown in Fig. 2. We further washed the crystalline powder with a solution of acetic acid in DCM to remove reaction by-products and other zirconium species. Although the powder can be handled in air, it is recommended to keep the clusters in a glovebox or desiccator for long term storage, since they react slowly with atmospheric water and become partially insoluble.

Zr12-propionate is similarly synthesized and purified as **Zr12**-acetate. We extended the synthetic strategy to longer carboxylic acid ligands (see Fig. 2), but the resulting clusters

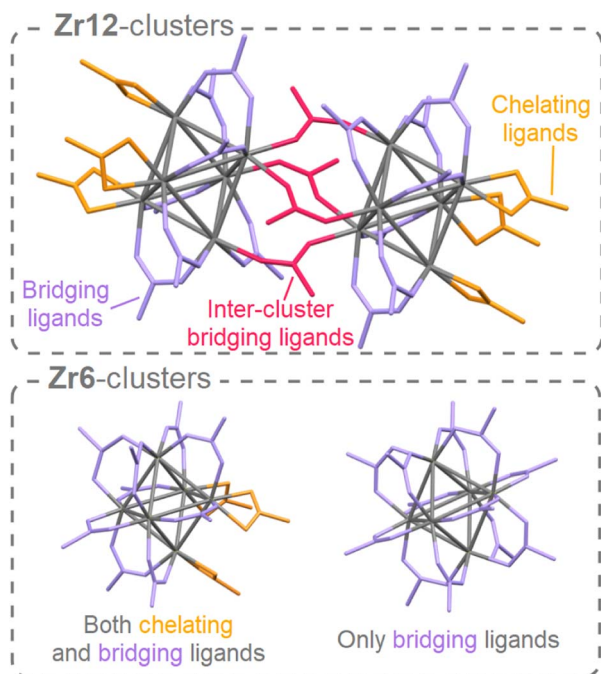
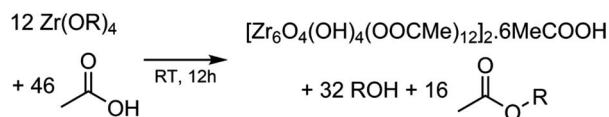


Fig. 1 The structure of the **Zr12** dimer and **Zr6** monomers. For the dimer, there are three different types of ligands: chelating, bridging, and inter-cluster bridging. The latter connects two monomers to form the dimer. A monomer can have either only bridging ligands or a combination of bridging and chelating ligands. For clarity, hydrogen atoms and oxygen atoms on the cluster core are omitted, and ligand chains are shortened to acetate.



Zr₁₂-acetate synthesis

Other ligands used for cluster synthesis

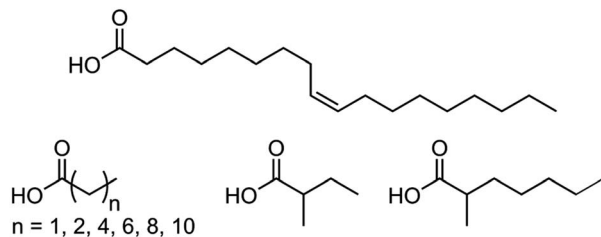


Fig. 2 Balanced chemical equation for the synthesis of Zr₁₂-acetate clusters from zirconium propoxide (or butoxide) and acetic acid. The final product has co-crystallized acetic acid and dichloromethane (the latter is not shown). Also the structures of the other carboxylic acids used here for cluster synthesis are shown.

cannot be crystallized anymore and therefore other purification techniques are required. Zr₁₂-butanoate is washed overnight in acetonitrile, while Zr₁₂-hexanoate, Zr₆-methylbutyrate and Zr₆-methylheptanoate clusters can be precipitated with acetonitrile and isolated by centrifugation. Dissolution in DCM and repeating the precipitation cycle twice, delivers the purified clusters. This type of purification is typical for colloidal nanocrystals,⁶⁰ and thus reinforces the idea of fatty acid capped clusters as atomically precise models for nanocrystals. For the clusters with even longer ligands (octyl, decyl, dodecyl and oleyl chains), the first precipitation is performed with acetonitrile, while acetone is the preferred anti-solvent for the two consecutive steps to precipitate the clusters, and separate it from, *e.g.*, the ester co-product. The physical appearance of the clusters ranges from semi-crystalline solids (acetate, propionate and methylbutanoate) to waxy solids (butanoate, hexanoate, decanoate and dodecanoate) or viscous oils (octanoate, oleate and methylheptanoate), see Fig. S1.†

2.2 Structural analysis by X-ray PDF

While the Zr₁₂-acetate and Zr₁₂-propionate clusters are known in the literature, the others are not. To prove their structure, we analyzed the clusters with X-ray total scattering and Pair Distribution Function (PDF) analysis. Total scattering takes into account both Bragg and diffuse scattering. The real space PDF is ideal to study nanomaterials, local effects and amorphous materials.^{61–63} We measured the experimental PDF of all the above clusters and ordered them according to increasing carbon contents, see Fig. 3. Focusing first on the PDF of acetate capped clusters and comparing it with the structure from single crystal XRD,⁴ we easily recognize the Zr–O distance at 2.2 Å and the Zr–Zr distances within one Zr₆ cluster at 3.5 Å and 5 Å. We label them as intra-Zr₆ distances. The longer range Zr–Zr distances are specific to the Zr₁₂ dimer and are present at 5.7 Å,

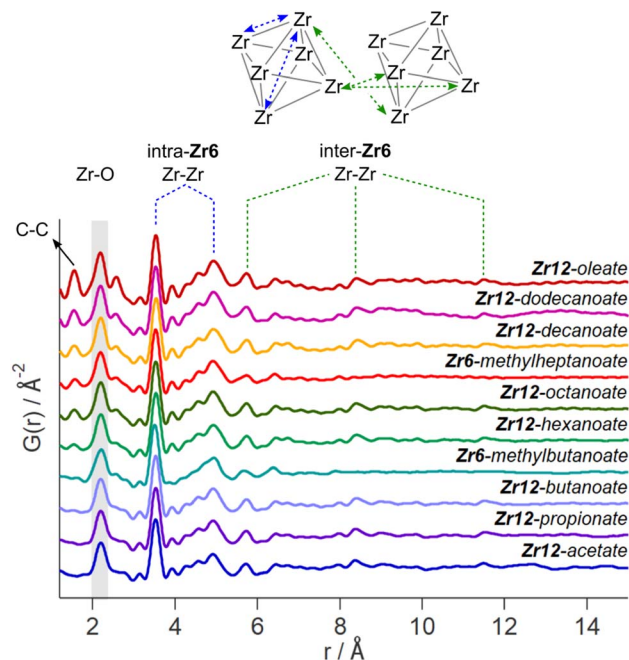


Fig. 3 Experimental PDF of zirconium oxo clusters with different capping ligands. The C–C, Zr–O, and Zr–Zr distances are assigned. We make a distinction between the Zr–Zr distances within one Zr₆ cluster and the Zr–Zr distances that are characteristic for the dimer.

8.4 Å, and 11.5 Å. They are labeled as inter-Zr₆ distances. When the chain length increases, the same features remain present in the PDF, except for the case of methylbutanoate and methylheptanoate where the inter-Zr₆ distances are missing. The peak at 5.7 Å does not completely disappear but changes shape and has a lower intensity. The residual intensity stems from intra-Zr₆ Zr–O distances (not the first coordination sphere but longer range correlations). The above observations suggest that both the Zr₆-methylbutanoate and Zr₆-methylheptanoate clusters are monomers while the others are dimers. This is in agreement with our earlier hypothesis that branching of the carboxylic acid at the alpha position is the deciding factor between the monomer and dimer.² On analyzing the theoretical structure of Zr₁₂-isobutyrate (SI), the steric hindrance is clear from the space filling representation. On the other hand, a higher degree of oligomerization was excluded based on the absence of features beyond 11.5 Å, and based on the crystal structure of Zr₁₂-acetate. With increasing chain lengths, we also observe the growth of a peak at 1.5 Å, which we assign to C–C distances in the ligand chain.

Apart from this model-free analysis, we sought to quantitatively fit the PDFs. We focus again first on Zr₁₂-acetate to refine our approach since a single crystal XRD structure is available for this cluster (CCDC-604528).⁴ To demonstrate the importance of the organic ligands to the PDF, we theoretically calculated the PDF of Zr₁₂-acetate, including either (i) no carbon atoms, (ii) only the carbonyl carbon atoms, or (iii) all carbon atoms, see Fig. S2.† The carbon atoms contribute significantly to the PDF, especially between 4 Å and 8 Å. This result stands in stark contrast to the regular practice of ignoring the ligands in the



PDF refinement of nanocrystals and larger clusters.^{7,50–59} The ligand effect is further demonstrated by fitting the **Zr12**-acetate PDF with various models, see Fig. 4. For each structure model, we refined thermal motion parameters while keeping the structure (*i.e.*, the atomic positions) unchanged. In the first attempt, similar to traditional approaches, we only used the **Zr6** cluster core as a model. The model could capture the basic Zr–O and intra-**Zr6** Zr–Zr distances but gave overall a poor agreement and a significant misfit in the Zr–O distance. Using the dimer structure, the inter-**Zr6** Zr–Zr distances are well described. By including the oxygen atoms from coordinated acetates, a significant reduction in R_w is achieved and the Zr–O peak is now accurately fitted. By including the carbon atoms from coordinated acetate, we obtain a decent fit, with $R_w = 0.14$.

