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# Assessing the roles of synthesis method and chemical composition in determining structure–property correlations in alloyed, ultrathin nanowire motifs for the methanol oxidation reaction†

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In the context of developing novel fuel cell catalysts, we have successfully synthesized in high yields not only ultrathin nanowires with compositions of Pt<sub>1</sub>Ru<sub>1</sub> and Pt<sub>3</sub>Ru<sub>1</sub> but also more complex spoke-like dendritic clusters of Pt<sub>1</sub>Ru<sub>1</sub> and Pt<sub>1</sub>Ru<sub>9</sub> in ambient pressure under relatively straightforward, solution-based reaction conditions, mediated by either cetyltrimethylammonium bromide (CTAB) or oleylamine (OAm), respectively. EXAFS analysis allowed us to determine the homogeneity of the as-prepared samples. Based on this analysis, only the Pt<sub>3</sub>Ru<sub>1</sub> sample was found to be relatively homogeneous. All of the other samples yielded results, suggestive of a tendency for the elements to segregate into clusters of 'like' atoms. We have also collected complementary HRTEM EDS mapping data, which support the idea of a segregation of elements consistent with the EXAFS results. We attribute the differences in the observed morphologies and elemental distributions within as-prepared samples to the presence of varying surfactants and heating environments, employed in these reactions. Methanol oxidation reaction (MOR) measurements indicated a correlation of specific activity (SA) values not only with intrinsic chemical composition and degree of alloying but also with the reaction process used to generate the nanoscale motifs in the first place. Specifically, the observed performance of samples tested decreased as a function of chemical composition (surfactant used in their synthesis), as follows: Pt<sub>3</sub>Ru<sub>1</sub> (CTAB) > Pt<sub>1</sub>Ru<sub>1</sub> (CTAB) > Pt<sub>1</sub>Ru<sub>1</sub> (OAm) > Pt<sub>1</sub>Ru<sub>9</sub> (OAm).

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## 1. Introduction

State-of-the-art direct alcohol fuel cell (DAFC) electrocatalysts primarily consist of nanostructured precious metals (*i.e.*, Pt) and metal-based alloys supported onto carbon supports; these metals tend to be both scarce and expensive. Moreover, typical Pt-based electrocatalysts lack the stability and durability

required for long-term FC applications, due to a mixture of factors, including CO poisoning, metal dissolution, and surface oxidation.<sup>1,2</sup> Hence, approaches towards creating a novel, highly stable, and well-performing catalyst have relied on either maximizing Pt loading, minimizing overall Pt content, or replacing platinum-group metals (PGMs) with more earth-abundant and less expensive alternatives.<sup>3</sup>

One such strategy involves the creation of ultrathin nanowires (NWs), measuring ~2 nm in average diameter, which are expected to maintain slightly contracted surfaces, the latter of which can weaken the interaction with O<sub>2</sub> and prevent their passivation by O<sub>2</sub>. Specifically, our group has previously noted that ultrathin NWs may be not only more chemically uniform but also more structurally monodisperse with fewer defect sites as compared with commercial bulk and nanoparticulate (NP) analogues.<sup>4</sup> In particular, ultrathin Pt NWs maximize the available surface area-to-volume ratio and minimize overall Pt loading, thereby yielding catalytically attractive entities.<sup>5</sup>

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Another parallel strategy involves the tuning of the chemical composition of Pt alone *via* alloy formation.<sup>6,7</sup> Specifically, the precise nature of the metal can modify the d band structure and electron density of Pt through a “ligand effect”. Indeed, both Pt–Pt bond distances and the intrinsic electronic properties of Pt itself are altered by the presence of the underlying, adjoining metal, all of which can manifest themselves as a “lattice strain effect”. These critically relevant ligand and lattice strain effects can reveal themselves as perceptible shifts in the peak positions within the Pt 4f region associated with chemically sensitive X-ray photoelectron spectroscopy and X-ray absorption fine structure (XAFS) spectra. As an example, within PtRu, Pt atoms are expected to present more vacant 5d electronic states as a consequence of the electron withdrawing effect of the adjoining Ru.<sup>8</sup>

As a matter of combining the desirable approaches of ultrathin NW motifs with alloy generation, we focus on the synthesis of ultrathin, alloyed NWs of Pt with Ru, so as to improve their overall CO tolerance.<sup>9–16</sup> Specifically, when alloyed at low-to-moderate Ru concentrations ( $\chi_{\text{Ru}} = 0\text{--}0.6$ ), the PtRu alloy forms a solid solution with Ru atoms occupying sites within the Pt's face centered cubic lattice.<sup>17</sup> Surface Ru sites can catalyze the oxidation of CO and other C-containing intermediates to CO<sub>2</sub> at lower potentials, thereby decreasing the amount of irreversible binding of CO to active Pt sites and promoting stability.<sup>18</sup> However, contemporary bimetallic PtRu alloys lack sufficient CO tolerance to be effective, long-lasting catalysts, thereby prompting a strong motivation to probe and understand correlations between the fabrication method and the resulting atomic structure.

Ultrathin PtRu NWs have been previously synthesized using hydrothermal methods.<sup>19,20</sup> These reactions resulted in the production of 2–3 nm diameter wires, incorporating different Pt and Ru ratios. With these protocols, surfactants and precursors were dissolved in water, mixed, and subsequently heated within a Teflon-lined steel autoclave. Although hydrothermal methods are certainly reliable, these relatively high-energy-consuming procedures often involve elevated temperatures (upwards of 150–200 °C) and relatively long reaction times (*i.e.*, 24 h or more) with autoclaves run at high pressures.

