

Cite this: *Chem. Sci.*, 2021, 12, 3660

All publication charges for this article have been paid for by the Royal Society of Chemistry

New atomically precise M_1Ag_{21} ($M = Au/Ag$) nanoclusters as excellent oxygen reduction reaction catalysts†

Xuejuan Zou,^{ac} Shuping He,^{ac} Xi Kang,^{id ac} Shuang Chen,^{bc} Haizhu Yu,^{id ac} Shan Jin,^{id *bc} Didier Astruc,^{id *d} and Manzhou Zhu^{id *abc}

By introducing 1,1'-bis-(diphenylphosphino)ferrocene (dppf) as an activating ligand, two novel nanoclusters, M_1Ag_{21} ($M = Au/Ag$), have been controllably synthesized and structurally characterized. The atomically precise structures of the M_1Ag_{21} nanoclusters were determined by SCXC and further confirmed by ESI-TOF-MS, TGA, XPS, DPV, and FT-IR measurements. The M_1Ag_{21} nanoclusters supported on activated carbon (C) are exploited as efficient oxygen reduction reaction (ORR) catalysts in alkaline solutions. Density functional theory (DFT) calculations verify that the catalytic activities of the two cluster-based systems originate from the significant ensemble synergy effect between the M_{13} kernel and dppf ligand in M_1Ag_{21} . This work sheds lights on the preparation of cluster-based electrocatalysts and other catalysts that are activated and modified by peripheral ligands.

Received 26th October 2020

Accepted 5th January 2021

DOI: 10.1039/d0sc05923d

rsc.li/chemical-science

1. Introduction

The increasing demand for energy and environmental sustainability has accelerated the research on efficient, low-cost and environmentally friendly alternative energy conversion systems.^{1,2} Therefore, the development of renewable energy technology becomes extremely important for which good oxygen reduction reaction (ORR) catalysts are key examples. 1,1'-Bis-(diphenylphosphino)-ferrocene (dppf),³⁻⁵ an excellent donor and bulky bisphosphine ligand as well as a derivative of ferrocene, has displayed excellent electrochemical performance in some dppf-containing complexes,⁶⁻⁹ indicating that dppf-functionalized complexes/nanoparticles may have more potential applications in electrochemistry.

Metal nanoclusters, with small sizes and atomically precise structures,¹⁰ have attracted much attention in various fields, such as catalysis, sensing, optics, and electrochemistry.¹¹⁻¹⁶

With the development of the synthetic chemistry of metal nanoclusters, a number of reports have indicated that it is possible to precisely control the size and shape of nanocluster structures, in other words, that means the surface ligands (*i.e.*, thiolates, phosphines, alkynyls, *etc.*) could be of great relevance to the geometrical structures and potential performances of nanoclusters.¹⁷⁻²⁴ For example, for different liganded- Au_{25} nanoclusters, spherical $Au_{25}(SR)_{18}$ (ref. ²⁵ and ²⁶) shows a larger total current density than rod-shaped $[Au_{25}(PPh_3)_{10}(SR')_5Cl_2]^{2+}$ (ref. ²⁷) for electrochemical CO_2 reduction;²⁸ however, higher conversion can be observed for rod-shaped Au_{25} than spherical Au_{25} in photocatalytic oxidation of benzyl amine reaction.²⁹ In addition, Jin *et al.*, observed that the $Au_{25}(SNAp)_{18}$ nanoclusters exhibit significantly improved thermal stability, antioxidant properties and better catalytic activity and selectivity in the Ullmann heterocoupling reactions than $Au_{25}(PET)_{18}$ nanoclusters.¹⁷ Similar results can also be seen in Au_{28} , Au_{38} and so on.³⁰⁻³² Besides, the bisphosphine ligand can also be used for regulating the structures and properties in metal nanoclusters. Recently, Zhu's group reported ligand-exchange induced transformation from bi-icosahedral $Pt_2Ag_{23}(PPh_3)_{10}Cl_7$ to monoicosahedral $Pt_1Ag_{24}(SPhMe_2)_{18}$, and then to bare-icosahedral $Pt_1Ag_{12}(dppm)_5(SPhMe_2)_2$; by introducing the dppm diphosphine ligand, the surface rigidity of the clusters has increased and promoted the enhancement of luminescence further.^{20,33,34} In view of the unique advantages of dppf, will these stereo-electronic ligand properties strongly and favorably impact the sizes,³⁵⁻³⁷ and electronic properties⁶⁻⁸ of metal nanoclusters, and therefore their performances in catalysis?

Herein, by introducing dppf as an activating ligand, two novel nanoclusters, $[Ag_{22}(dppf)_3(SAdm)_{12}](BPh_4)_2$ (**Ag22**) and

^aDepartment of Chemistry, Centre for Atomic Engineering of Advanced Materials, Key Laboratory of Chemistry for Inorganic/Organic Hybrid Functionalized Materials, Anhui University, Hefei, Anhui, 230601, P. R. China. E-mail: zmx@ahu.edu.cn

^bInstitutes of Physical Science and Information Technology, Anhui University, Hefei, Anhui, 230601, P. R. China. E-mail: jinshan@ahu.edu.cn

^cKey Laboratory of Structure and Functional Regulation of Hybrid Materials, Anhui University, Ministry of Education, Hefei, 230601, P. R. China

^dUniversité de Bordeaux, 33405 Talence Cedex, France. E-mail: didier.astruc@u-bordeaux.fr

† Electronic supplementary information (ESI) available: Synthesis, characterization, and electrochemical measurements. Details and crystal data of $Ag_{22}(dppf)_3(SAdm)_{12}(BPh_4)_2$ and $Au_1Ag_{21}(dppf)_3(SAdm)_{12}(BPh_4)_2$ (cif). CCDC 2031749 and 2031750. For ESI and crystallographic data in CIF or other electronic format see DOI: 10.1039/d0sc05923d



[Au₁Ag₂₁(dppf)₃(SAdm)₁₂](BPh₄)₂ (**Au₁Ag₂₁**), are reported as their bis-BPh₄ salts. The atomically precise structures of these two nanoclusters are determined by single crystal X-ray crystallography (SCXC), and further confirmed by electrospray ionization time-of-flight mass spectrometry (ESI-TOF-MS), thermogravimetric analysis (TGA), X-ray photoelectron spectroscopy (XPS), and Fourier transform infrared (FT-IR) measurements. Both nanoclusters have the typical icosahedral M₁₃ (M = Au/Ag) kernel protected by three Ag₃(SR)₄ motifs and three dppf ligands. The incorporation of dppf not only regulates the structures of these two nanoclusters, but also drastically improves the ORR performances in alkaline media. Indeed, both nanoclusters supported on activated carbon (C) exhibit excellent ORR activity, and **Au₁Ag₂₁(dppf)₃(SAdm)₁₂/C** shows a better performance than **Ag₂₂(dppf)₃(SAdm)₁₂/C**. To the best of our knowledge, this is the first study of the ORR properties of dppf-functionalized metal nanoclusters, and the observations in this work will provide some guidance for the future development of nanoclusters in catalysis and electrocatalysis.