The crystal structure of **Zr12**-acetate contains also hydrogen bonded ligands and co-crystallized DCM molecules. When the hydrogen bonded ligands are explicitly included, only a marginally better fit is obtained, see Fig. S3.† Alternatively, we added an exponentially decaying sinusoidal function as a second phase, resulting in a slightly (but significantly) better fit with an R_w value of 0.11, see Fig. S4.† This second phase describes the disordered but still structured intensity and has previously been used to model solvent restructuring, or a regular array of high/low density regions.^{64,65} While it is gratifying to obtain an excellent fit for the data, we sought to determine whether we could distinguish the **Zr12**-acetate

structure from other reported cluster structures (determined by single crystal XRD). We thus attempted to fit the experimental PDF of **Zr12**-acetate clusters with the reported structures of **Zr12**-propionate, **Zr6**-trimethylacetate, **Zr6**-methacrylate, **Zr6**-isobutyrate, **Zr6**-acetate, **Zr4**-methacrylate, **Zr4**-formate isopropoxide, **Zr3**-acetate isopropoxide, **Zr3**-acetate *tert*-butoxide, **Zr10**-salicylate, and **Zr26**-formate, see Fig. S5.†^{3,4,32,66–70} For these refinements, we removed the excess carbons in the structures to arrive at models with acetate ligands. The models based on **Zr3**, **Zr4**, **Zr10** and **Zr26** all showed very poor agreement with the data. The **Zr6** structure models also delivered poor fits ($R_w = 0.24–0.31$). The second best agreement was obtained for the **Zr12**-propionate model ($R_w = 0.18$). While this model is structurally very similar to the **Zr12**-acetate model, the Zr–Zr distances are slightly larger in the propionate model (Table S6†). We conclude that PDF is able to pick up the correct structure from a series of possible cluster structures, based on the goodness of fit.

The developed strategy is now applied to the other oxo clusters. Fig. 5 shows the PDF refinement for four selected oxo clusters, capped with either butanoate, methylbutanoate, octanoate or oleate. In each case, we use a cluster model along with a decaying sine wave. For **Zr12**-butanoate and **Zr12**-octanoate, we find that the **Zr12**-acetate structure model gives a satisfactory fit (Fig. 5), which is clearly better than the refinement with a **Zr6**-acetate structural model (Fig. S6†). The latter structure has all bridging ligands and was reported after crystallization from an

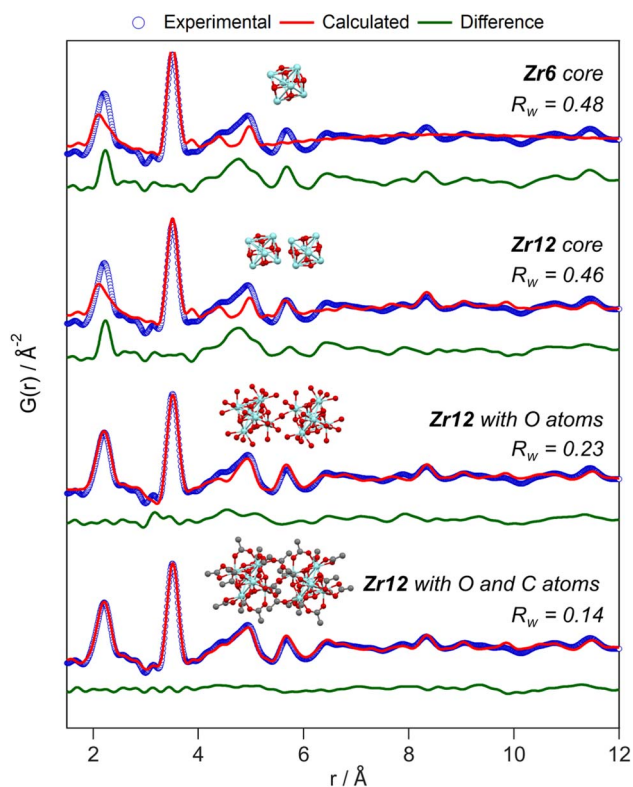


Fig. 4 PDF refinement of the **Zr12**-acetate cluster with various models, derived from the reported crystal structure.⁴ The best fit is obtained when including the oxygen and carbon atoms from the acetate ligands. The refined parameters are given in Table S1.†

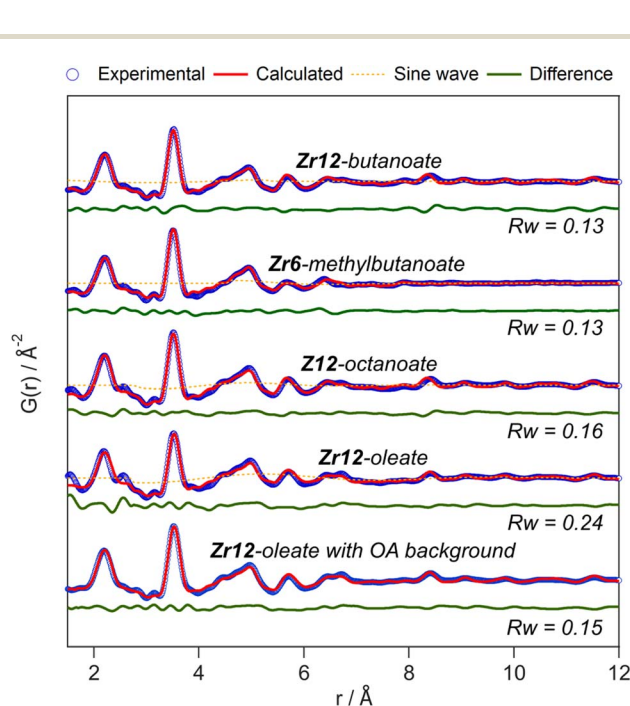


Fig. 5 PDF refinement for **Zr12**-butanoate and **Zr12**-octanoate using the **Zr12**-acetate structure model. PDF refinement of **Zr6**-methylbutanoate using the **Zr6**-acetate structure model. Finally, the PDF refinement of **Zr12**-oleate using the **Zr12**-propionate structure model, with or without background correction in reciprocal space. If applied in the refinement, the exponentially dampening sine wave is shown (orange dotted lines). The refined parameters are given in Table S3.†



aqueous solution.⁶⁶ In contrast, the **Zr6**-methylbutanoate cluster, is best described by using the **Zr6** model and gives a worse fit for the **Zr12** model. The **Zr12**-oleate cluster was best described by using the **Zr12**-propionate structure model with slightly larger Zr–Zr distances. Again, the **Zr6** model delivered a worse fit (Fig. S6†). We can thus distinguish the **Zr6** structures from the **Zr12** structures, based on the refinement. However, we notice that the R_w increases with increasing chain length (indicating a worse fit). The origin lies in the fact that we did not explicitly include all the carbons of the long ligands in the models. The short-range C–C distances (at 1.5 and 2.6 Å) are not fitted in the refinement. However, the sinusoidal function seems to describe the long range C–C distances quite well, presumably because the ligand chain is disordered. This hypothesis is confirmed by the increase in the relative amplitude of the sine wave from **Zr12**-acetate to **Zr12**-oleate (Tables S1 and S3†). The improvement in R_w upon including the sinusoid is small for acetate (0.03) and large for oleate (0.09) (see Fig. S7†).

A more effective way of dealing with long ligands is measuring the free ligand explicitly as a background and subtracting its scattering from the data in reciprocal space. For **Zr12**-oleate, this leads to a significant improvement in the fit. The short range C–C distances disappeared from the PDF (Fig. 5) and the sine wave is completely obsolete (Table S3†). The goodness of fit is now comparable to that of the case of the clusters with shorter chains. Note that the presence of long ligands reduces the signal-to-noise ratio during the data collection (due to a lower fraction of oxo clusters in the X-ray beam). We conclude that, after initial training on the acetate clusters, our PDF analysis is able to confirm the structure of these previously unreported clusters. Our synthetic procedures indeed lead to the formation of soluble **Zr6** oxo clusters, forming monomers or dimers depending on the steric hindrance of the ligand. We further emphasize that good refinements can only be achieved by including the binding group of the ligand explicitly in the model (even after background subtraction). This is much more important for the small clusters discussed here, than for larger clusters or nanocrystals. For the latter two, there is a larger core of strongly scattering atoms, and relatively fewer, weakly scattering ligands. However, small nanocrystals have typically worse a goodness of fit compared to larger ones and it might be worth considering including ligands explicitly to obtain better refinements.⁵⁰

2.3 The organic ligand shell

For PDF modelling we need to include the carboxylate head group, but PDF is in general not very sensitive to the organic fraction. Detailed information on the presence of uncoordinated ligands or coordination modes is hard to obtain from PDF. We turn now to FTIR, TGA and NMR. In Fig. 6, we show the FTIR spectra of selected oxo clusters (the complete FTIR spectra of all clusters are presented in Fig. S8†). The CH_2 stretches ($2800\text{--}3000\text{ cm}^{-1}$) gain intensity with increasing chain length. The small, sharp peak at around 3450 cm^{-1} and the broad peak between 3500 and 3000 cm^{-1} were assigned to the $\mu_3\text{-OH}$