To mitigate for these potential concerns, herein, we focus on arguably milder solution-based methods to generate our PtRu alloys. As an initial example, in a typical room-temperature, surfactant-mediated synthesis scheme, inspired by previous work by our group and others,<sup>12,21–24</sup> the precursors (*e.g.* H<sub>2</sub>PtCl<sub>6</sub> & RuCl<sub>3</sub>) are co-reduced within the spatial confines of worm-like micellar pores of cetyltrimethylammonium bromide (CTAB) formed in a water-chloroform microemulsion by the addition of NaBH<sub>4</sub>. The chemical composition of as-prepared NWs can be reliably and predictably varied (Pt<sub>1-x</sub>Ru<sub>x</sub>, 0.3 ≥ *x* ≥ 0) by changing the ratio of Pt and Ru content in the added precursor solution. Using this soft template approach, we have successfully reported on the use of the CTAB method to produce a multitude of stable ultrathin NWs, characterized by varying compositions including but not limited to Pt, Pd, Pd<sub>1-x</sub>Ni<sub>x</sub>, Pd<sub>1-x</sub>Cu<sub>x</sub>, Pd<sub>1-x</sub>Au<sub>x</sub>, Pt<sub>1-x</sub>Sn<sub>x</sub>, Pt<sub>1-x</sub>Fe<sub>x</sub>, Pt<sub>1-x-y</sub>Ru<sub>x</sub>Fe<sub>y</sub>, and Pd<sub>1-x</sub>Pt<sub>x</sub>.<sup>25–38</sup>

As a second example, representing a means of comparison with the CTAB method, we have developed a ‘new’ procedure that we are seeking to extend to fabricating RuM (M = arbitrary composition including non-PGMs) NWs. In this context, we have previously used the OAm method to produce ultrathin NW motifs of RuCo alloys.<sup>39</sup> In a typical protocol for Ru NWs, RuCl<sub>3</sub> was dissolved in oleylamine (OAm) and oleic acid (OA), and heated at 350 °C under an air-sensitive environment (*i.e.*, argon) for 1 h. It is worth noting that oleylamine acted as both a reducing agent and a surfactant, whereas oleic acid behaved as a supplementary surfactant agent. In terms of a specific chemical role in the reaction, it has been previously proposed that both oleylamine and oleic acid acting synergistically in concert can behave as capping ligands to effectively guide and enable the formation of NWs through a process of oriented attachment.<sup>40–43</sup>

While these two solution-based synthesis protocols are known to be effective at producing ultrathin NWs, what is far less clear is a general understanding of how the structure and chemical composition of as-prepared anisotropic NW products are necessarily contingent upon and vary depending on precisely how they were made. We will therefore probe and address these questions with not only (i) PtRu NWs, synthesized by CTAB *versus* OAm/OA methods, but also as an added bonus, (ii) within PtRu NWs of varying compositions generated by the CTAB technique. Our analyses will be based on a combination of various characterization techniques, including transmission electron microscopy (TEM), high-resolution TEM (HRTEM), energy dispersive X-ray spectroscopy (EDS), and X-ray absorption fine structure (XAFS). As a subset of XAFS, the extended (EXAFS) signal incorporates crucial information about coordination numbers, interatomic distances, and the nature of disorder within systems (due to both static and dynamic displacements of all atoms from their average positions).<sup>44</sup> Not surprisingly, it has been extensively used for studying local structure and composition in bimetallic nanoalloys.<sup>29,45–47</sup> In particular, the study of the EXAFS spectra can yield valuable data about the number, type of, and distance to the atoms surrounding the central, X-ray absorbing atom.

In terms of prior EXAFS work, especially with respect to analogous NP systems, the structural evolution and atomic distribution of the “microemulsion lyotropic liquid-crystalline”-templated synthesis of mesoporous PtRu NPs after electroreduction were probed for varied duration times, using X-ray absorption near edge structure (XANES) and EXAFS spectroscopy, complemented by EDS and field-emission TEM.<sup>48</sup> Moreover, we have been involved with a comprehensive structural and architectural evaluation based in part on EXAFS and XANES analysis,<sup>49</sup> which revealed that 4.4 nm PtRu (1:1) alloyed NPs actually consisted of crystalline homogeneous random alloys with little twinning in a typical face-centered cubic (fcc) cell; specifically, the Pt atoms were predominantly metallic, whereas the Ru atoms were partially oxidized and were presumably located on the NP surface.

In the context of actual PtRu anodes used for operational direct methanol fuel cells (DMFCs),<sup>50</sup> the catalyst was found to



be essentially metallic with half of the Ru incorporated into a fcc Pt alloy lattice and the remaining half in an amorphous phase. EXAFS analysis suggested that the fcc lattice was not fully disordered and that the phases in which the Ru resided within the catalysts were dissimilar to what would have been expected in conventional oxides. Furthermore, in a separate report,<sup>51</sup> catalyst restructuring was noted during *in situ* DMFC experiments through the observation of increases in the total metal coordination numbers; specifically, EXAFS and XANES showed that the highly oxidized Pt of an as-received commercial Johnson-Matthey PtRu (1 : 1) catalyst is fully reduced to its metallic state, concomitant with the reduction of a substantial amount of oxidized Ru, upon exposure to an actual operating DMFC environment. Finally, in a third study, XANES and EXAFS data confirmed that a highly alloyed state of PtRu NPs is responsible for their superior electrocatalytic performance as compared with typical commercial electrocatalysts,<sup>52</sup> thereby collectively emphasizing the importance of properly assessing chemical composition at the nanoscale.

As we observed at the time, while the quantitative EXAFS models themselves tend to be mutually consistent in their overall conclusions, they still needed to be improved upon and optimized to acquire a thorough and accurate understanding of local atomic structure within relatively complex systems, such as ultrathin PtRu NW alloys. Hence, the novelty of our current study involves not only (i) demonstrable advances in effective solution-based PtRu NW synthesis using our in-house OAm/OA protocol but also more significantly, (ii) the application of EXAFS and HRTEM EDS in order to compare and contrast the discrete effects of (a) synthesis method and (b) variable chemical composition upon the resulting structure and growth of these alloyed anisotropic PtRu motifs.

As an additional novelty, what we have done herein is to correlate the MOR performance of our novel nanoscale motifs not only with carefully tailored chemical compositions but also ultimately with the synthesis reaction process and associated surfactant used to generate these nanomaterials. This was accomplished primarily through a careful analysis of acquired EXAFS and HRTEM data, which suggested not only that the greater Pt content, the better the catalyst performed but also that even with an identical  $Pt_xRu_y$  chemical composition, variable size and morphology do matter significantly for stability under MOR conditions.