2. Experimental methods

2.1. Materials

All the following reagents and solvents are commercially available, and all of them were used without further purification: silver nitrate (AgNO₃, 99%), hydrogen tetrachloroaurate(III) (HAuCl₄·3H₂O, 99.99%), 1,1'-bis(diphenylphosphino)ferrocene (dppf, 97%, Sigma-Aldrich), 1-adamantanethiol (1-AdmSH, 97%, Sigma-Aldrich), sodium borohydride (NaBH₄, 98%, Sigma-Aldrich), sodium tetraphenylborate (NaBPh₄, 98%), methyl sulfide (C₂H₆S, 99%, Sigma-Aldrich), methanol (CH₃OH, HPLC, Aldrich), acetonitrile (CH₃CN, HPLC, Aldrich), dichloromethane (CH₂Cl₂, HPLC grade, Aldrich), toluene (Tol, HPLC grade, Aldrich), and *n*-hexane (hex, HPLC grade, Aldrich). Pure water was obtained from Wahaha Co. Ltd.

2.2. Synthesis of Ag₂₂(dppf)₃(SAdm)₁₂(BPh₄)₂ nanocluster

60 mg AgNO₃ (0.35 mmol) was dissolved in 5 mL methanol using ultrasound treatment, and this solution was added into 15 mL toluene. After 15 min, 50 mg 1,1'-bis(diphenylphosphino)ferrocene (0.09 mmol) and 50 mg 1-adamantanethiol (0.3 mmol) were simultaneously added into the solution; 30 min later, 1 mL of an aqueous solution of NaBH₄ (25 mg mL⁻¹, (0.66 mmol)) was added, and the reaction mixture was stirred for 5 hours at room temperature. The crude product was obtained by rotary evaporation, then washed three times with *n*-hexane and dissolved in CH₂Cl₂ again. Then, 20 mg NaBPh₄ (0.058 mmol) in 2 mL CH₃OH was added to the solution in order to substitute the anion in the cluster for crystallization. A red precipitate was collected by centrifugation (5 min at ~7000 rpm), then washed twice with excess *n*-hexane and again collected by centrifugation. Red crystals were formed from a CH₂Cl₂/hex (volume ratio = 1 : 3) or toluene/methanol solution of the clusters at room temperature after three days. The yield of the **Ag₂₂** nanocluster (crystals) was 10% (about 9.7 mg) based on Ag atoms.

2.3. Synthesis of the dppf-Au complexes

Methyl sulfide (2 mL, 27 mmol) was slowly added to CH₃OH solution (15 mL) of 1 mmol (400 mg) HAuCl₄·3H₂O until a milky white precipitation formed; then the solution was stirred for half an hour and dried for later use. This product was dissolved in 15 mL CH₂Cl₂ again, and 555 mg (1 mmol) of dppf was added to the solution. The reaction mixture was stirred for 2 h, then the dppf-Au complex obtained was dried, and about 280 mg of complex was obtained for use.

2.4. Synthesis of the Au₁Ag₂₁(dppf)₃(SAdm)₁₂(BPh₄)₂ nanocluster

The experimental procedure for the synthesis of the Au₁Ag₂₁(dppf)₃(SAdm)₁₂(BPh₄)₂ nanocluster was the same as that used for the synthesis of Ag₂₂(dppf)₃(SAdm)₁₂(BPh₄)₂ except that 20 mg dppf-Au was used instead of 50 mg dppf. Comparing with the **Ag₂₂** nanocluster, more impurities were present in the crude product, and the yield of pure **Au₁Ag₂₁** nanocluster (crystals) was only 5% (about 5.2 mg) based on Ag atoms.

2.5. Preparation of electrocatalysts of Ag₂₂(dppf)₃(SAdm)₁₂/C and Au₁Ag₂₁(dppf)₃(SAdm)₁₂/C

100 mg of activated carbon was first dispersed into 20 mL CH₂Cl₂. Then, 10 mg **Ag₂₂** or 10 mg **Au₁Ag₂₁** nanocluster was dissolved in 10 mL CH₂Cl₂, and this solution was added dropwise into the above activated carbon dispersion at constant stirring at ambient temperature for 10 h until the colour of the supernatant disappeared. CH₂Cl₂ was removed by rotary evaporation, forming catalysts for which the nanoclusters were supported on activated carbon with a loading of 10%. These catalysts are named **Ag₂₂(dppf)₃(SAdm)₁₂/C** and **Au₁Ag₂₁(dppf)₃(SAdm)₁₂/C**.

2.6. Preparation of electrocatalysts of Ag₂₂ and Au₁Ag₂₁ nanoclusters with different loadings

100 mg of activated carbon was first dispersed into 20 mL CH₂Cl₂. Then, 5 mg or 20 mg or 30 mg **Ag₂₂** nanocluster was dissolved in 10 mL CH₂Cl₂, and this solution was added dropwise into the above activated carbon dispersion at constant stirring at ambient temperature for 10 h until the colour of the supernatant disappeared. CH₂Cl₂ was removed by rotary evaporation, forming catalysts for which the nanoclusters were supported on activated carbon with a loading of 5%, 20%, and 30%. These catalysts are named **Ag₂₂(dppf)₃(SAdm)₁₂/C-5%**, **Ag₂₂(dppf)₃(SAdm)₁₂/C-20%**, and **Ag₂₂(dppf)₃(SAdm)₁₂/C-30%**. Also, the preparation of catalysts of **Au₁Ag₂₁(dppf)₃(SAdm)₁₂/C-5%**, **Au₁Ag₂₁(dppf)₃(SAdm)₁₂/C-20%**, and **Au₁Ag₂₁(dppf)₃(SAdm)₁₂/C-30%** is similar to the method above except for replacing **Ag₂₂** with **Au₁Ag₂₁**.

2.7. Preparation of electrocatalysts of Ag₂₂(dppf)₃(SAdm)₁₂/C and Au₁Ag₂₁(dppf)₃(SAdm)₁₂/C at different temperatures

Ag₂₂(dppf)₃(SAdm)₁₂/C or **Au₁Ag₂₁(dppf)₃(SAdm)₁₂/C** was calcined in a tube furnace for two hours in Ar at different temperatures. These systems are named **Ag₂₂(dppf)₃(SAdm)₁₂/**



C-150 °C, $\text{Ag}_{22}(\text{dppf})_3(\text{SAdm})_{12}/\text{C-200 °C}$, $\text{Ag}_{22}(\text{dppf})_3(\text{SAdm})_{12}/\text{C-300 °C}$, $\text{Au}_1\text{Ag}_{21}(\text{dppf})_3(\text{SAdm})_{12}/\text{C-150 °C}$, $\text{Au}_1\text{Ag}_{21}(\text{dppf})_3(\text{SAdm})_{12}/\text{C-200 °C}$, and $\text{Au}_1\text{Ag}_{21}(\text{dppf})_3(\text{SAdm})_{12}/\text{C-300 °C}$.

2.8. X-ray crystallography

The data collection for $\text{Ag}_{22}(\text{dppf})_3(\text{SAdm})_{12}(\text{BPh}_4)_2$ and $\text{Au}_1\text{Ag}_{21}(\text{dppf})_3(\text{SAdm})_{12}(\text{BPh}_4)_2$ was carried out on a Stoe Stadivari diffractometer under a liquid nitrogen flow at 150 K (for $\text{Ag}_{22}(\text{dppf})_3(\text{SAdm})_{12}(\text{BPh}_4)_2$) and 120 K (for $\text{Au}_1\text{Ag}_{21}(\text{dppf})_3(\text{SAdm})_{12}(\text{BPh}_4)_2$), using graphite-monochromatized Cu K α radiation ($\lambda = 1.54186 \text{ \AA}$). Data reductions and absorption corrections were performed using the SAINT and SADABS programs, respectively. The structure was solved by direct methods and refined with full-matrix least squares on F^2 using the SHELXTL software package. All non-hydrogen atoms were refined anisotropically, and all the hydrogen atoms were set in geometrically calculated positions and refined isotropically using a riding model.