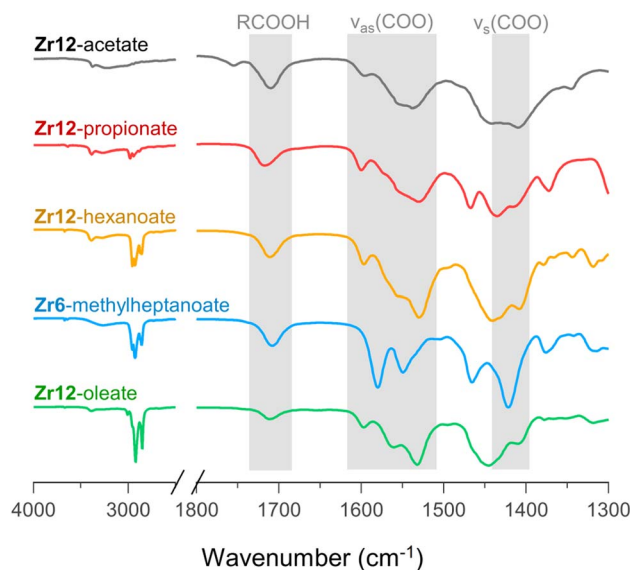


Fig. 6 FTIR spectra of **Zr12**-acetate, -propionate, -hexanoate, and -oleate and **Zr6**-methylheptanoate. The weak band at 1750 cm^{-1} in the spectrum of **Zr12**-acetate is assigned to a small amount of acetic acid that is not involved in hydrogen bonding of any kind.⁷¹ An ester impurity is ruled out because of the absence of any signal at around 4 ppm in NMR, (see Fig. S10†).

moieties on the cluster core, which are involved in hydrogen bonding. While all these samples were extensively purified by multiple precipitation/re-dissolution cycles, it is clear that there is still protonated carboxylic acid in the sample (observed at 1710 cm^{-1}), presumably hydrogen bonded to the cluster. Such hydrogen bonding is also observed for the co-crystallized acetic acid in the single-crystal structure of **Zr12**-acetate, see Fig. S9.† Our efforts to purify the clusters further and remove the hydrogen bonded acid were not successful and even compromised the integrity of the cluster core, see the ESI.† Since attenuated total reflection (ATR) FTIR is not quantitative, we used TGA to quantify the amount of excess carboxylic acid. We find about 1.1–3.7 excess ligand molecules per **Zr6** monomer, see the ESI.† There is no clear trend among the different clusters. Only **Zr12**-acetate stands out with 5.5 excess ligands per monomer.

Returning to the FTIR spectrum, the bands in the region $1500\text{--}1600\text{ cm}^{-1}$ are assigned to the asymmetric stretch of the carboxylate, $\nu_{\text{as}}(\text{COO})$. While the symmetric stretch, $\nu_{\text{s}}(\text{COO})$, is expected at around $1400\text{--}1450\text{ cm}^{-1}$, there is significant overlap of CH_2 and CH_3 deformation bands which are expected at around $1440\text{--}1460$. Focusing here on the qualitative differences, we first draw attention to the spectrum of **Zr6**-methylheptanoate. In contrast to the IR spectrum of **Zr6**-pivalate,⁷² $\nu_{\text{as}}(\text{COO})$ is split into two bands (1580 and 1550 cm^{-1}). While **Zr6**-pivalate has all bridging carboxylates, the band splitting for **Zr6**-methylheptanoate indicates that part of the methylheptanoate ligands are in the chelating mode. When comparing the pattern of the $\nu_{\text{as}}(\text{COO})$ bands in **Zr6**-methylheptanoate with those of **Zr12**-oleate, we find striking differences. For **Zr6**-methylheptanoate, the most intense band is at 1580 cm^{-1} while



for **Zr12**-oleate, the most intense band is at 1532 cm^{-1} . In addition, **Zr12**-oleate has a band with the highest wavenumber, 1597 cm^{-1} . Typically the wavelength difference between $\nu_{\text{as}}(\text{COO})$ and $\nu_{\text{s}}(\text{COO})$ is used as a diagnostic tool to distinguish between chelating ($\Delta\nu < 110\text{ cm}^{-1}$), bridging ($\Delta\nu = 140\text{--}190\text{ cm}^{-1}$) or monodentate ($\Delta\nu = 200\text{--}320\text{ cm}^{-1}$) binding modes.^{73,74} This is an empirical rule-of-thumb and in the case of multiple bands it is difficult to assign the paired $\nu_{\text{s}}(\text{COO})$. With these limitations in mind, we tentatively calculate $\Delta\nu = 190\text{ cm}^{-1}$ for the band at 1597 cm^{-1} , and assign it to the ligands that bridge the two clusters. More importantly, the same $\nu_{\text{as}}(\text{COO})$ pattern of **Zr12**-oleate is retrieved for all the **Zr12** clusters (see also Fig. S8†). Furthermore, the $\nu_{\text{as}}(\text{COO})$ pattern of **Zr6**-methylheptanoate is quasi identical to that of **Zr6**-methylbutanoate. This is an exciting observation since it provides a very fast method to distinguish between the **Zr6** and **Zr12** clusters.

The ^1H NMR spectra of the selected oxo clusters are shown in Fig. 7A. For **Zr12**-propionate, hexanoate and oleate, we recognize three different resonances for the alpha CH_2 moiety, indicating three different distinguishable carboxylate environments (see also Fig. S13† for the other **Zr12** clusters). On the other hand, **Zr6**-methylheptanoate has only two alpha CH resonances (see Fig. S13† for **Zr6**-methylbutanoate). It is

striking that the NMR pattern seems to roughly agree with the $\nu_{\text{as}}(\text{COO})$ pattern in FTIR. For **Zr12**-acetate, we find four methyl resonances. Diffusion ordered spectroscopy (DOSY), shows that three resonances (1.7–2 ppm) have the same diffusion coefficient, while the fourth (2.1 ppm) diffuses more quickly, see Fig. 7B. The resonance at 2.1 ppm thus corresponds to excess acetic acid, as also determined by FTIR and TGA. The measured diffusion coefficient ($D = 1106\text{ }\mu\text{m s}^{-1}$) is lower than the reference of free acetic acid ($D = 1851\text{ }\mu\text{m s}^{-1}$), and we conclude that the excess acetic acid is in fast exchange between two states: (i) free and (ii) hydrogen bonded to the cluster.⁷⁵ The signals between 5 and 7 ppm in all cluster spectra are assigned to the exchangeable protons on the cluster core itself since the resonances disappear upon addition of D_2O , see Fig. S14.† Apart from **Zr12**-acetate and -propionate, all the spectra show significant line broadening, similar to nanocrystal-bound ligands.⁷⁶ The longer the ligands, the broader the alpha CH_2 resonance, and the sharper the methyl resonance becomes. Homogeneous line broadening depends on the rotational mobility and it is expected that the methyl group has increased mobility in a longer ligand.⁷⁶ The alpha resonance is closest to the surface. Its mobility is not enhanced in a longer ligand and instead, the total rotational mobility of the cluster decreases since it becomes a bigger object. In contrast to nanocrystal-bound ligands, we find little contribution of heterogeneous broadening to the resonances, see Table S8.†

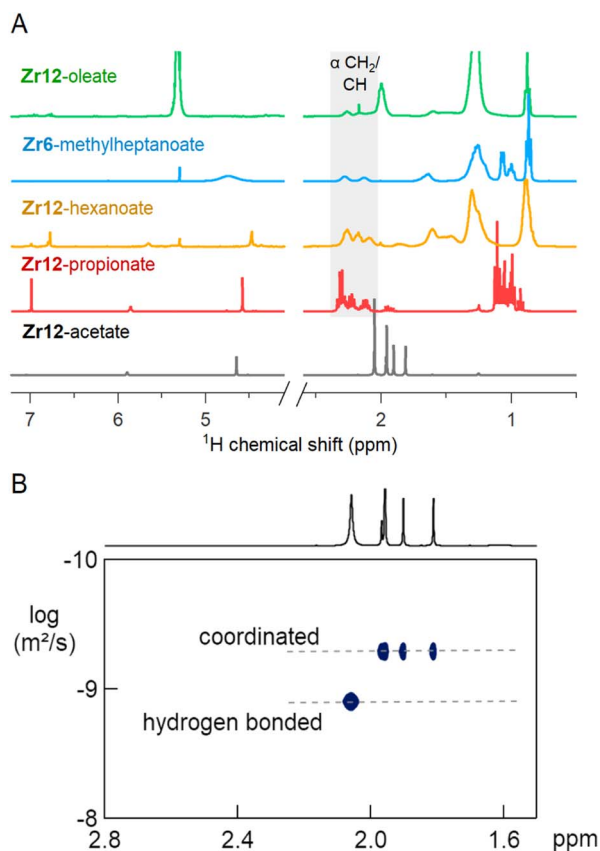


Fig. 7 (A) NMR spectra in CDCl_3 of **Zr12**-acetate, -propionate, -hexanoate and -oleate and **Zr6**-methylheptanoate. (B) DOSY of the **Zr12**-acetate cluster with one faster diffusing species and a set of three resonances pertaining to a slowly diffusing species (the cluster).