## 2. Experimental

### 2.1. Materials

All chemicals were used without further purification. Specifically, dihydrogen hexachloroplatinate(IV) hexahydrate ( $H_2PtCl_6$ , 99.9%), ruthenium chloride ( $RuCl_3$ , 99.9%), oleic acid (OA, 90%), and anhydrous ethanol denatured (99%) were purchased from Beantown Chemical. Hexadecyltrimethylammonium bromide (CTAB, 98%) and oleylamine (OAm, 70%) were obtained through Millipore Sigma. Chloroform (ACS grade) was acquired from VWR. Sodium borohydride ( $NaBH_4$ , 98%) was bought

from Alfa Aesar. Perchloric acid (Optima grade) and methanol (Optima grade) were separately purchased from Fisher Scientific.

### 2.2. Synthesis of nanowires

**2.2.1. Soft-template, CTAB-based method.** A total of 10 mL of aqueous solutions of  $H_2PtCl_6$  and  $RuCl_3$  with concentrations of 20 mM were combined in stoichiometric quantities. This solution was subsequently added to 5 mL of chloroform, containing 40 mM CTAB and 40 mL of water. The mixture was then stirred for 30 minutes, after which 0.2 g of  $NaBH_4$  in 5 mL of water was added with more stirring for another 20 minutes. The product was ultimately collected by centrifugation and washed with ethanol.

**2.2.2. Oleylamine/oleic acid method.** In a typical protocol for Ru NWs, 0.25 mmol of  $RuCl_3$  was dissolved in 7.5 mL of oleylamine and 7.5 mmol of oleic acid, and heated at 350 °C under argon for 1 h. The reaction was performed at atmospheric pressure. The solution was then allowed to cool to room temperature, after which about 20 mL of methanol was later added. The product was collected after centrifugation and subsequently washed several times with methanol, hexane, and ethanol, respectively.

As an extension of that prior work in our lab for creating sustainable solution-based alternatives for nanoscale synthesis, we employed the following modification. Specifically, for the ultrathin NW synthesis of  $Pt_1Ru_1$ , 0.125 mmol  $H_2PtCl_6$  and 0.125 mmol  $RuCl_3$  were dispersed in 7.5 mL of oleic acid and 7.5 mL of oleylamine. The contents of the flask were degassed under vacuum at 120 °C. The sample was then heated at 300 °C under the flow of Ar gas for 1 h, while stirring. The precipitate was ultimately collected by centrifugation, and washed with hexane and ethanol. For the analogous synthesis of ultrathin NWs of  $Pt_1Ru_9$ , 0.025 mmol of  $H_2PtCl_6$  and 0.225 mmol of  $RuCl_3$  were used.

### 2.3. Electrochemical measurements

As-prepared samples of PtRu nanowires and dendritic clusters were supported onto Vulcan carbon XC-72R and rendered into catalyst inks by dispersing these dry powders into 200 proof ethanol to create a 1.5 mg mL<sup>-1</sup> solution. A glassy carbon rotating disk electrode (GC-RDE, Pine Instruments, 5 mm) was prepared by polishing it using aluminum oxide powder (average particle size of 0.3 μm). Four 5 μL drops of the catalyst ink were then loaded onto the glassy carbon electrode, prior to allowing aliquots of this solution to dry under vacuum. One 5 μL drop of an ethanolic 0.025% Nafion solution was subsequently utilized to seal in the catalyst.

A standard three-electrode electrochemical cell was assembled with a Pt counter electrode and an Ag/AgCl<sub>3</sub> reference electrode. Electrochemical measurements of PtRu/C were performed in the presence of a 0.1 M perchloric acid (Optima grade) electrolyte, which had been generated from high-purity type 1 water with a measured resistance of 18.2 MΩ cm. Open circuit potentiometry (OCP) was carried out using pure hydrogen gas to assess the potential of the reference electrode. Cyclic



voltammogram (CV) curves were collected within an Ar-saturated electrolyte solution, at a scan rate of 20 mV s<sup>-1</sup>. The measured catalytic activity toward the oxidation of small organic molecules was determined by acquiring linear sweep voltammograms at a scan rate of 20 mV s<sup>-1</sup> within a solution mixture, containing de-oxygenated 0.1 M perchloric acid and 0.5 M of methanol. Chronoamperometry measurements indicative of stability were obtained with the same methanol/electrolyte solution by ramping up the voltage from 0.0 V to 0.7 V vs. RHE and then monitoring the current for a period of 3600 s.

## 2.4. Characterization

**2.4.1. X-ray absorption fine structure (XAFS) measurements and data analysis.** All XAFS experiments were performed at the QAS (7-BM) beamline at the National Synchrotron Light Source II (NSLS-II) located in Brookhaven National Laboratory (BNL). All data were collected using the transmission mode. A double-crystal Si(111) monochromator was utilized to collect measurements at the Pt L<sub>3</sub>-edge (11564 eV) and Ru K-edge (22117 eV). Reference spectra for the corresponding metal foils were taken during each measurement to be used for energy calibration and data alignment. Data were processed and analyzed using the Athena and Artemis software packages.<sup>53</sup> The Athena software was needed to assign the photoelectron energy origin,  $E_0$ , and to perform edge-step normalization and background subtraction of the measured X-ray absorption coefficient data. The background-subtracted and edge step-normalized absorption coefficient data were then converted to  $k$  space. The  $k^2$ -weighted data were Fourier transformed (FT) to  $r$ -space, and EXAFS fitting was performed in  $r$ -space using Artemis.

Data analysis was first completed on the EXAFS data of elemental metal foils, wherein the coordination number ( $N$ ) of the first shell was set to be equal to 12. The passive electron reduction factors ( $S_0^2$ ) were varied in the fit. For the bimetallic systems, multiple-edge analysis was done to fit the signals, measured from each of the alloying constituent component's absorption edge, simultaneously. Data for the monometallic samples were simulated using FEFF calculations, performed using fcc structures for Pt and the hcp structure for Ru. To calculate FEFF theory for the heterometallic samples incorporating elements A and B, the atoms of type B were put into a first nearest neighbor position within the coordinate list with respect to the atoms of the type A. The  $S_0^2$  parameters for the NW components were subsequently fixed to be equal to those obtained for their respective bulk foil counterparts.