2.9. Electrochemical measurements for the oxygen reduction reaction

All electrochemical measurements were performed on the CHI700E electrochemical workstation in a standard three-electrode system at room temperature. A saturated Ag/AgCl electrode and platinum wire were used as the reference electrode and counter electrode, respectively. A rotating disk electrode (RDE, geometric area = 0.196 cm^2) modified with catalyst ink was used as the working electrode in the ORR test. Catalyst ink was prepared by dispersing 10 mg catalyst in 1 mL ethanol with 10 μL Nafion, which was sonicated for 30 min to form a homogeneous solution. All potential values in this work were calibrated to the reversible hydrogen electrode (RHE) scale according to the following equation:

$$E_{\text{RHE}} = E(\text{Ag}/\text{AgCl}) + 0.966.$$

For the oxygen reduction reaction, the working electrode was prepared by dropping 10 μL catalyst ink on the RDE. 0.1 M KOH was used as the electrolyte and was saturated with pure Ar or O_2 . A scan rate of 0.05 V s^{-1} was used for all the experiments.

3. Results and discussion

3.1. Crystallization, structural analysis, and characterization of two nanoclusters

The detailed synthesis methods are provided in the Experimental methods, and the crystals of both $\text{Ag}_{22}(\text{dppf})_3(\text{SAdm})_{12}(\text{BPh}_4)_2$ and $\text{Au}_1\text{Ag}_{21}(\text{dppf})_3(\text{SAdm})_{12}(\text{BPh}_4)_2$ nanoclusters were obtained *via* crystallization in toluene/methanol after three days. SCXC was applied to determine the structures of $[\text{Ag}_{22}(\text{dppf})_3(\text{SAdm})_{12}](\text{BPh}_4)_2$ and $[\text{Au}_1\text{Ag}_{21}(\text{dppf})_3(\text{SAdm})_{12}](\text{BPh}_4)_2$. The overall structures are shown in Fig. S1.† There are two PPh_4 -counterions in the unit cell of Ag_{22} or $\text{Au}_1\text{Ag}_{21}$, indicating the +2-charge state of both nanoclusters.

The free electron counts of both nanoclusters are calculated as $8e$ (*i.e.*, 22 (metal) -12 (thiol ligands) -2 (charge) = $8e$).

The detailed structural analysis is shown in Fig. 1; the icosahedral M_1Ag_{12} ($\text{M} = \text{Au}/\text{Ag}$) kernel is protected by three $\text{Ag}_3(\text{SR})_4$ motifs and three dppf ligands. Taking Ag_{22} as an example, the distances of the central Ag with the other 12 Ag atoms in the icosahedral kernel vary from 2.773 to 2.789 Å , with an average distance of 2.779 Å (Fig. 2A-I), and the average distance of the interlayer twelve Ag atoms is 2.924 Å (Fig. 2A-II). Both sets of three $\text{Ag}_3(\text{SR})_4$ motifs and three dppf ligands are arranged in a trillion-shape mode (Fig. 1A). Detailed bond length comparison of Ag_{22} and $\text{Au}_1\text{Ag}_{21}$ is shown in Fig. 2. Two pairs of enantiomers of these nanoclusters are shown in Fig. S2 and S3.†

The UV-vis spectra of the purified Ag_{22} and $\text{Au}_1\text{Ag}_{21}$ nanoclusters are shown in Fig. 3a and b. Two broad optical absorptions at 406 and 448 nm, along with four shoulder peaks at 480, 515, 565 and 650 nm, are found for Ag_{22} . By comparison, two main peaks at 387 and 430 nm, and two shoulder peaks at 483

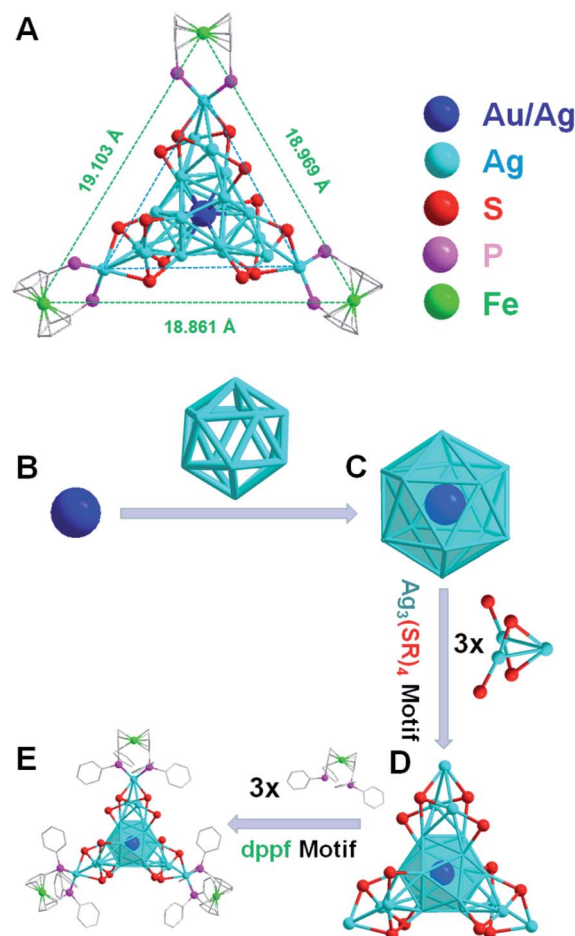


Fig. 1 (A) Total structure of $[\text{M}_1\text{Ag}_{21}(\text{dppf})_3(\text{SAdm})_{12}]^{2+}$. (B) Innermost Au/Ag atom. (C) M_1Ag_{12} kernel with an icosahedral configuration ($\text{M} = \text{Au}/\text{Ag}$). (D) M_1Ag_{12} kernel protected by three $\text{Ag}_3(\text{SR})_4$ motifs. (E) Three dppf architectures occupy the vertex positions of the nanocluster. For clarity, H and some C atoms are omitted. Color labels: blue/light blue = Au/Ag; red = S; purple = P; green = Fe.



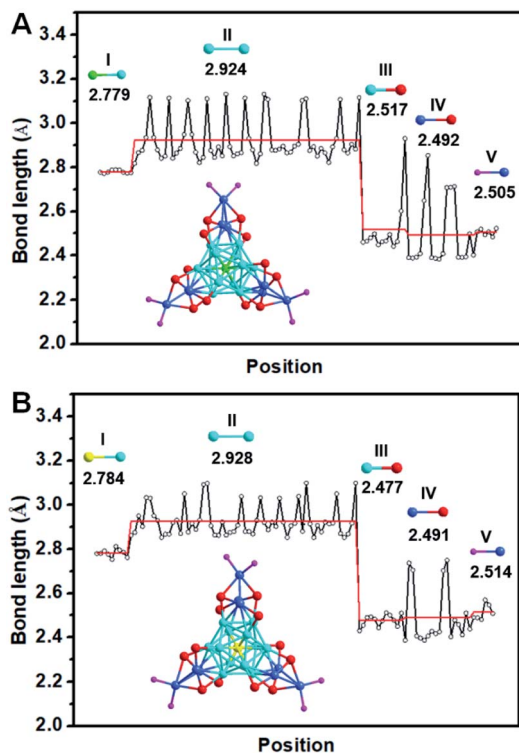


Fig. 2 Bond lengths in (A) Ag_{22} and (B) $\text{Au}_1\text{Ag}_{21}$ nanoclusters. (I) $M_{\text{kernel}}-\text{Ag}_{\text{kernel shell}}$, (II) $\text{Ag}_{\text{kernel shell}}-\text{Ag}_{\text{kernel shell}}$, (III) $\text{Ag}_{\text{kernel shell}}-\text{S}$, (IV) $\text{Ag}_{\text{shell}}-\text{S}$, (V) $\text{P}-\text{Ag}_{\text{shell}}$. $M = \text{Au}/\text{Ag}$. Color labels: yellow = Au; green, blue, light blue = Ag; red = S; purple = P.