2.4 The molecular formula by mass spectrometry

We sought to use electrospray ionization high resolution mass spectrometry (ESI-HR-MS) to obtain the mass of the clusters. Unfortunately, our carboxylate capped clusters show limited solubility in typical ESI-HR-MS solvents such as methanol and acetonitrile. **Zr12**-acetate clusters dissolve in methanol but no clusters were detected in ESI-HR-MS and NMR analysis provides evidence for cluster degradation in methanol, see Fig. S16 and S17.† The limited solubility of **Zr12**-acetate did not allow for any further HR-MS analysis. On the other end, the **Zr12**-oleate clusters were too non-polar. The other clusters could be dissolved in tetrahydrofuran (THF) and were analyzed with ESI-HR-MS after adding acetonitrile as co-solvent (full spectra and assignments in Fig. S18–S25†). The **Zr6** clusters give reasonably clean spectra where the molecular ion can be recognized as $\text{Zr}_6\text{O}_4(\text{OH})_3(\text{L})_{12}^+$ (the cluster minus an OH^- group). The example of **Zr6**-methylbutanoate is shown in Fig. 8. The **Zr12** clusters appear less stable and the ion $[\text{Zr}_6\text{O}_4(\text{OH})_7(\text{L})_9]\text{H}^+$ was observed in most cases (see Fig. 8). This ion is the proton adduct of the **Zr12** dimer after substituting six carboxylates for six hydroxides. The **Zr12** cluster indeed features 6 chelating ligands, which are most labile according to the literature.⁴ It is thus plausible that such an exchange takes place, further catalyzed by a coordinating solvent like THF. Interestingly, more fragmentation was seen for longer ligands and the monomeric form is even dominant for **Zr12**-decanoate and **Zr12**-dodecanoate. Despite the limitations, MS thus confirms the successful formation of oxo clusters with long carboxylate ligands. It is



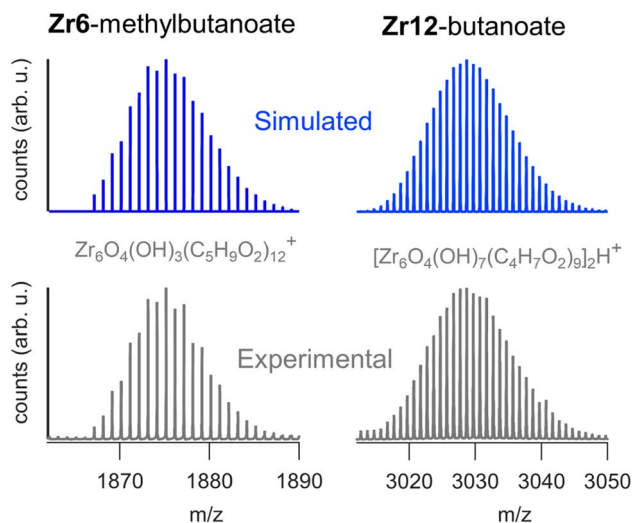


Fig. 8 ESI-HR-MS analysis of the dimeric **Zr12**-butanoate cluster and of the monomeric **Zr6**-methylbutanoate cluster. Both the experimental and simulated spectra are shown.

however not the correct technique to distinguish the dimer from the monomer structure.

2.5 Ligand exchange

Ligand exchange where all carboxylates are replaced by another type of carboxylate is an interesting way to change the cluster structure and/or functionality. Pucherberger *et al.* previously studied the exchange between methacrylate and propionate/acetate on zirconium oxo clusters by NMR. They inferred that, at room temperature, the monomer and dimer cannot be converted into one another and that the inter-cluster bridging ligands are unavailable for exchange.⁴ Here, we re-evaluate the ligand exchange process under more forcing conditions and apply our characterization toolbox to elucidate the cluster structure. Starting from the **Zr12**-acetate cluster, we stir it with 1.5 equivalents of a new ligand (a carboxylic acid from Fig. 2) before we evaporate all volatile species at 70 °C. Since acetic acid has a reasonably low boiling point, it is readily removed from the reaction mixture, forcing the ligand exchange to proceed to completion (confirmed by the absence of acetate signals in the ¹³C NMR, see Fig. S26†). The clusters are then purified to remove excess high boiling ligand. PDF analysis of the clusters shows that the dimer is obtained for butanoate, hexanoate, octanoate, decanoate, dodecanoate and oleate, see Fig. S27.† The **Zr6** monomer is retrieved for methylbutanoate or methylheptanoate, showing that the dimer can be broken up under appropriate conditions and with the right structure-directing ligand. Also, the IR and NMR spectra look identical to the ones from the clusters that are directly synthesized from zirconium propoxide, see Fig. S28 and S29.† We also treated the **Zr6**-methylbutanoate cluster twice with 3 equivalents of hexanoic acid. Methylbutanoic acid does not evaporate as easily as acetic acid and a higher excess of new ligand was needed. The resulting cluster has the typical IR and NMR signatures of a **Zr12** cluster, see Fig. S30 and S31.† Also the HRMS shows similar

results compared to the HRMS of bottom up synthesized **Zr12**-hexanoate, see Fig. S21 and S32.† The results of the ligand exchanges are summarized in Fig. 9.

2.6 Hafnium oxo clusters

To generalize our methodology to hafnium oxo clusters, we reacted hafnium butoxide with acetic acid, methylbutanoic acid, or oleic acid. After purification, the resulting clusters were characterized by PDF, FTIR, TGA, and NMR. The acquisition and refinement of PDF data are more straightforward for the hafnium oxo clusters since hafnium has a higher scattering cross section, see Fig. 10A. All pairs that have at least one hafnium atom are enhanced in intensity, and all other pairs (*e.g.*, C-C) lose relative intensity. Using a structural model of **Hf12**-acetate (CCDC-604534),⁴ we can readily describe the PDF data of the **Hf12**-acetate and **Hf12**-oleate clusters. The goodness of fit is even better than for the respective **Zr12** clusters that were refined in an equivalent way, see Fig. 5. In the absence of a **Hf6** structure model, we constructed a **Hf6** cluster model from the **Hf12** structure. This model accurately described the PDF data, confirming that the **Hf6**-methylbutanoate cluster is a monomer. The hafnium oxo clusters thus follow the same structure-directing rules as the zirconium oxo clusters. Also, the FTIR spectra (Fig. 10B) and NMR spectra (Fig. S33†) show the expected patterns, similar to those in the case of zirconium. Ligand exchanges can be performed from acetate to oleate or methylbutanoate ligands, forming the expected **Hf12** or **Hf6** cluster respectively, based on FTIR and NMR data (Fig. S34 and S35†). Finally, TGA shows that there is again free carboxylic acid present on top of the charge balancing ligand shell (Fig. S36 and Table S10†).

2.7 Catalysis

Finally, we take the perspective that these fatty acid capped oxo clusters are the smallest conceivable nanocrystal prototypes.^{2,30} While technically not a crystal anymore, such a cluster represents the lower limit of scaling down a ligand capped metal oxide nanocrystal. Hence, it has a maximized surface-to-volume ratio. Maximizing surface area is particularly important in heterogeneous catalysis and we thus hypothesized that oxo clusters (diameter = 0.5 nm) would have superior catalytic performance over metal oxide nanocrystals (diameter = 2–10 nm). To test this, we take the previously reported esterification

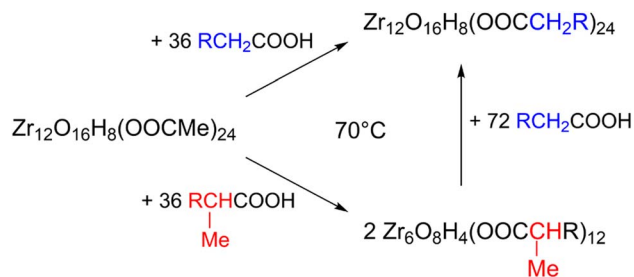


Fig. 9 Exchange reaction of the **Zr12**-acetate cluster with hexanoic acid, and methylbutanoic acid.



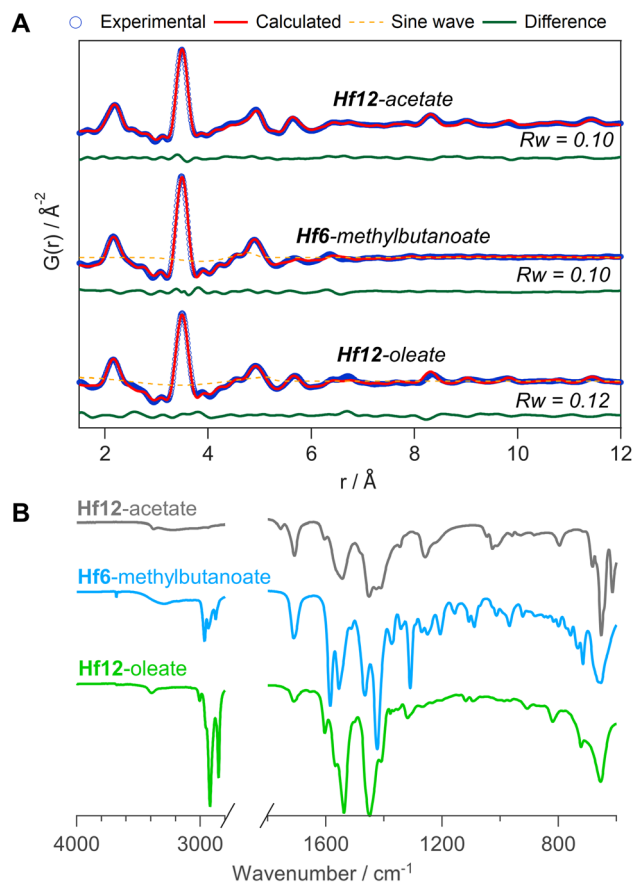


Fig. 10 (A) PDF fit for Hf12-acetate, Hf6-methylbutanoate and Hf12-oleate clusters with exponentially dampening sine wave contribution. The refined parameters are indicated in Table S9.† (B) FTIR spectra of the hafnium oxo clusters synthesized *via* bottom up.

of oleic acid with ethanol as a catalytic model system.⁴¹ We synthesized 5.5 nm tetragonal zirconia nanocrystals, and functionalized their surface with oleate ligands,⁷⁷ see also Fig. S37 and S38.† **Zr12-oleate** clusters are the atomically precise, smallest conceivable prototype of such nanocrystals. Given that both clusters and nanocrystals have the same oleate ligand, the only difference is the inorganic core. In addition, the ligand is identical to the catalytic substrate (oleic acid), thus avoiding competition between the ligand and substrate for the surface sites,⁴¹ or the formation of side products. To compare the efficiency of the two catalysts, we added in both cases 10 mol% Zr to the reaction mixture. We monitored the reactions for the first 5 hours by taking aliquots and quantifying the ester product *via* NMR, see Fig. 11.