For the bimetallic NWs, the fits were performed for both edges concurrently, and the following constraints were applied.<sup>54</sup> In particular, the heterometallic bond lengths were set to be equal ( $R_{A-B} = R_{B-A}$ ) along with the mean squared bond length disorders ( $\sigma_{A-B}^2 = \sigma_{B-A}^2$ ), whereas the homometallic bond lengths (*i.e.*,  $R_{A-A}$  and  $R_{B-B}$ ) and mean squared bond length disorders (*i.e.*,  $\sigma_{A-A}^2$  and  $\sigma_{B-B}^2$ ) were varied independently. The coordination numbers were also tuned independently for all samples.

**2.4.2. Transmission electron microscopy (TEM).** Transmission electron microscopy (TEM) images were collected with the JEOL 1400 transmission electron microscope, operating at an accelerating voltage of 120 kV. The instrument was also equipped with a 2048 × 2048 Gatan CCD camera. As-prepared nanowire samples were prepared for TEM imaging by dispersing in ethanol *via* sonication. Small aliquots of the solutions were then drop cast onto lacey carbon-coated copper grids (1 or 2 drops). TEM images were quantitatively analyzed with the use of the ImageJ software to record measurements of average size and standard deviation.

**2.4.3. High resolution transmission electron microscopy (HRTEM).** To gain complementary chemical insights, high angle annular dark field (HAADF) images and HRTEM EDS mapping were obtained on an FEI Talos F200X instrument, working at 200 kV and equipped with a four-quadrant 0.9-sr energy dispersive X-ray spectrometer (EDS) for both elemental and compositional mapping.

**2.4.4. Thermogravimetric analysis (TGA).** The actual loading amounts of PtRu samples deposited onto Vulcan XC-72R carbon were confirmed by TGA using a TA Instruments Q500. Powders were dried and then loaded into a clean 50 μL Pt pan. These were subsequently heated to 600 °C at a rate of 10°/min and held at this temperature for 10 minutes. This relatively high temperature was chosen to ensure the complete degradation of any organic components, including the Vulcan carbon supports. Compressed air and anhydrous nitrogen were used as carrier gases with a flow rate of 40 mL min<sup>-1</sup> and 120 mL min<sup>-1</sup>, respectively, throughout these measurements.

## 3. Results

### 3.1. Structural and chemical overview

TEM images and the corresponding size distributions associated with all samples of not only Pt<sub>1</sub>Ru<sub>1</sub> and Pt<sub>3</sub>Ru<sub>1</sub> prepared using the CTAB method but also Pt<sub>1</sub>Ru<sub>1</sub> and Pt<sub>1</sub>Ru<sub>9</sub>, generated with the OAm/OA protocol are shown in Fig. 1. Subsequent HRTEM images and lattice spacing for all of the four samples produced have been provided in ESI,† Fig. S1. It is evident that synthesis in the presence of CTAB resulted in a web-like, entangled framework of smaller constituent wire-like subunits, whereas the samples obtained with the OAm/OA fabrication protocol incorporated a more globular mass consisting of localized spoke-like, dendritic clusters. As such, even qualitatively speaking, the isolated NW samples appear to highlight the importance and relevance of the synthesis method in dictating the resulting morphology.

The corresponding EDS elemental maps of the CTAB and OAm/OA-derived samples are provided in Fig. 2 and 3, respectively. They highlight that in all of the samples, the ensembles of the atoms appear to be strongly disordered, forming nanoporous-like features, with the dimensions of the smallest units ranging from ~1 to 5 nm (Fig. 1). Interestingly, for both of the Pt:Ru 1:1 samples, produced from the CTAB and OAm/OA-derived protocols, as shown in Fig. 2A and 3A, respectively,



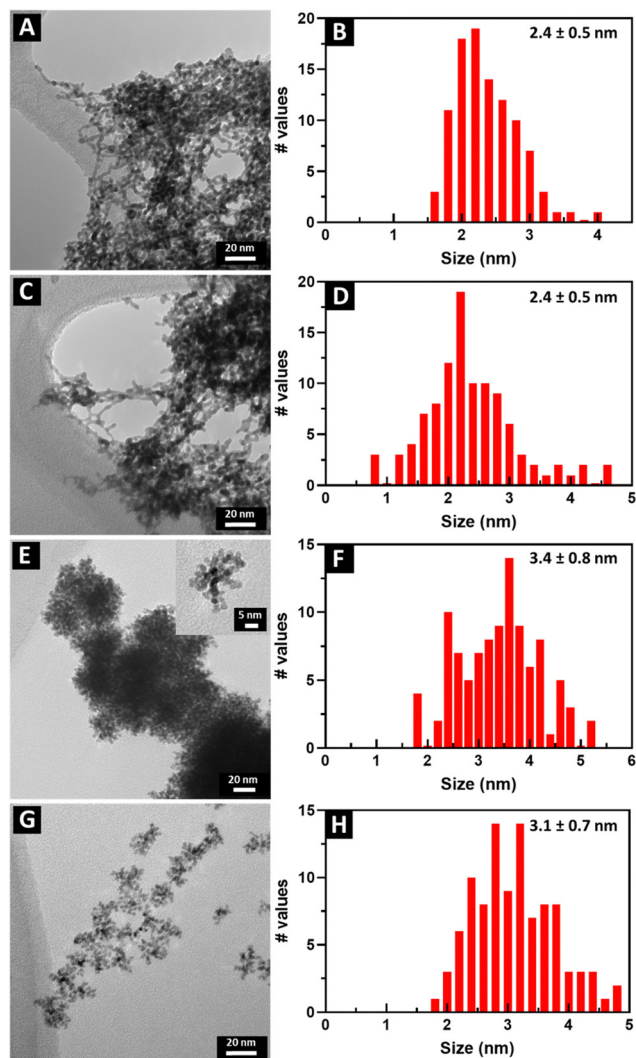


Fig. 1 TEM images and size distributions are shown of a range of chemically distinctive ultrathin NWs, prepared by different synthesis methods. (A) and (B)  $\text{Pt}_1\text{Ru}_1$  and (C) and (D)  $\text{Pt}_3\text{Ru}_1$  were both prepared with CTAB. (E) and (F)  $\text{Pt}_1\text{Ru}_1$  with the inset showing an individual cluster as well as (G) and (H)  $\text{Pt}_1\text{Ru}_9$  were generated with OAm and OA.

the elemental maps seem to show a greater propensity of Pt regions to concentrate and localize within the core, whereas the Ru regions give the impression of being more diffuse with a higher concentration at the outer periphery of the structure. In broad strokes, these observations are generally corroborated by the analysis of local compositional motifs *via* the EXAFS data modeling, as discussed in greater detail below.