and 615 nm are observed for $\text{Au}_1\text{Ag}_{21}$. The experimental optical absorption spectra are also used to calculate the optical energy gaps (Fig. S4 and S5[†]) that are 1.49 eV and 1.69 eV for Ag_{22} and $\text{Au}_1\text{Ag}_{21}$, respectively. The chemical formulas of both nanoclusters are confirmed by high-resolution ESI-TOF-MS, XPS, and TGA. Firstly, ESI-MS spectra confirmed the purity of Ag_{22} and $\text{Au}_1\text{Ag}_{21}$ (Fig. 3c and d). An intense peak at m/z of 3021.72 Da is shown in Fig. 3c, the magnification of this peak evidencing a +2-charge state of Ag_{22} , because this set of peaks shows a characteristic isotopic pattern with peaks separated by an m/z of 0.5 Da. The complete match of experimental and simulated isotope patterns verifies the measured formulas of Ag_{22} . Also, the intense peak of $\text{Au}_1\text{Ag}_{21}$ matches well with its theoretical value (Fig. 3d). TGA was also employed to test the purity of Ag_{22} and $\text{Au}_1\text{Ag}_{21}$ (Fig. 3e and f). The experimental value of 56.65% shows a little difference from the theoretical value of 58.45% for Ag_{22} ; for $\text{Au}_1\text{Ag}_{21}$, the experimental value is determined to be 56.33%, close to its theoretical value of 57.62%. The theoretical value of 58.45 wt% weight loss for $[\text{Ag}_{22}(\text{dppf})_3(\text{SAdm})_{12}]\text{Cl}_2$ was calculated *via* $[\text{M}(12\text{SR} + 2\text{Cl} + 3\text{dppf} - 3\text{Fe})/\text{M}(22\text{Ag} + 12\text{SR} + 2\text{Cl} + 3\text{dppf})] \times 100\%$. The theoretical value of 57.62 wt% weight loss for $[\text{Au}_1\text{Ag}_{21}(\text{dppf})_3(\text{SAdm})_{12}]\text{Cl}_2$ was calculated *via* $[\text{M}(12\text{SR} + 3\text{dppf} + 2\text{Cl} - 3\text{Fe})/\text{M}(\text{Au} + 21\text{Ag} + 12\text{SR} + 3\text{dppf} + 2\text{Cl})] \times 100\%$. XPS was performed to probe the compositions of these nanoclusters (Fig. 3g and h), and all the elements in both nanoclusters have been clearly marked. FT-IR spectroscopy was used to characterize the nanoclusters (Fig. S6[†]). The peaks of 2848

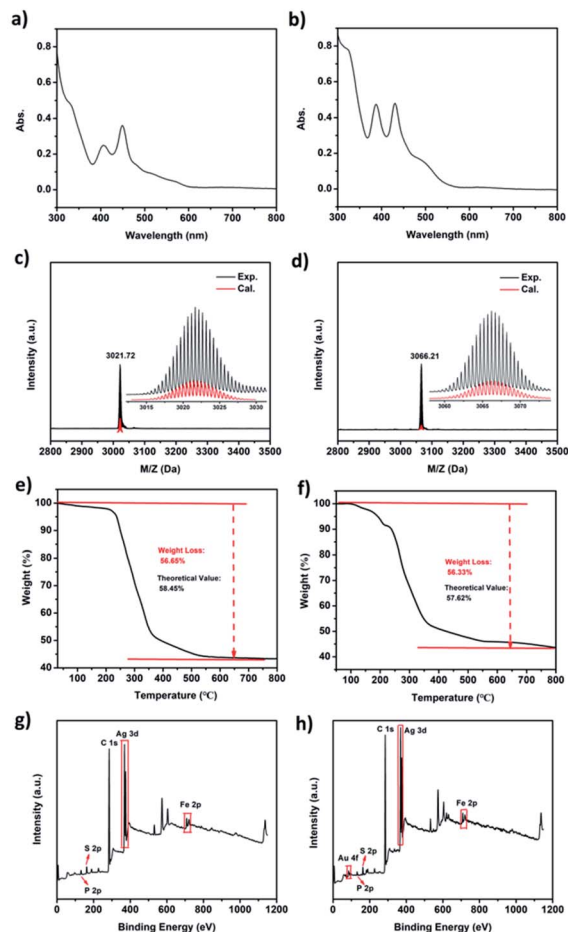


Fig. 3 The (a and b) UV-vis spectra; (c and d) ESI-MS spectra; (e and f) TGA spectra and (g and h) XPS spectra of $\text{Ag}_{22}(\text{dppf})_3(\text{SAdm})_{12}(\text{BPh}_4)_2$ and $\text{Au}_1\text{Ag}_{21}(\text{dppf})_3(\text{SAdm})_{12}(\text{BPh}_4)_2$.

cm^{-1} and 2905 cm^{-1} are assigned to 1-AdmSH, and the 3051 cm^{-1} peak belongs to dppf. DPV measurements were then conducted. For Ag_{22} , two oxidation peaks at 0.869 (O1) and 1.334 V (O2), and three reduction peaks at -1.167 (R1), -1.392 (R2) and -1.770 V (R3) are observed (Fig. S7[†]). For $\text{Au}_1\text{Ag}_{21}$, two oxidation peaks at 0.738 (O1) and 1.253 V (O2), and two reduction peaks at -1.538 (R1) and -1.750 (R2) are detected. Generally, the HOMO-LUMO gap corresponds to the difference between O1 – R1 and O2 – O1.³⁸ In this context, the electrochemical HOMO-LUMO gaps are 1.571 V ($2.036 \text{ V} - 0.465 \text{ V}$) and 1.761 V ($2.276 \text{ V} - 0.515 \text{ V}$) for Ag_{22} and $\text{Au}_1\text{Ag}_{21}$, respectively; they are consistent with the optical energy gaps.

3.2. CV and LSV measurements for the two nanoclusters

The peak potentials (E_p) and peak current density (j_p), as well as the onset potential (E_{onset}) are the criteria to evaluate the ORR activity of a nanomaterial. Cyclic voltammetry (CV) and linear sweep voltammetry (LSV) measurements were then carried out to study the ORR performances of the activated carbon-supported Ag_{22} and $\text{Au}_1\text{Ag}_{21}$ nanoclusters (see the Experimental methods for the preparation of activated carbon-supported nanoclusters).