The oxo clusters are clearly more catalytically active than the nanocrystals, confirming our hypothesis. An initial rate analysis shows that the rate of the reaction when using **Zr12-oleate** is 0.3 mM min^{-1} while for ZrO_2 nanocrystals it is 0.06 mM min^{-1} . The clusters thus feature about a five-fold higher reaction rate, which we relate to a higher surface-to-volume ratio. The cluster has every zirconium atom available at its surface. In contrast, we estimate that 19% (about one fifth) of the zirconium atoms in the nanocrystals are available for catalysis. The other 81% is

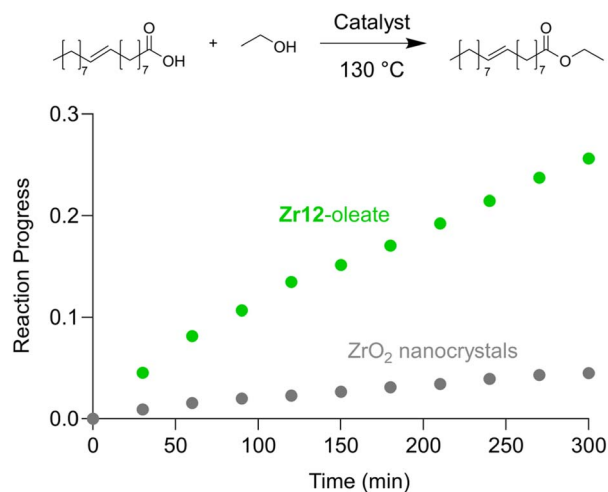


Fig. 11 Catalytic esterification of oleic acid with ethanol, catalyzed by either **Zr12-oleate** (green) or ZrO_2 nanoparticles (gray). In both cases, 10 mol% Zr, with respect to the carboxylic acid substrate, was added.

buried under the surface. This estimation agrees well with the five-fold rate enhancement of the clusters. Zirconium and especially hafnium are less abundant elements so ensuring that all metal atoms are available for catalysis and not lost in the bulk is an important step towards a more sustainable catalyst design.⁷⁸ These results thus show that fatty acid capped oxo clusters are an exciting material class and they can possess advantages over their larger relatives; oxide nanocrystals. One can thus consider (oxo) clusters for many more catalytic reactions that are currently performed with nanocrystal catalysts. As atomically precise nanocrystal prototypes, oxo clusters are also better suited for mechanistic studies compared to nanocrystals which feature polydispersity and a more irregular surface structure.^{34,79}

3 Conclusion

We have synthesized fatty acid capped zirconium and hafnium oxo clusters with a library of linear and branched carboxylic acids. To prove the structure of the oxo clusters, we used X-ray PDF analysis. We showed that the structure models need to include carboxylate binding groups to obtain excellent refinements. Given the larger scattering power of hafnium, the refinements were even more straightforward for the hafnium oxo clusters. We were thus able to prove the structure of our fatty acid capped oxo clusters, despite the fact that they cannot be crystallized. Given the lack of crystallization, we developed purification protocols akin to the ones for colloidal nanocrystals, reinforcing the notion that these fatty acid capped oxo clusters are atomically precise nanocrystal prototypes, of the smallest conceivable scale.

We further comprehensively characterized the oxo clusters by FTIR, NMR, TGA and ESI-HR-MS. We find that the cationic cluster cores are charge balanced by chelating and bridging carboxylate ligands and that a small excess of carboxylic acid is hydrogen bonded to the cluster core. Using our characterization



tools, we studied carboxylate for carboxylate ligand exchange and found that the **Zr6/Hf6** monomer and the **Zr12/Hf12** dimer structure can be converted into one another, depending on the structure of the final ligand. Finally, we compared the catalytic performance of the clusters with that of larger oxide nanocrystals. We found that, for the same amount of zirconium added, the oxo clusters show a five times higher esterification rate than oxide nanocrystals, due to their higher surface area. Since the oxo clusters are the limit of downscaling oxide nanocrystals, we thus propose them here as appealing materials, or at least as atomically precise model systems.

4 Experimental

4.1 Materials

Zirconium propoxide (70 w% in 1-propanol) and hafnium butoxide (99%) was provided by Sigma-Aldrich and stored in a Straus flask upon arrival. Acetic acid (>99%) was purchased from Sigma-Aldrich and vacuum distilled after which it was stored in a Schlenk flask. Zirconium isopropoxide isopropanol complex (99.9%), propionic acid (>99.5%), butyric acid (99%), hexanoic acid (99%), octanoic acid (99%), decanoic acid (>99.5%), dodecanoic acid (98%), oleic acid (90%), methylbutyric acid (98%), methylheptanoic acid (>98.5%), pivalic acid (99%) and benzyl alcohol (anhydrous, 99.8%) were bought from Sigma-Aldrich and used without any further purification. Acetone, diethylether (BHT stabilized) and dichloromethane (DCM) were bought from Biosolve and used without any further purification. HPLC grade acetonitrile (ACN) was bought from VWR and used without any further purification. All yields reported here are calculated without any co-precipitated molecules unless otherwise specified. Centrifugation was always performed at 5000 rcf for 3 minutes, unless otherwise specified. After purification all clusters are stored in a desiccator.

4.2 Zr12-acetate synthesis

A 20 mL vial was equipped with a septum and cycled three times between argon and vacuum. Zirconium propoxide (2.25 mL, 5 mmol, 1 eq.) was added to the vial, together with dry DCM (5.463 mL). Under stirring, distilled acetic acid (2.287 mL, 40 mmol, 8 eq.) was injected, reaching a total reaction volume of 10 mL and thus a zirconium concentration of 0.5 M. After 12 hours at room temperature (19–24 °C), the crystalline powder is isolated by filtration and further washed with 50 mL of a DCM : acetic acid mixture (4 : 1). Finally, the white powder was dried overnight under high vacuum. The powder was stored in a desiccator. The cluster was obtained as a white solid, with a 98% yield, calculated based on the molecular formula: $\text{Zr}_{12}\text{O}_8(\text{OH})_8(\text{CH}_3\text{COO})_{24} \cdot 6\text{CH}_3\text{COOH} \cdot 3.5\text{DCM}$.⁴

4.3 Zr12-Propionate synthesis

Zirconium propoxide (2.35 mL, 5.22 mmol, 1 eq.) was mixed with propionic acid (3.13 mL, 41.8 mmol, 8 eq.). After overnight reaction the solid was washed with a 1-propanol : propionic acid mixture (ratio 1 : 3.4). After washing, the solid was dried for 8 hours on a Schlenk line. **Zr12-propionate** was obtained as

a white solid with a yield of 51%, calculated based on the molecular formula: $\text{Zr}_{12}\text{O}_8(\text{OH})_8(\text{CH}_3\text{CH}_2\text{COO})_{24} \cdot 6\text{CH}_3\text{CH}_2\text{COOH}$.⁴

4.4 Zr12-Butanoate, hexanoate and octanoate

For butanoate, hexanoate and octanoate capped clusters, zirconium propoxide (2.25 mL, 5 mmol, 1 eq.) was first mixed with DCM, and afterwards 8 equivalents of acid was added. The amount of DCM is adjusted to maintain a Zr concentration of 0.5 M. After stirring for 24 hours at 30 °C the clusters were purified. The butanoate capped cluster was purified by evaporating the solvent using the Schlenk line after which a white waxy solid was obtained. Acetonitrile (15 mL) was added and the clusters were macerated overnight. After centrifugation for 4 minutes at 5000 rcf the supernatant was discarded and the clusters were dried on the Schlenk line for 24 h yielding a white solid. For hexanoate and octanoate clusters the solution was divided over 2 large centrifuge tubes (± 5 mL reaction mixture in each) and acetonitrile (10 mL) was added to each centrifuge tube in order to precipitate the clusters. *Via* centrifugation (5000 rcf, 4 min), the cluster (viscous oil) was separated from the supernatant and both oils were redissolved in DCM (1 mL). To each tube 6 mL acetonitrile was added to precipitate the cluster. This last step was repeated twice and finally, the clusters were dried overnight under vacuum. **Zr12-hexanoate** was obtained as a waxy solid and **Zr12-octanoate** was obtained as a viscous liquid. **Zr12-butanoate** was obtained with a yield of 49%. **Zr12-hexanoate** was obtained as a waxy solid with a yield of 71%. **Zr12-octanoate** was obtained as a viscous oil with a yield of 52%.