### 3.2. Visual examination of the raw XAFS data

Pt  $L_3$ -edge XANES data for  $\text{Pt}_1\text{Ru}_1$  and  $\text{Pt}_3\text{Ru}_1$  prepared using a CTAB surfactant are shown in Fig. 4A. The  $k^2$ -weighted EXAFS data in  $k$ -space and  $r$ -space in which the Fourier transforms were performed for the  $k^2$ -weighted EXAFS spectra in the  $k$ -range from 2 to 15  $\text{\AA}^{-1}$  using the Hanning window function and  $dk = 2 \text{\AA}^{-1}$ , are shown in Fig. 4B and C, respectively. The associated Ru K-edge data for the same samples are highlighted

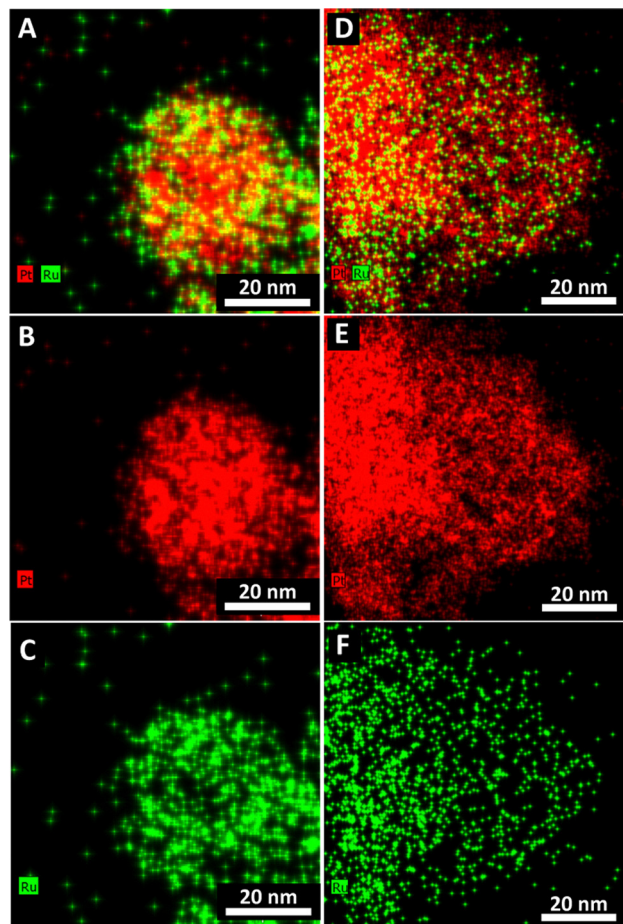


Fig. 2 EDS elemental maps of PtRu generated using the CTAB method.  $\text{Pt}_1\text{Ru}_1$  maps are provided as follows: (A) the composite of Pt and Ru, (B) Pt, and (C) Ru. Analogous  $\text{Pt}_3\text{Ru}_1$  maps are shown, as (D) the composite of Pt and Ru, (E) Pt, and (F) Ru.

in Fig. 5A–C. The Fourier transforms (Fig. 5C) were performed for the  $k^2$ -weighted EXAFS spectra (Fig. 5B) in the  $k$ -range from 2 to 15  $\text{\AA}^{-1}$ . The corresponding Pt  $L_3$  edge SXANES and EXAFS data for  $\text{Pt}_1\text{Ru}_1$  and  $\text{Pt}_1\text{Ru}_9$  prepared using the OAm/OA surfactant are illustrated in ESI<sup>†</sup> Fig. S2. The associated Ru K-edge data are provided in ESI<sup>†</sup> Fig. S3.

### 3.3. Results of EXAFS data analysis

Table S1 (ESI<sup>†</sup>) presents the Fourier transform parameters, fitting ranges, constraints used in the multiple – edge fits, and the resultant  $r$ -factors for each analyzed data set. Fourier transform magnitudes of the  $k^2$ -weighted EXAFS data and fits for the  $\text{Pt}_3\text{Ru}_1$  (originating from CTAB) samples at the Pt  $L_3$ -edge and Ru K-edge are given in Fig. 6A and B, respectively. The corresponding data and the fits for not only the reference Pt and Ru foils but also samples of  $\text{Pt}_1\text{Ru}_1$  (derived from CTAB) in addition to  $\text{Pt}_1\text{Ru}_1$  and  $\text{Pt}_1\text{Ru}_9$  (produced from OAm/OA) are shown in ESI<sup>†</sup> Fig. S4A, B, S5, S6 and S7, respectively.

The best fit values of the amplitude factor ( $S_0^2$ ) for Pt and Ru edges were obtained from the analysis of their respective metal