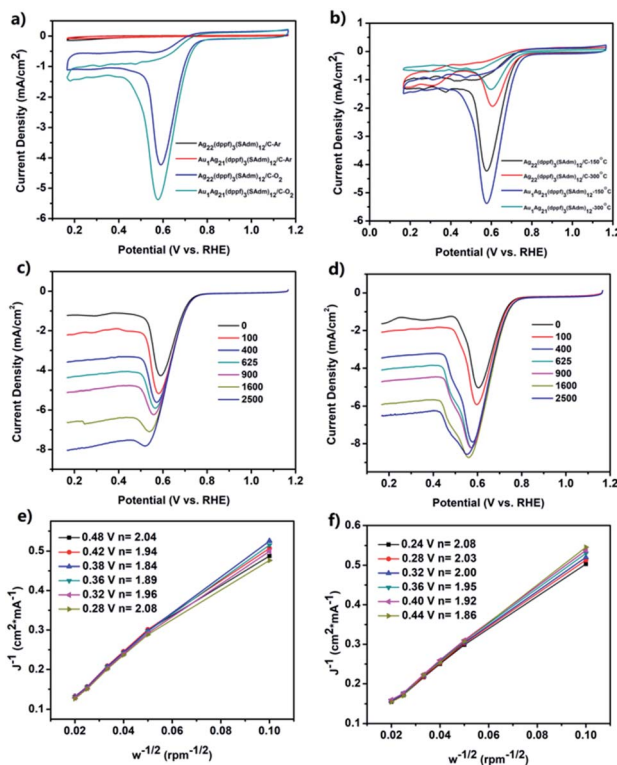


Fig. 4 (a) CV curves of $\text{Ag}_{22}(\text{dppf})_3(\text{SAdm})_{12}/\text{C}$ and $\text{Au}_1\text{Ag}_{21}(\text{dppf})_3(\text{SAdm})_{12}/\text{C}$. (b) CV curves of $\text{Ag}_{22}(\text{dppf})_3(\text{SAdm})_{12}/\text{C}$ and $\text{Au}_1\text{Ag}_{21}(\text{dppf})_3(\text{SAdm})_{12}/\text{C}$ after calcination at 150 and 300 °C. LSV curves of (c) $\text{Ag}_{22}(\text{dppf})_3(\text{SAdm})_{12}/\text{C}$ and (d) $\text{Au}_1\text{Ag}_{21}(\text{dppf})_3(\text{SAdm})_{12}/\text{C}$. Koutecky–Levich plots for the ORR on (e) Ag_{22}/C and (f) $\text{Au}_1\text{Ag}_{21}(\text{dppf})_3(\text{SAdm})_{12}/\text{C}$ at different potentials. All these experiments were performed in 0.1 M KOH saturated with Ar/O_2 , and the scan rate was 0.05 V s⁻¹.

Fig. 4a depicts CV measurements acquired with the catalyst-modified electrodes. In Ar-saturated KOH solution (black and red curves), no redox peak was observed within the potential range of 0.166 to 1.166 V. However, well-defined voltammetric peaks emerged in O_2 -saturated 0.1 M KOH solution, showing the obvious activity of $\text{Ag}_{22}(\text{dppf})_3(\text{SAdm})_{12}/\text{C}$ and $\text{Au}_1\text{Ag}_{21}(\text{dppf})_3(\text{SAdm})_{12}/\text{C}$ in the ORR. The peak potentials (E_p) are about 0.59 and 0.57 V for $\text{Ag}_{22}(\text{dppf})_3(\text{SAdm})_{12}/\text{C}$ and $\text{Au}_1\text{Ag}_{21}(\text{dppf})_3(\text{SAdm})_{12}/\text{C}$, respectively. The peak current density (j_p) of $\text{Ag}_{22}(\text{dppf})_3(\text{SAdm})_{12}/\text{C}$ (4.23 mA cm⁻²) and $\text{Au}_1\text{Ag}_{21}(\text{dppf})_3(\text{SAdm})_{12}/\text{C}$ (5.38 mA cm⁻²) is also observed.

These features indicate the feasibility of exploiting activated carbon-supported nanoclusters as highly efficient ORR catalysts. Owing to the positive synergetic effect between metals, $\text{Au}_1\text{Ag}_{21}(\text{dppf})_3(\text{SAdm})_{12}/\text{C}$ displays a better activity than $\text{Ag}_{22}(\text{dppf})_3(\text{SAdm})_{12}/\text{C}$. The loading amount effect for ORR activity was then investigated. Based on the E_p and j_p values, a general reactivity trend for both nanoclusters is that 10% loading amount results in the best ORR activity (Fig. S9 and Table S1†). Other contributing factors, such as capping ligands, were also probed. The corresponding test results are shown in Fig. 4b, S10–S12 and Table S2.† The nanoclusters were loaded on activated carbon with uniform dispersion, and typical TEM images of $\text{Ag}_{22}(\text{dppf})_3(\text{SAdm})_{12}/\text{C}$ and $\text{Au}_1\text{Ag}_{21}(\text{dppf})_3(\text{SAdm})_{12}/\text{C}$ before

and after calcination at 150 and 300 °C are shown in Fig. S12.† These results clearly indicate that the calcination processes induce the aggregation of nanoclusters and further result in decreasing ORR activity. Interestingly, this finding is opposed to previous reports on $\text{Pd}_5(\text{C}_{12}\text{H}_{25}\text{S})_{13}$.³⁹ The CV curve of commercial Pt/C is shown in Fig. S13.† The comparison of the ORR activity of $\text{M}_1\text{Ag}_{21}(\text{dppf})_3(\text{SAdm})_{12}/\text{C}$ with materials in other reported studies in alkaline solutions is shown in Table S3;† the activities of $\text{M}_1\text{Ag}_{21}(\text{dppf})_3(\text{SAdm})_{12}/\text{C}$ are better than those of most non-Pt ORR electrocatalysts.

In order to gain further insights into the ORR process of the two nanoclusters, linear sweep voltammetry (LSV) measurements were performed at different rotation rates with a scan rate of 0.05 V s⁻¹ in O_2 -saturated 0.1 M KOH solution using a rotating disk electrode (RDE). All the LSV curves for $\text{Ag}_{22}(\text{dppf})_3(\text{SAdm})_{12}/\text{C}$ and $\text{Au}_1\text{Ag}_{21}(\text{dppf})_3(\text{SAdm})_{12}/\text{C}$ are shown in Fig. 4c and d. A general observed trend is that the ORR current density increases with increasing electrode rotation rate. The onset potential (E_{onset}) is found at approximately 0.80 and 0.82 V for $\text{Ag}_{22}(\text{dppf})_3(\text{SAdm})_{12}/\text{C}$ and $\text{Au}_1\text{Ag}_{21}(\text{dppf})_3(\text{SAdm})_{12}/\text{C}$, respectively, close to that obtained in CV measurements (Fig. 4a; 0.79 and 0.80 V). Kinetic information on the ORR for both nanoclusters was obtained according to the Koutecky–Levich (K–L) equations. Fig. 4e and f depict the corresponding K–L plots (j^{-1} vs. $\omega^{-1/2}$) at various electrode potentials. The data exhibit good linearity, and the slopes remain approximately constant over the potential range of 0.28 to 0.48 V in Fig. 4e, thus suggesting consistent electron transfer for oxygen reduction at different electrode potentials for $\text{Ag}_{22}(\text{dppf})_3(\text{SAdm})_{12}/\text{C}$. The linearity of the plots is usually taken as an indication of first-order reaction kinetics with respect to dissolved O_2 . The corresponding numbers (n) of transferred electrons vary from 1.84 to 2.08, indicating that the dominant two-electron ORR process occurs with $\text{Ag}_{22}(\text{dppf})_3(\text{SAdm})_{12}/\text{C}$. This is the same for $\text{Au}_1\text{Ag}_{21}(\text{dppf})_3(\text{SAdm})_{12}/\text{C}$ (Fig. 4f) for which the values of n are 1.86–2.08 at the potential range of 0.24–0.44 V, close to the theoretical value ($n = 2$). Moreover, the stability of these two systems has also been tested (Fig. S14–S17†). For $\text{Ag}_{22}(\text{dppf})_3(\text{SAdm})_{12}/\text{C}$, after 1000 CV cycles, the limiting current density decreased by only 0.13 mA cm⁻², and after another 1000 cycles, the current density decreased by 0.55 mA cm⁻². The stability of $\text{Au}_1\text{Ag}_{21}(\text{dppf})_3(\text{SAdm})_{12}/\text{C}$ is a little poorer than that of $\text{Ag}_{22}(\text{dppf})_3(\text{SAdm})_{12}/\text{C}$, indicating a decrease of 1.17 mA cm⁻² after 2000 cycles compared to the initial value. Meanwhile, the onset potential (E_{onset}) and half-wave potential ($E_{1/2}$) remained unchanged after 1000 or 2000 cycles, which indicates that the electrocatalysts are recyclable and stable.