4.5 Zr12-Decanoate, dodecanoate and oleate

For decanoic and dodecanoic acid, zirconium propoxide (2.25 mL, 5 mmol, 1 eq.) was mixed with 8 equivalents of acid and diluted with DCM to 0.33 M and 0.25 M respectively in Zr. Decanoic and dodecanoic acids are solids and therefore the concentration had to be lowered in order to dissolve these acids. For oleic acid (which is a liquid), the volume is already 15 mL without DCM. Therefore, no DCM was added and a 40 mL flask was used. After stirring for 24 hours at 30 °C, DCM was evaporated using the Schlenk line for both **Zr12-decanoate** and **Zr12-dodecanoate**, while the oleate sample can be purified as such. The solution for each cluster was divided over 2 centrifuge tubes and ACN (10 mL per tube) was added to precipitate the clusters. *Via* centrifugation (5000 rcf, 4 min), the cluster precipitate was separated from the supernatant and redissolved in DCM (1 mL). To this 6 mL acetone was added to precipitate the cluster. This last step was repeated twice and finally, the clusters were dried under vacuum. **Zr12-decanoate** and **Zr12-dodecanoate** were obtained as waxy solids with a yield of 80% and 93% respectively. Finally, **Zr12-oleate** was obtained as a viscous oil with a yield of 83%.

4.6 Zr6-Methylbutanoate and methylheptanoate

Zirconium propoxide (2.25 mL, 5 mmol, 1 eq.) was mixed with 8 equivalents of acid and diluted with DCM to 0.5 M in Zr. After stirring for 48 hours at 30 °C, the solution was dried under



vacuum and dissolved again in 1 mL DCM. Afterwards, the solution is divided over 2 centrifuge tubes and acetonitrile (4 mL per tube) was added to precipitate the clusters. *Via* centrifugation (5000 rcf, 5 min), the cluster precipitate was separated from the supernatant and redissolved in DCM (1 mL each). To both tubes acetonitrile (double the volume) was added to precipitate the cluster. This last step was repeated twice and finally, the clusters were dried overnight under vacuum. **Zr6**-methylbutanoate was obtained as a white solid with a yield of 93%. **Zr6**-methylheptanoate was obtained as a viscous oil with a yield of 55%.

4.7 Zr12-Acetate ligand exchange

The **Zr12**-acetate cluster (200 mg, 1.4 mmol acetate, 1 eq.) was weighed into a 20 mL vial. The incoming carboxylic acid (1.5 equivalents) was added together with 1 mL of DCM and stirred for 60 min, after which a clear solution was obtained. Subsequently, the solution was further stirred under vacuum at 70 °C for 1 hour. The sample transforms from a heterogeneous mixture into a viscous liquid. 1 purification step was performed as described above for the respective cluster. Yields: butyric acid = 82%, methylbutyric acid = 41%, hexanoic acid = 97.2%, octanoic acid = 95%, and oleic acid = 100%.

4.8 Zr6-Methylbutanoate ligand exchange

The **Zr6**-methylbutanoate cluster (200 mg, 1.27 mmol methylbutanoate, 1 eq.) was weighed into a 20 mL vial. 2 mL DCM together with 3 equivalents of hexanoic acid were added and the solution was stirred for 1 hour at room temperature. Afterwards, the solution is dried under vacuum for 1 hour at 70 °C. One purification cycle was performed by dissolving the sample in 0.5 mL DCM and precipitating it by adding 2 mL ACN. This turbid solution was centrifuged at 5000 rcf for 4 minutes. The whole process was repeated once to ensure full exchange. After the second exchange step the sample was purified twice as described above. **Zr12**-hexanoate was obtained as a viscous liquid with a yield of 56.5%.

4.9 Hf12-Acetate, oleate and methylbutanoate

Hafnium *n*-butoxide (1.9 mL, 5 mmol, 1 eq.) was mixed with 8 equivalents of acetic, oleic or methylbutyric acid. Prior to mixing dry DCM was added to the hafnium precursor in order to set $[Hf] = 0.5$ M in the whole reaction mixture. **Hf12**-acetate and **Hf12**-oleate were reacted for 24 h at 30 °C. For **Hf12**-oleate no dry DCM was added. The **Hf6**-methylbutanoate synthesis was kept at 30 °C for 48 h. Purification was performed according to the zirconium clusters for the respective acids. **Hf12**-acetate and **Hf6**-methylbutanoate were obtained as white solids with yields of 100% and 50%, respectively. **Hf12**-oleate was obtained as a viscous oil with a yield of 37%.

4.10 Hf12-Acetate ligand exchange

The **Hf12**-acetate cluster (200 mg, 1.26 mmol acetate, 1 eq.) was weighed into a 20 mL vial. 1.5 equivalents (1.89 mmol) of the new carboxylic acid, 597 μ L oleic acid or 301 μ L

methylheptanoic acid, was added together with 1 mL of DCM and stirred for 60 min, after which a clear solution was obtained. Subsequently, the solution was further stirred under vacuum at 70 °C for 1 hour. The sample transforms from a heterogeneous mixture into a viscous liquid. Purification was performed as described above for the respective cluster. The exchanged clusters were obtained as a viscous oil with a yield of 70% for the oleate exchanged cluster and 37% for the methylheptanoic acid exchanged clusters.

4.11 Nanocrystal synthesis

The zirconia nanocrystals were synthesized according to Garnweitner *et al.* with few modifications.⁷⁷ $Zr(OiPr)_4 \cdot iPrOH$ (6.6 g, 17 mmol) was mixed with benzyl alcohol (60 mL, 580 mmol) in a Parr bomb inside a nitrogen filled glovebox. The Parr bomb was heated for 48 h at 210 °C. After the reaction, a white powder was obtained *via* centrifugation. This was washed with 60 mL diethyl ether followed by centrifugation three times (in 4 centrifuge tubes with 15 mL each). The white solid is dispersed in 40 mL toluene (4 centrifuge tubes of 10 mL each). To each centrifuge tube oleic acid was added (600 μ L, 1.9 mmol) to functionalize the nanocrystal surface and the centrifuge tubes were stirred and sonicated for 30 minutes. This procedure resulted in a cloudy solution of nanocrystals. The insolubles were removed by centrifugation and a transparent solution of the soluble nanocrystals in the supernatant was collected in 2 centrifuge tubes and the nanocrystals were precipitated using a 1 : 1 ratio of nanocrystal solution and acetone. The collected nanocrystals were then purified *via* three precipitation/redissolving cycles; the nanocrystals in 10 mL toluene were split off into two centrifuge tubes and precipitated by adding 15 mL acetone to each of the two tubes. The nanocrystals were obtained as a white solid with a yield of 43%.

4.12 Catalytic experiments

10 mol% (in Zr) of the catalyst was dispersed in a 4 mL solution of 200 mM oleic acid and 400 mM EtOH in *o*-dichlorobenzene. 10 mol% Zr corresponds to 54.36 mg of the **Zr12**-oleate cluster (0.0067 mmol clusters or 0.08 mmol Zr) and 9.86 mg ZrO_2 (0.08 mmol Zr). The reaction mixture is then heated to 130 °C for 5 h in a closed vial. Aliquots are taken every 30 min.

4.13 General instrumentation

Nuclear magnetic resonance (NMR) measurements were recorded at 298 K on a Bruker UltraShield 500 spectrometer operating at a frequency of 500.13 MHz. The IR spectra were recorded on a PerkinElmer spectrum 2 ATR-FTIR with a diamond crystal.

4.14 Mass spectrometry

ESI mass spectra were acquired using a Bruker maXis4G or maXis II high resolution mass spectrometer equipped with an electrospray ionization source. The cluster was dissolved in a mixture of ACN : DCM, in which the amount of ACN was maximized while avoiding precipitation of the clusters. The



samples were directly introduced into the instrument at a rate of $6 \mu\text{L min}^{-1}$ using a syringe pump. The heated capillary temperature was $200 \text{ }^\circ\text{C}$ and the capillary voltage was 3.6 kV . The samples were analyzed in positive ion mode. DataAnalysis 4.4 from Bruker was used to process the raw data.

4.15 Synchrotron X-ray total scattering experiments

The samples were prepared in a 1 mm polyamide kapton tube and were measured either at beamline 11-ID-BM at the Advanced Photon Source, Argonne National Laboratory, USA or at beamline P21.1 at DESY in Hamburg, Germany. X-ray total scattering data were collected at room temperature in rapid acquisition mode, using a PerkinElmer digital X-ray flat panel amorphous silicon detector (2048×2048 pixels and $200 \times 200 \mu\text{m}$ pixel size) with a sample-to-detector distance of 180 mm (11-ID-BM) or 380 mm (P21.1). The incident wavelength of the X-rays was $\lambda = 0.2110 \text{ \AA}$ (11-ID-BM) or 0.1220 \AA (P21.1). Calibration of the experimental setup was performed using a Ni standard.