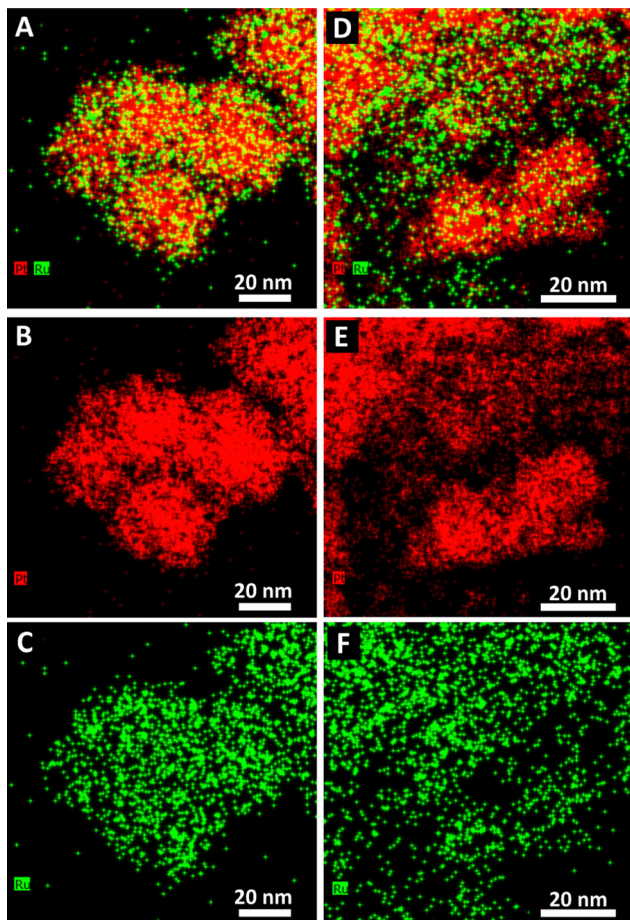


Fig. 3 EDS elemental maps of PtRu prepared using the OAm/OA method. Pt<sub>1</sub>Ru<sub>1</sub> maps are given as (A) the composite of Pt and Ru, (B) Pt, and (C) Ru. Corresponding Pt<sub>1</sub>Ru<sub>9</sub> elemental maps are shown in (D) the composite of Pt and Ru, (E) Pt, and (F) Ru.

foils: 0.84(3) and 0.79(7), respectively. The best fit results of the Pt edge and Ru edge data analyses, together with the results obtained for Pt and Ru foil references, are summarized in ESI,<sup>†</sup> Table S2.

## 4. Discussion

### 4.1. EXAFS

To assess the local compositional motifs of a given bimetallic alloy, Cowley's short range order parameter,  $\alpha$ , is often used. It is expressed as:<sup>44,47,54,55</sup>

$$\alpha_{AB} = 1 - \frac{N_{AB}/N_A}{x_B} \quad (1)$$

where  $N_{AB}$  represents the first nearest neighboring A–B coordination number;  $N_A$  is the total coordination number for A-metal neighbors; and  $x_B$  denotes the concentration of the atomic species B. Because the determination of the concentration of atomic species may differ from those used during the sample preparation, we prefer to rely on a self-consistent method of calculating  $x_A$  and  $x_B$ , which is based on the heterometallic

coordination numbers, namely  $N_{AB}$  and  $N_{BA}$ , independently obtained by EXAFS analysis (ESI,<sup>†</sup> Table S2):<sup>47</sup>

$$\frac{x_B}{x_A} = \frac{N_{AB}}{N_{BA}}; x_B = \frac{1}{1 + N_{BA}/N_{AB}}; x_A = 1 - x_B. \quad (2)$$

The calculated values of the  $N_{Pt}$ ,  $N_{Ru}$ , the alloy concentrations of  $x_A$  and  $x_B$ , obtained using eqn (2), and the short-range order (SRO) parameters obtained using eqn (1), are provided in Table 1.

In our previous work,<sup>56</sup> we have highlighted the limits of applicability of the SRO and include a more detailed discussion in the ESI.<sup>†</sup> That is, its interpretation as an indicator of either positive (when  $\alpha_{AB} > 0$ ) or negative (when  $\alpha_{AB} < 0$ ) tendency to clustering of 'like' (A–A and B–B) atoms is fundamentally limited to the analysis of relatively homogeneous alloys only, *i.e.*, those bimetallic systems in which the relative number of neighbors of different atom types to each atom is approximately the same, regardless of the atom's location. Using that definition, even in a bimetallic nanoparticle (in which the number of nearest neighbors to atoms on the surface is smaller than in the interior), different components can be homogeneously mixed. That characteristic of homogeneity is convenient, because it can be seamlessly verified in terms of observable EXAFS results.

For example, if the total coordination numbers,  $N_A = N_{AA} + N_{AB}$  and  $N_B = N_{BA} + N_{BB}$  are relatively similar, the alloy is homogeneous. Otherwise, there is a macroscopic segregation of components A and B (*e.g.*, cluster-by-cluster or core-shell *etc.*). Notably, in a core-shell situation, the comparison between the values of  $N_A$  and  $N_B$  will communicate which element is more under-coordinated and, hence, preferentially localized on the surface. Only when the alloy is found to be relatively homogeneous can we ponder the next question: What is the short-range order in the alloy?

Therefore, prior to interpreting the SRO parameters, it is important to first investigate the relationship between the total coordination numbers  $N_{Pt}$  and  $N_{Ru}$  of the two different elements. Table 1 shows that their values are significantly different for both Pt<sub>1</sub>Ru<sub>1</sub> samples. Namely, they are, first, larger in the CTAB sample as compared with the OAm-derived one, and, second,  $N_{Pt} > N_{Ru}$  in both types of samples. It should also be noted that as the composition of the nanowires changes using both the CTAB and OAm-based methods, the  $N_{Pt}$  remains constant, but conversely, the  $N_{Ru}$  does change. For example, with the CTAB protocol, as the relative amount of Ru increases from Pt<sub>3</sub>Ru<sub>1</sub> to Pt<sub>1</sub>Ru<sub>1</sub>, the corresponding  $N_{Ru}$  value decreases from 7.1 to 4.3. Likewise, as the chemical composition of the sample created using the OAm method evolves from Pt<sub>1</sub>Ru<sub>1</sub> to Pt<sub>1</sub>Ru<sub>9</sub>, we note that the coordination numbers associated with Ru decrease from 8.1 to 4.6. This observation indicates both a reduction in the metal–metal coordination and a decrease in the homogeneity of the mixture.