3.3. Possible mechanism rationalizing the increase of ORR activities

Then, several control experiments were carried out to investigate the origin of the higher catalytic activities of these two systems. CV curves of $\text{Ag}_{25}(2,4\text{-DMBT})_{18}/\text{C}$, $\text{Au}_{25}(\text{PET})_{18}/\text{C}$, $\text{Ag}_{44}(\text{SPhF}_2)_{30}/\text{C}$, and $\text{Ag}_{50}(\text{dppm})_6(\text{TBBM})_{30}/\text{C}$ are shown in Fig. 5a; these nanoclusters possess sizes similar to that of M_1Ag_{21} . Their activities are not as good as that of



Table 1 Comparison of the ORR activities of different samples calculated by CV and LSV measurements in 0.1 M KOH solution

Sample	E_p^a (V)	j_p^a (mA cm $^{-2}$)	E_{onset}^b (V)	$E_{1/2}^b$ (V)	j_p^b (mA cm $^{-2}$) at 0.57 V
Ag ₂₂ (dppf) ₃ (SAdm) ₁₂ /C	0.59	4.23	0.82	0.63	6.75
Au ₁ Ag ₂₁ (dppf) ₃ (SAdm) ₁₂ /C	0.57	5.38	0.86	0.66	8.39
Ag ₂₅ (2,4-DMBT) ₁₈ /C	0.59	2.96	0.79	0.62	4.01
Au ₂₅ (PET) ₁₈ /C	0.56	2.98	0.80	0.63	4.58
Au ₁ Ag ₂₂ (SAdm) ₁₂ /C	0.63	1.21	0.81	0.64	2.75
Ag ₂₉ (PPh ₃) ₄ (BDT) ₁₂ /C	0.63	2.54	0.78	0.67	3.79
Ag ₄₄ (SPhF ₂) ₃₀ /C	0.55	3.06	0.77	0.62	5.37
Ag ₅₀ (dppm) ₆ (TBBM) ₃₀ /C	0.61	3.28	0.81	0.66	5.27
Ag-dppf complexes	0.62	2.83	0.82	0.68	3.15
Ag-dppf nanoparticles	0.58	2.62	0.79	0.62	3.28
Ag-SAdm complexes	0.59	2.65	0.81	0.65	3.52
Ag-SAdm nanoparticles	0.59	2.85	0.78	0.63	4.59
dppf/C	0.61	2.23	0.80	0.66	3.93
Activated carbon (C)	0.66	2.51	0.82	0.70	3.27
Commercial Pt/C	0.70	5.12	0.98	0.81	8.40

^a E_p and j_p were determined by CV (Fig. 4a and 5a-c). ^b E_{onset} , $E_{1/2}$, and j_p at 0.57 V were determined from rotating disk voltammograms with a rotation speed of 2500 rpm (Fig. 4c and d, and ESI Fig. S18-S20).

M₁Ag₂₁(dppf)₃(SAdm)₁₂/C. In addition, the performances of the smaller complexes or larger nanoparticles were also compared (Fig. 5b and Table 1). These experiments exclude the effects of size on the ORR activity of these systems, *i.e.*, **M₁Ag₂₁** nanoclusters showed better performances than various other nanoclusters. In order to probe whether it is a cluster core or a ferrocenyl ligand that is responsible for the redox wave in ORR catalysis, the activities of dppf/C, activated carbon (C), Ag₂₉(PPh₃)₄(BDT)₁₂/C, and Au₁Ag₂₂(SAdm)₁₂/C were further compared. As shown in Fig. 5c and Table 1, the blank support and individual dppf/C had poor activities, and the difference between AdmSH and PPh₃ was negligible in light of the activities of Ag₂₉(PPh₃)₄(BDT)₁₂/C and Au₁Ag₂₂(SAdm)₁₂/C. Moreover, the XPS spectra of Au 4f and Ag 3d in **Au₁Ag₂₁** and **Au₁Ag₂₁(dppf)₃(SAdm)₁₂/C** (Fig. 5d and e), and Ag 3d in **Ag₂₂** and **Ag₂₂(dppf)₃(SAdm)₁₂/C** (Fig. 5f) indicate that there is no charge transfer between the two metals. Notably, the XPS spectra of Ag 3d shift to higher energies upon deposition on C, which is ascribed to ionization of surface Ag (for example, formation of Ag-O oxide) as described in other reports. This shows an increase of electron density (upshift of the d-band center) in Ag, further resulting in improvement of the O₂ reduction activities.⁴⁰⁻⁴² The aforementioned experimental results clearly confirm that dppf does play an important role in improving the ORR performance. As a result, the activated carbon-supported **Ag₂₂** and **Au₁Ag₂₁**, with a loading of 10% and without pyrolysis, emerged as the optimal ORR catalysts, with better performance than other similar-sized nanoclusters without the dppf ligand (*i.e.*, Ag₂₉, Ag₄₄, Ag₅₀, Au₁Ag₂₂, Ag₂₅, and Au₂₅), larger Ag nanoparticles and dppf-containing complexes. Earlier, a large number of theoretical calculations have shown that the origin of the electrocatalytic activity is mainly due to the strain effect, ligand effect, geometric effect or ensemble effect based on various models (the micro-kinetic model, the activity volcano plots, origin of adsorbate scaling relationships).^{43,44} Taken together, in these two systems, not only in Ag with dppf ligands,

but also in the nanocluster **Au₁Ag₂₁**, the ensemble effect is the key factor explaining the excellent ORR performances. The ensemble effect shifts the d-band center of the whole systems downward in energy, leads to weaker binding of intermediates