4.16 Analysis of synchrotron X-ray total scattering data

Raw 2D data were corrected for geometrical effects and polarization, and then azimuthally integrated to produce 1D scattering intensities *versus* the magnitude of the momentum transfer Q (where $Q = 4\pi \sin \theta / \lambda$ for elastic scattering) using pyFAI and xpdtools.^{80,81} The program xPDFsuite with PDFgetX3 was used to perform the background subtraction, further corrections, and normalization to obtain the reduced total scattering structure function $F(Q)$, and Fourier transformation to obtain the pair distribution function (PDF), $G(r)$.^{82,83} For data reduction, the following parameters were used after proper background subtraction: $Q_{\text{min}} = 0.8 \text{ \AA}^{-1}$, $Q_{\text{max}} = 22 \text{ \AA}^{-1}$, and $R_{\text{poly}} = 0.9 \text{ \AA}$. We used the actual chemical composition of each cluster (inorganic core + charge balancing ligands) for data reduction. Modeling and fitting were carried out using Diffpy-CMI.⁸⁴ The Debye scattering equation was used to generate the calculated PDF from discrete structure models. The structure models are supplied as xyz files in the ESI.† The refinements were carried out by refining the scale factor, isotropic atomic displacement parameters (Uiso), and delta2 (coefficient for the $1/r^2$ contribution to the peak sharpening). The exponentially dampening sine-wave contribution was calculated according to the following equation:

$$y = Ae^{-\left(\frac{r-r_0}{2\sigma_{\text{eff}}}\right)^2} \sin\left(2\pi\left(\frac{r}{\lambda} - \phi\right)\right)$$

A – amplitude of oscillation, r – the distance in the PDF, λ – wavelength, ϕ – phase shift, σ – effective dampening with $\sigma_{\text{eff}} = \sigma/a$ for $r < r_0$ and $\sigma_{\text{eff}} = \sigma \times a$ for $r > r_0$, a is the asymmetry parameter. and r_0 is not a physical parameter in real space and is used to describe different dampening behaviors.⁶⁴

Data availability

The data underlying the figures is available on the Zenodo platform, DOI: [10.5281/zenodo.7373015](https://doi.org/10.5281/zenodo.7373015). In the ESI,† further

characterization *via* PDF, NMR, MS and FTIR is available. The ESI† also contains the structure models that we used for the PDF refinements.

Author contributions

D. Van den Eynden: synthetic and analytic methodology, investigation, writing – original draft, visualization. R. Pokratath: PDF analysis, TGA data acquisition, writing – original draft, visualization. J. Pulparayil Mathew: validation, investigation (catalysis). E. Goossens: TGA data acquisition. K. De Buysser: supervision. J. De Roo: conceptualization, supervision, writing – review and editing, visualization.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

The authors thank Prof. Dr Kirsten Jensen and Dr Francisco De Azambuja for fruitful discussions and scientific assistance. The authors thank Michael Pfeffer for MS analysis. The authors thank SNF NCCR molecular system engineering (182895), the University of Basel, Research Foundation–Flanders (Project No. 1S11721N) and the SNF Eccellenza funding scheme (project number: 194172) for funding. We acknowledge DESY (Hamburg, Germany), a member of the Helmholtz Association HGF, for the provision of experimental facilities. Parts of this research study were carried out at PETRA III and we would like to thank Dr Soham Banerjee for assistance in using beamline P21.1. This research used resources of the Advanced Photon Source, a U.S. Department of Energy (DOE) Office of Science user operated facility for the DOE Office of Science by the Argonne National Laboratory under Contract No. DE-AC02-06CH11357. We would like to thank Dr Leighanne Gallington for assistance in using beamline 11-ID-B (proposal: GUP-79568).

Notes and references

- W. Baek, H. Chang, M. S. Bootharaju, J. H. Kim, S. Park and T. Hyeon, *JACS Au*, 2021, **1**, 1849–1859.
- D. Van den Eynden, R. Pokratath and J. De Roo, *Chem. Rev.*, 2022, **122**, 10538–10572.
- G. Kickelbick and U. Schubert, *Chem. Ber.*, 1997, **130**, 473–478.
- M. Puchberger, F. R. Kogler, M. Jupa, S. Gross, H. Fric, G. Kickelbick and U. Schubert, *Eur. J. Inorg. Chem.*, 2006, **2006**, 3283–3293.
- T. Frot, S. Cochet, G. Laurent, C. Sassoye, M. Popall, C. Sanchez and L. Rozes, *Eur. J. Inorg. Chem.*, 2010, **2010**, 5650–5659.
- G. Schmid, *Chem. Soc. Rev.*, 2008, **37**, 1909–1930.
- A. N. Beecher, X. Yang, J. H. Palmer, A. L. LaGrassa, P. Juhas, S. J. L. Billinge and J. S. Owen, *J. Am. Chem. Soc.*, 2014, **136**, 10645–10653.



- 8 D. C. Gary, S. E. Flowers, W. Kaminsky, A. Petrone, X. Li and B. M. Cossairt, *J. Am. Chem. Soc.*, 2016, **138**, 1510–1513.
- 9 B. S. Mitchell, S. M. Krajewski, J. A. Kephart, D. Rogers, W. Kaminsky and A. Velian, *JACS Au*, 2022, **2**, 92–96.
- 10 J.-Y. Huang, H. Xu, E. Peretz, D.-Y. Wu, C. K. Ober and T. Hanrath, *Chem. Mater.*, 2019, **31**, 10017–10022.
- 11 H. Han, S. Kallakuri, Y. Yao, C. B. Williamson, D. R. Nevers, B. H. Savitzky, R. S. Skye, M. Xu, O. Voznyy, J. Dshemuchadse, L. F. Kourkoutis, S. J. Weinstein, T. Hanrath and R. D. Robinson, *Nat. Mater.*, 2022, **21**, 518–525.
- 12 H. Chang, B. H. Kim, H. Y. Jeong, J. H. Moon, M. Park, K. Shin, S. I. Chae, J. Lee, T. Kang, B. K. Choi, J. Yang, M. S. Bootharaju, H. Song, S. H. An, K. M. Park, J. Y. Oh, H. Lee, M. S. Kim, J. Park and T. Hyeon, *J. Am. Chem. Soc.*, 2019, **141**, 7037–7045.
- 13 D. C. Gary, M. W. Terban, S. J. L. Billinge and B. M. Cossairt, *Chem. Mater.*, 2015, **27**, 1432–1441.
- 14 S. D. Pike, E. R. White, M. S. P. Shaffer and C. K. Williams, *Nat. Commun.*, 2016, **7**, 13008.
- 15 C. Falaise, H. A. Neal and M. Nyman, *Inorg. Chem.*, 2017, **56**, 6591–6598.
- 16 S. Gross, *J. Mater. Chem.*, 2011, **21**, 15853–15861.
- 17 U. Schubert, *Chem. Soc. Rev.*, 2011, **40**, 575–582.
- 18 J. Moons, F. de Azambuja, J. Mihailovic, K. Kozma, K. Smiljanic, M. Amiri, T. Cirkovic Velickovic, M. Nyman and T. N. Parac-Vogt, *Angew. Chem., Int. Ed.*, 2020, **59**, 9094–9101.
- 19 Y. Zhang, F. de Azambuja and T. N. Parac-Vogt, *Catal. Sci. Technol.*, 2022, **12**, 3190–3201.
- 20 L. Wu, J. Liu, M. Vockenhuber, Y. Ekinci and S. Castellanos, *Eur. J. Inorg. Chem.*, 2019, **2019**, 4136–4141.
- 21 S. Kataoka and K. Sue, *Eur. J. Inorg. Chem.*, 2022, **2022**, e202200050.
- 22 J. A. Sommers, L. Palys, N. P. Martin, D. B. Fast, M. Amiri and M. Nyman, *J. Am. Chem. Soc.*, 2022, **144**, 2816–2824.
- 23 T. Krämer, F. Tuna and S. D. Pike, *Chem. Sci.*, 2019, **10**, 6886–6898.
- 24 J. H. Cavka, S. Jakobsen, U. Olsbye, N. Guillou, C. Lamberti, S. Bordiga and K. P. Lillerud, *J. Am. Chem. Soc.*, 2008, **130**, 13850–13851.
- 25 M. H. Beyzavi, R. C. Klet, S. Tussupbayev, J. Borycz, N. A. Vermeulen, C. J. Cramer, J. F. Stoddart, J. T. Hupp and O. K. Farha, *J. Am. Chem. Soc.*, 2014, **136**, 15861–15864.
- 26 A. V. Dighe, L. Huelsenbeck, R. R. Bhawnani, P. Verma, K. H. Stone, M. R. Singh and G. Giri, *JACS Au*, 2022, **2**, 453–462.
- 27 Z. Hu, Y. Wang and D. Zhao, *Chem. Soc. Rev.*, 2021, **50**, 4629–4683.
- 28 L. Feng, J. Pang, P. She, J.-L. Li, J.-S. Qin, D.-Y. Du and H.-C. Zhou, *Adv. Mater.*, 2020, **32**, 2004414.
- 29 Y. Zhang, F. de Azambuja and T. N. Parac-Vogt, *Coord. Chem. Rev.*, 2021, **438**, 213886.
- 30 U. Schubert, *Coord. Chem. Rev.*, 2022, **469**, 214686.
- 31 U. Schubert, *Coord. Chem. Rev.*, 2017, **350**, 61–67.
- 32 F. R. Kogler, M. Jupa, M. Puchberger and U. Schubert, *J. Mater. Chem.*, 2004, **14**, 3133–3138.
- 33 G. Kickelbick, P. Wiede and U. Schubert, *Inorg. Chim. Acta*, 1999, **284**, 1–7.
- 34 Z. Hens and J. De Roo, *J. Am. Chem. Soc.*, 2020, **142**, 15627–15637.
- 35 J. De Roo, F. Van den Broeck, K. De Keukeleere, J. C. Martins, I. Van Driessche and Z. Hens, *J. Am. Chem. Soc.*, 2014, **136**, 9650–9657.
- 36 K. De Keukeleere, S. Coucke, E. De Canck, P. Van Der Voort, F. Delpesch, Y. Coppel, Z. Hens, I. Van Driessche, J. S. Owen and J. De Roo, *Chem. Mater.*, 2017, **29**, 10233–10242.
- 37 H. L. Nguyen, F. Gándara, H. Furukawa, T. L. H. Doan, K. E. Cordova and O. M. Yaghi, *J. Am. Chem. Soc.*, 2016, **138**, 4330–4333.
- 38 Y. Yang, H. Qin, M. Jiang, L. Lin, T. Fu, X. Dai, Z. Zhang, Y. Niu, H. Cao, Y. Jin, F. Zhao and X. Peng, *Nano Lett.*, 2016, **16**, 2133–2138.
- 39 D. Doblás, T. Kister, M. Cano-Bonilla, L. González-García and T. Kraus, *Nano Lett.*, 2019, **19**, 5246–5252.
- 40 J. De Roo, Y. Justo, K. De Keukeleere, F. Van den Broeck, J. C. Martins, I. Van Driessche and Z. Hens, *Angew. Chem., Int. Ed.*, 2015, **54**, 6488–6491.
- 41 J. De Roo, I. Van Driessche, J. C. Martins and Z. Hens, *Nat. Mater.*, 2016, **15**, 517–521.
- 42 J. De Roo, Z. Zhou, J. Wang, L. Deblock, A. J. Crosby, J. S. Owen and S. S. Nonnenmann, *Chem. Mater.*, 2018, **30**, 8034–8039.
- 43 Y. Okada, K. Ishikawa, N. Maeta and H. Kamiya, *Chem.–Eur. J.*, 2018, **24**, 1853–1858.
- 44 L. Deblock, E. Goossens, R. Pokratath, K. De Buysser and J. De Roo, *JACS Au*, 2022, **2**, 711–722.
- 45 J. Kreutzer, M. Puchberger, C. Artner and U. Schubert, *Eur. J. Inorg. Chem.*, 2015, **2015**, 2145–2151.
- 46 A. E. Platero-Prats, A. Mavrandonakis, L. C. Gallington, Y. Liu, J. T. Hupp, O. K. Farha, C. J. Cramer and K. W. Chapman, *J. Am. Chem. Soc.*, 2016, **138**, 4178–4185.
- 47 H. Xu, S. Sommer, N. L. N. Broge, J. Gao and B. B. Iversen, *Chem.–Eur. J.*, 2019, **25**, 2051–2058.
- 48 T. D. Bennett, T. K. Todorova, E. F. Baxter, D. G. Reid, C. Gervais, B. Bueken, B. Van de Voorde, D. De Vos, D. A. Keen and C. Mellot-Draznieks, *Phys. Chem. Chem. Phys.*, 2016, **18**, 2192–2201.
- 49 F. C. N. Firth, M. W. Gaultois, Y. Wu, J. M. Stratford, D. S. Keeble, C. P. Grey and M. J. Cliffe, *J. Am. Chem. Soc.*, 2021, **143**, 19668–19683.
- 50 X. Yang, A. S. Masadeh, J. R. McBride, E. S. Božin, S. J. Rosenthal and S. J. L. Billinge, *Phys. Chem. Chem. Phys.*, 2013, **15**, 8480–8486.
- 51 R. Pokratath, D. Van den Eynden, S. R. Cooper, J. K. Mathiesen, V. Waser, M. Devereux, S. J. L. Billinge, M. Meuwly, K. M. O. Jensen and J. De Roo, *JACS Au*, 2022, **2**, 827–838.
- 52 S. Banerjee, C.-H. Liu, K. M. O. Jensen, P. Juhas, J. D. Lee, M. Tofanelli, C. J. Ackerson, C. B. Murray and S. J. L. Billinge, *Acta Crystallogr., Sect. A: Found. Adv.*, 2020, **76**, 24–31.