These results are in full agreement with the TEM size distributions and the EDS elemental maps (Fig. 1–3) we have previously discussed. In particular, Fig. 1B and F show that the characteristic size of the metal regions in the Pt<sub>1</sub>Ru<sub>1</sub> sample is



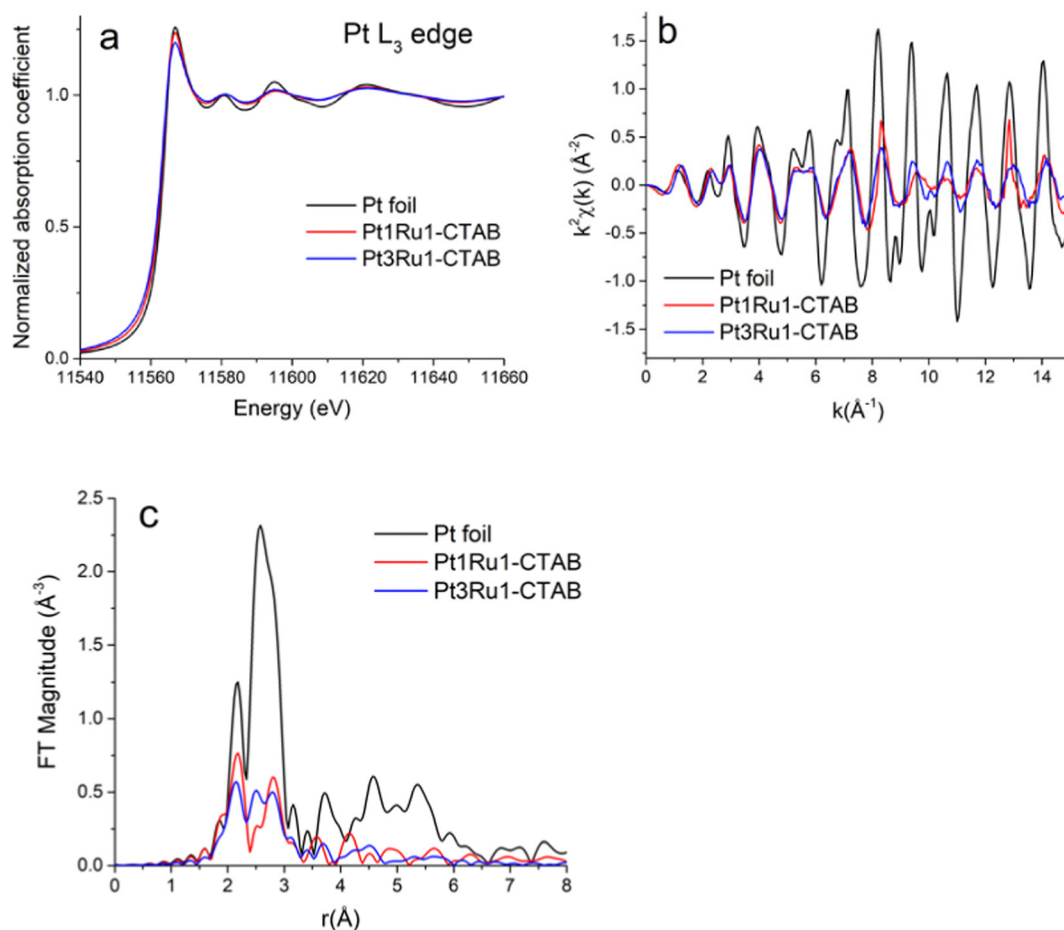


Fig. 4 (A) Pt L<sub>3</sub>-edge XANES data. (B)  $k^2$ -weighted EXAFS data in  $k$ -space. (C) Fourier transform magnitudes of the  $k^2$ -weighted EXAFS data for Pt foil, Pt<sub>1</sub>Ru<sub>1</sub> (produced with CTAB), and Pt<sub>3</sub>Ru<sub>1</sub> (produced with CTAB).

smaller for the CTAB-derived samples, as compared with their OAm/OA-produced analogues. That finding is in full agreement with the relative decrease of the metal-metal coordination numbers of the former samples as compared with the latter ones (Table 1). In both samples, the elemental maps show much smaller Ru regions on the surfaces of the larger Pt ones (Fig. 2A and 3A), in full agreement with the relationship ( $N_{\text{Pt}} > N_{\text{Ru}}$ ) obtained from EXAFS data analysis (Table 1). The only alloy sample that displays relative homogeneity ( $N_{\text{Pt}} \approx N_{\text{Ru}}$ ) is Pt<sub>3</sub>Ru<sub>1</sub>, derived from CTAB (Table 1). It can also be characterized as quasi-random, because the SRO parameter is consistent with zero (Table 1).

We note that the coordination numbers in all samples are much smaller than those predicted from the average sizes obtained by TEM (Fig. 1). Indeed, for the sizes in the range of 2.4–3.4 nm, the average coordination numbers of metal atoms,<sup>57</sup>  $N_{\text{MM}} = x_{\text{A}}N_{\text{AM}} + x_{\text{B}}N_{\text{BM}}$ , should be of the order of 10 or larger, assuming, as a simple model, an ideal cuboctahedral geometry.<sup>58</sup> This estimate is significantly larger than the coordination numbers reported in Table 1. The explanation can likely be ascribed to the strongly disordered, nanoporous-like nature of the metal ensembles (Fig. 2 and 3), which contain much larger numbers of under-coordinated atoms than those

predicted from conventional close-packed models typical of the bulk.

#### 4.2. Probing the effects of synthesis methods

To understand the differences between the OAm/OA and CTAB synthetic protocols, as noted, we synthesized ultrathin NWs of Pt<sub>1</sub>Ru<sub>1</sub> using both procedures, with equimolar amounts of the H<sub>2</sub>PtCl<sub>6</sub> and RuCl<sub>3</sub> precursors. Many aspects of the synthesis methods can impact the growth of alloyed nanomaterials. Important reaction variables that we can reliably alter include but are not limited to the reaction temperature, surfactant identity, and reduction mechanism. Each of these parameters will be explored in the context of the growth of the wires and particles. As mentioned in Section 3.1, the CTAB method produced a network of tiny nanowires. In this methodology, CTAB forms worm-like micelles at the interface between water and chloroform. The long non-polar tails of CTAB make the surfactant readily soluble in chloroform, thereby enabling the formation of micelles with the hydrophobic tails exposed to chloroform and the more hydrophilic polar heads positioned away from the organic solvent.