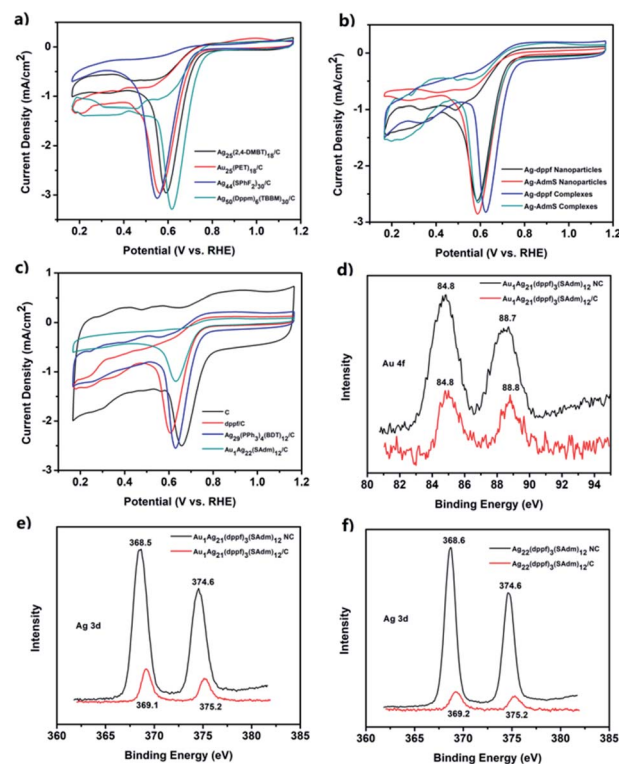


Fig. 5 CV curves of (a) Ag₂₅(2,4-DMBT)₁₈/C, Au₂₅(PET)₁₈/C, Ag₄₄(SPhF₂)₃₀/C and Ag₅₀(dppm)₆(TBBM)₃₀/C; (b) Ag-dppf nanoparticles/C, Ag-dppf complexes/C, Ag-SAdm nanoparticles/C, and Ag-SAdm complexes; (c) activated carbon, dppf/C, Ag₂₉(PPh₃)₄(BDT)₁₂/C, and Au₁Ag₂₂(SAdm)₁₂/C. XPS spectra of (d) Au 4f and (e) Ag 3d of the **Au₁Ag₂₁** nanocluster and **Au₁Ag₂₁(dppf)₃(SAdm)₁₂/C**; (f) Ag 3d of the **Ag₂₂** nanocluster and **Ag₂₂(dppf)₃(SAdm)₁₂/C**.



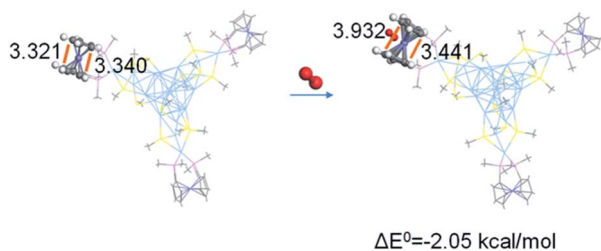


Fig. 6 Corresponding distances in the optimized geometry of $\text{Ag}_{22}(\text{dmpf})_3(\text{SMe})_{12}$ and $\text{Ag}_{22}(\text{dmpf})_3(\text{SMe})_{12}-\text{O}_2$.

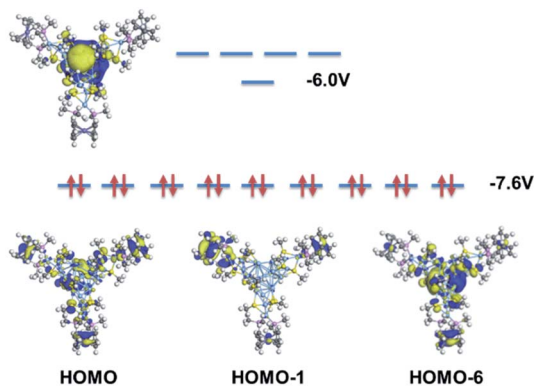


Fig. 7 Frontier orbital analysis of $\text{Ag}_{22}(\text{dmpf})_3(\text{SMe})_{12}$.

and therefore increases the ORR activity. The parallel computational screening of Au(Rh) and Ag alloys as ORR electrocatalysts has been previously performed by the Henkelman group.⁴⁵

Herein, DFT calculations were performed on the interaction of dioxygen and the modelling of Ag_{22} by simplifying bis(diphenylphosphino)ferrocene (dppf) and SAdm as bis(dimethylphosphino)ferrocene (dmpf) and SMe, respectively. The geometry optimization with DFT calculations (at the GGA:PBE/DND level of theory) on the DMol³ suite of programs implies an exothermic ligation of dioxygen to ferrocene by 2.1 kcal mol⁻¹, as expected. Comparison of the optimized geometry of $\text{Ag}_{22}(\text{dmpf})_3(\text{SMe})_{12}$ and $\text{Ag}_{22}(\text{dmpf})_3(\text{SMe})_{12}-\text{O}_2$ indicates opened cyclopentadienyl groups and a short Fe–O distance (3.154 Å, Fig. 6). In addition, the Mulliken charge analysis shows charge transfer from the $\text{Ag}_{22}(\text{dmpf})_3(\text{SMe})_{12}$ cluster to the dioxygen moiety (the Mulliken charge of O_2 in $\text{Ag}_{22}(\text{dmpf})_3(\text{SMe})_{12}-\text{O}_2$ is -0.135), supporting the reduction of dioxygen by the concerned Ag_{22} cluster.

Meanwhile, according to the frontier orbital analysis (Fig. 7), the HOMO of Ag_{22}^{2+} shows 9 degenerate molecular orbitals. Three of them correspond to the supramolecular P orbitals (centered at the metallic core, see HOMO-6 for example), and the other 6 orbitals mainly distributed on the three ferrocene groups (see HOMO and HOMO-1 for example; the full details are provided in Fig. S21†). The extensive orbital contributions of the ferrocene group to the HOMOs support the proposal that the ferrocene-containing ligand enhances the ORR activity.

4. Conclusions

In summary, by introducing the dppf ligand, two novel nanoclusters, $\text{Ag}_{22}(\text{dppf})_3(\text{SAdm})_{12}(\text{BPh}_4)_2$ and $\text{Au}_1\text{Ag}_{21}(\text{dppf})_3(\text{SAdm})_{12}(\text{BPh}_4)_2$, have been controllably synthesized. The atomically precise structures of the two nanoclusters were determined by SCXC and further confirmed by ESI-TOF-MS, TGA, XPS, DPV, and FT-IR measurements. Both nanoclusters supported on activated carbon (C) exhibited excellent ORR activity. DFT calculations have confirmed that the presence of the dppf ligand is the underlying cause of improved ORR activity of these two nanoclusters. This study provides new insights into the highly selective 2e-reduction of dioxygen to hydrogen peroxide in nanocluster-based catalysis, which hopefully provides some reference value for future catalytic and ORR studies utilizing nanoclusters.⁴⁶

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

Financial support by the National Natural Science Foundation of China (21901001, 21631001), CNRS and University of Bordeaux is gratefully acknowledged.

Notes and references

- R. Borup, J. Meyers, B. Pivovar, Y. S. Kim, R. Mukundan, N. Garland, D. Myers, M. Wilson, F. Garzon, D. Wood, P. Zelenay, K. More, K. Stroh, T. Zawodzinski, J. Boncella, J. E. McGrath, M. Inaba, K. Miyatake, M. Hori, K. Ota, Z. Ogumi, S. Miyata, A. Nishikata, Z. Siroma, Y. Uchimoto, K. Yasuda, K. Kimijima and N. Iwashita, *Chem. Rev.*, 2007, **107**, 3904–3951.
- M. K. Debe, *Nature*, 2012, **486**, 43–51.
- C. Wei, R. R. Rao, J. Peng, B. Huang, I. E. L. Stephens, M. Risch, Z. J. Xu and Y. Shao-Horn, *Adv. Mater.*, 2019, **31**, 1806296.
- D. Astruc, *Eur. J. Inorg. Chem.*, 2017, 6–29.
- A. Togni and T. Hayashi, *J. Organomet. Chem.*, 1995, **498**, C27–C28.
- D. G. MacDonald, C. Kübel and J. F. Corrigan, *Inorg. Chem.*, 2011, **50**, 3252–3261.
- D.-S. Park, M. A. Jabbar, H. Park, H. M. Lee, S.-C. Shin and Y.-B. Shim, *Bull. Korean Chem. Soc.*, 2007, **28**, 1996–2002.
- S. G. Kim, D. H. Kim, D. M. Kang, M. A. Jabbar, K. S. Min, Y.-B. Shim and S. C. Shin, *Bull. Korean Chem. Soc.*, 2005, **26**, 1603–1606.
- S. H. Li, X. Liu, W. Hu, M. Chen and Y. Zhu, *J. Phys. Chem. A*, 2020, **124**, 6061–6067.
- X. Zou, S. Jin, S. Wang, M. Zhu and R. Jin, *Nano Futures*, 2018, **2**, 045004.
- Q. Tang, G. Hu, V. Fung and D.-e. Jiang, *Acc. Chem. Res.*, 2018, **51**, 2793–2802.