- 53 S. R. Cooper, R. O. Candler, A. G. Cosby, D. W. Johnson, K. M. O. Jensen and J. E. Hutchison, *ACS Nano*, 2020, **14**, 5480–5490.
- 54 S. K. Pradhan, Y. Mao, S. S. Wong, P. Chupas and V. Petkov, *Chem. Mater.*, 2007, **19**, 6180–6186.
- 55 R. U. Ichikawa, A. G. Roca, A. López-Ortega, M. Estrader, I. Peral, X. Turrillas and J. Nogués, *Small*, 2018, **14**, 1800804.
- 56 E. Dhaene, R. Pokratath, O. Aalling-Frederiksen, K. M. O. Jensen, P. F. Smet, K. De Buysser and J. De Roo, *ACS Nano*, 2022, **16**, 7361–7372.
- 57 K. M. O. Jensen, P. Juhas, M. A. Tofanelli, C. L. Heinecke, G. Vaughan, C. J. Ackerson and S. J. L. Billinge, *Nat. Commun.*, 2016, **7**, 11859.
- 58 A. S. Anker, T. L. Christiansen, M. Weber, M. Schmiele, E. Brok, E. T. S. Kjær, P. Juhás, R. Thomas, M. Mehring and K. M. O. Jensen, *Angew. Chem., Int. Ed.*, 2021, **60**, 20407–20416.
- 59 C. B. Williamson, D. R. Nevers, A. Nelson, I. Hadar, U. Banin, T. Hanrath and R. D. Robinson, *Science*, 2019, **363**, 731–735.
- 60 Y. Shen, M. Y. Gee and A. B. Greytak, *Chem. Commun.*, 2017, **53**, 827–841.
- 61 S. J. L. Billinge, *Philos. Trans. R. Soc., A*, 2019, **377**, 20180413.
- 62 S. J. L. Billinge and I. Levin, *Science*, 2007, **316**, 561–565.
- 63 T. L. Christiansen, S. R. Cooper and K. M. O. Jensen, *Nanoscale Adv.*, 2020, **2**, 2234–2254.
- 64 M. Zobel, R. B. Neder and S. A. J. Kimber, *Science*, 2015, **347**, 292–294.
- 65 Z. Chen, Z. Chen, O. K. Farha and K. W. Chapman, *J. Am. Chem. Soc.*, 2021, **143**, 8976–8980.
- 66 C. Hennig, S. Weiss, W. Kraus, J. Kretzschmar and A. C. Scheinost, *Inorg. Chem.*, 2017, **56**, 2473–2480.
- 67 G. Trimmel, S. Gross, G. Kickelbick and U. Schubert, *Appl. Organomet. Chem.*, 2001, **15**, 401–406.
- 68 G. Kickelbick and U. Schubert, *J. Chem. Soc., Dalton Trans.*, 1999, **8**, 1301–1306.
- 69 B. Nateghi, I. Boldog, K. V. Domasevitch and C. Janiak, *CrystEngComm*, 2018, **20**, 5132–5136.
- 70 T. J. Boyle, L. Ottley and M. Rodriguez, *Polyhedron*, 2005, **24**, 1727–1738.
- 71 X. Qian, J. Li and X. Peng, *Chem. Mater.*, 2022, **34**, 7006–7014.
- 72 P. Piszczek, A. Radtke, A. Grodzicki, A. Wojtczak and J. Chojnacki, *Polyhedron*, 2007, **26**, 679–685.
- 73 J. Zhang, H. Zhang, W. Cao, Z. Pang, J. Li, Y. Shu, C. Zhu, X. Kong, L. Wang and X. Peng, *J. Am. Chem. Soc.*, 2019, **141**, 15675–15683.
- 74 G. B. Deacon and R. J. Phillips, *Coord. Chem. Rev.*, 1980, **33**, 227–250.
- 75 B. Fritzing, R. K. Capek, K. Lambert, J. C. Martins and Z. Hens, *J. Am. Chem. Soc.*, 2010, **132**, 10195–10201.
- 76 J. De Roo, N. Yazdani, E. Drijvers, A. Lauria, J. Maes, J. S. Owen, I. Van Driessche, M. Niederberger, V. Wood, J. C. Martins, I. Infante and Z. Hens, *Chem. Mater.*, 2018, **30**, 5485–5492.
- 77 G. Garnweitner, L. M. Goldenberg, O. V. Sakhno, M. Antonietti, M. Niederberger and J. Stumpe, *Small*, 2007, **3**, 1626–1632.
- 78 H. E. Suess and H. C. Urey, *Rev. Mod. Phys.*, 1956, **28**, 53–74.
- 79 E. Drijvers, J. De Roo, J. C. Martins, I. Infante and Z. Hens, *Chem. Mater.*, 2018, **30**, 1178–1186.
- 80 G. Ashiotis, A. Deschildre, Z. Nawaz, J. P. Wright, D. Karkoulis, F. E. Picca and J. Kieffer, *J. Appl. Crystallogr.*, 2015, **48**, 510–519.
- 81 Z. X. Wright, *J. Synchrotron Radiat.*, 2017, **24**(2), 506–508.
- 82 P. Juhas, T. Davis, C. L. Farrow and S. J. L. Billinge, *J. Appl. Crystallogr.*, 2013, **46**, 560–566.
- 83 X. Yang, P. Juhas, C. L. Farrow and S. J. Billinge, 2014, arXiv preprint arXiv:1402.3163.
- 84 P. Juhas, C. L. Farrow, X. Yang, K. R. Knox and S. J. L. Billinge, *Acta Crystallogr., Sect. A: Found. Adv.*, 2015, **71**, 562–568.