- 12 M. Cao, R. Pang, Q.-Y. Wang, Z. Han, Z.-Y. Wang, X.-Y. Dong, S.-F. Li, S.-Q. Zang and T. C. W. Mak, *J. Am. Chem. Soc.*, 2019, **141**, 14505–14509.
- 13 X. Kang and M. Zhu, *Chem. Soc. Rev.*, 2019, **48**, 2422–2457.
- 14 B. Nieto-Ortega and T. Bürgi, *Acc. Chem. Res.*, 2018, **51**, 2811–2819.
- 15 K. Kwak and D. Lee, *Acc. Chem. Res.*, 2019, **52**, 12–22.
- 16 X. Du and R. Jin, *ACS Nano*, 2019, **13**, 7383–7387.
- 17 G. Li, H. Abroshan, C. Liu, S. Zhuo, Z. Li, Y. Xie, H. J. Kim, N. L. Rosi and R. Jin, *ACS Nano*, 2016, **10**, 7998–8005.
- 18 X. Wan, J. Wang, Z. Nan and Q.-M. Wang, *Sci. Adv.*, 2017, **3**, e1701823.
- 19 T. Higaki, Y. Li, S. Zhao, Q. Li, S. Li, X.-S. Du, S. Yang, J. Chai and R. Jin, *Angew. Chem., Int. Ed.*, 2019, **58**, 8291–8302.
- 20 X. Kang, L. Xiong, S. Wang, Y. Pei and M. Zhu, *Chem. Commun.*, 2017, **53**, 12564–12567.
- 21 T. Higaki, Q. Li, M. Zhou, S. Zhao, Y. Li, S. Li and R. Jin, *Acc. Chem. Res.*, 2018, **51**, 2764–2773.
- 22 W. Chang, P.-Y. Lee, J.-H. Liao, K. K. Chakrahari, S. Kahlal, Y.-C. Liu, M.-H. Chiang, J.-Y. Saillard and C. W. Liu, *Angew. Chem., Int. Ed.*, 2017, **56**, 10178–10182.
- 23 Y. Chen, C. Zeng, D. R. Kauffman and R. Jin, *Nano Lett.*, 2015, **15**, 3603–3609.
- 24 R. Qin, K. Liu, Q. Wu and N. Zheng, *Chem. Rev.*, 2020, **120**, 11810–11899.
- 25 M. Zhu, C. M. Aikens, F. J. Hollander, G. C. Schatz and R. Jin, *J. Am. Chem. Soc.*, 2008, **130**, 5883–5885.
- 26 M. W. Heaven, A. Dass, P. S. White, K. M. Holt and R. W. Murray, *J. Am. Chem. Soc.*, 2008, **130**, 3754–3755.
- 27 Y. Shichibu, Y. Negishi, T. Watanabe, N. K. Chaki, H. Kawaguchi and T. Tsukuda, *J. Phys. Chem. C*, 2007, **111**, 7845–7847.
- 28 S. Zhao, N. Austin, M. Li, Y. Song, S. D. House, S. Bernhard, J. C. Yang, G. Mpourmpakis and R. Jin, *ACS Catal.*, 2018, **8**, 4996–5001.
- 29 H. Chen, C. Liu, M. Wang, C. Zhang, N. Luo, Y. Wang, H. Aroshan, G. Li and F. Wang, *ACS Catal.*, 2017, **7**, 3632–3638.
- 30 Y. Chen, C. Liu, Q. Tang, C. Zeng, T. Higaki, A. Das, D.-e. Jiang, N. L. Rosi and R. Jin, *J. Am. Chem. Soc.*, 2016, **138**, 1482–1485.
- 31 S. Tian, Y.-Z. Li, M.-B. Li, J. Yuan, J. Yang, Z. Wu and R. Jin, *Nat. Commun.*, 2015, **6**, 8667.
- 32 Z. Li, C. Liu, H. Abroshan, D. R. Kauffman and G. Li, *ACS Catal.*, 2017, **7**, 3368–3374.
- 33 X. Kang, S. Jin, L. Xiong, X. Wei, M. Zhou, C. Qin, Y. Pei, S. Wang and M. Zhu, *Chem. Sci.*, 2020, **11**, 1691–1697.
- 34 Q. Li, T.-Y. Luo, M. G. Taylor, S. Wang, X. Zhu, Y. Song, G. Mpourmpakis, N. L. Rosi and R. Jin, *Sci. Adv.*, 2017, **3**, e1603193.
- 35 Z.-R. Wen, Z.-J. Guan, Y. Zhang, Y.-M. Lin and Q.-M. Wang, *Chem. Commun.*, 2019, **55**, 12992–12995.
- 36 X.-Y. Li, Z. Wang, H. Su, S. Feng, M. Kurmoo, C.-H. Tung, D. Sun and L.-S. Zheng, *Nanoscale*, 2017, **9**, 3601–3608.
- 37 L.-Y. Yao and V. W.-W. Yam, *J. Am. Chem. Soc.*, 2016, **138**, 15736–15742.
- 38 X. Kang, S. Chen, S. Jin, Y. Song, Y. Xu, H. Yu, H. Sheng and M. Zhu, *ChemElectroChem*, 2016, **3**, 1261–1265.
- 39 Z. Zhuang and W. Chen, *J. Colloid Interface Sci.*, 2019, **538**, 699–708.
- 40 L. Wang, Z. Tang, W. Yan, H. Yang, Q. Wang and S. Chen, *ACS Appl. Mater. Interfaces*, 2016, **8**, 20635–20641.
- 41 X. Xie, M. Wei, L. Du, Y. Nie, X. Qi, Y. Shaob and Z. Wei, *J. Mater. Chem. A*, 2017, **5**, 15390–15394.
- 42 L. Sumner, N. A. Sakthivel, H. Schrock, K. Artyushkova, A. Dass and S. Chakraborty, *J. Phys. Chem. C*, 2018, **122**, 24809–24817.
- 43 J. A. Trindell, Z. Duan, G. Henkelman and R. M. Crooks, *Chem. Rev.*, 2020, **120**, 814–850.
- 44 Y. Jiao, Y. Zheng, M. Jaroniec and S. Z. Qiao, *Chem. Soc. Rev.*, 2015, **44**, 2060–2086.
- 45 H. Li, L. Luo, P. Kunal, C. S. Bonifacio, Z. Duan, J. C. Yang, S. M. Humphrey, R. M. Crooks and G. Henkelman, *J. Phys. Chem. C*, 2018, **122**, 2712–2716.
- 46 Y. Du, H. Sheng, D. Astruc and M. Zhu, *Chem. Rev.*, 2020, **120**, 526–622.