When water, containing dissolved Pt<sup>4+</sup> and Ru<sup>3+</sup> ions, is introduced to the chloroform, an immiscible emulsion is



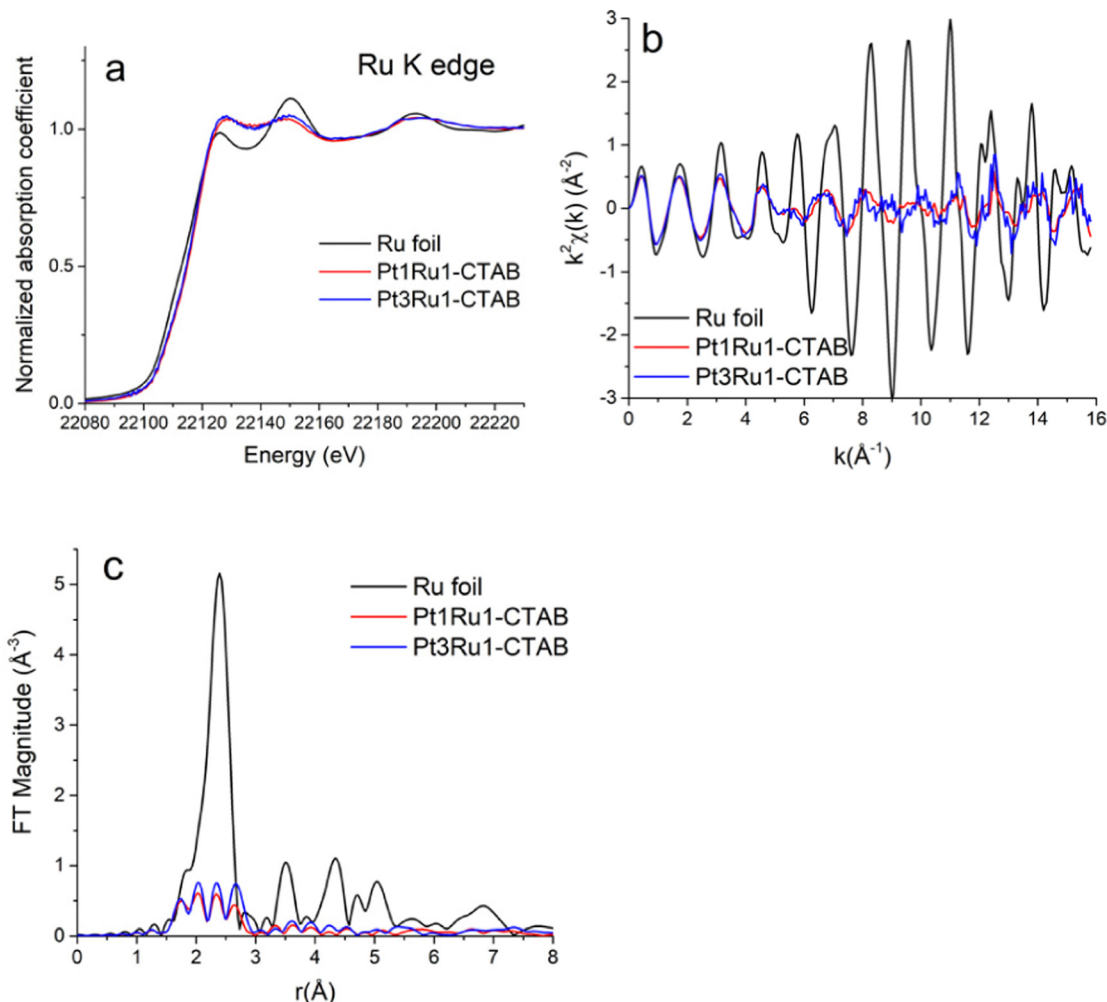


Fig. 5 (A) Ru K-edge XANES data. (B)  $k^2$ -weighted EXAFS data in  $k$ -space. (C) Fourier transform magnitudes of the  $k^2$ -weighted EXAFS data for Ru foil, Pt<sub>1</sub>Ru<sub>1</sub> (produced with CTAB), and Pt<sub>3</sub>Ru<sub>1</sub> (generated with CTAB).

subsequently formed. With the addition of vigorous stirring between water and chloroform, the micelles themselves are impacted by the presence of the metal ions. Specifically, at the interface of the two solvents, the dissolved metal ions are able to coordinate onto the polar heads of CTAB. When sodium borohydride ( $\text{NaBH}_4$ ) is then added into the reaction medium, it reduces  $\text{Pt}^{4+}$  and  $\text{Ru}^{3+}$  to  $\text{Pt}^0$  and  $\text{Ru}^0$ , respectively. The role of CTAB therefore is to spatially confine the growth of the reduced metals to the localized area contained within the micelles themselves, thereby allowing for the formation of isolated motifs of ultrathin nanowires.<sup>37</sup>

As mentioned, the OAm/OA procedure yielded a more globular nano-porous agglomerate structure. We postulate that OAm can simultaneously act as a solvent, a reducing agent, and a surfactant, whereas oleic acid functions as an additional “shape-control” ligand.<sup>39</sup> The interplay between these two species can be complex and is not fully understood. Nevertheless, it has been shown in the past that both OAm and OA are capping agents that can control the morphology of nanomaterials by binding onto their external surfaces and selectively

directing their growth; it is not surprising therefore that a particulate morphology was generated in this case.<sup>40,41,59</sup>

XRD data are provided in Fig. 7. The relatively broad XRD peaks can be ascribed to the relatively small sizes of the individual particles and wires. For samples of Pt<sub>3</sub>Ru<sub>1</sub> CTAB, Pt<sub>1</sub>Ru<sub>1</sub> CTAB, and Pt<sub>1</sub>Ru<sub>1</sub> OAm, we observed the expected pattern for fcc platinum with no apparent impurity phases of ruthenium. For the Pt<sub>1</sub>Ru<sub>0</sub> sample, we noted the pattern for hcp ruthenium with no platinum phases present. The lack of any obvious impurity phases is a good indication of successful alloying, as it suggests that there was no segregation of crystal phases.

As indicated earlier, both the CTAB and OAm/OA-mediated reactions utilize surfactants to control the growth of as-prepared PtRu nanomaterials. Surfactants can be designated into several broad categories, encompassing ionic (*i.e.*, anionic/cationic), charged non-ionic, non-charged, and polymeric species.<sup>60,61</sup> The interactions between the surfactants and the growing particles can fundamentally impact how they form and grow, because of the extent to which these organic molecules



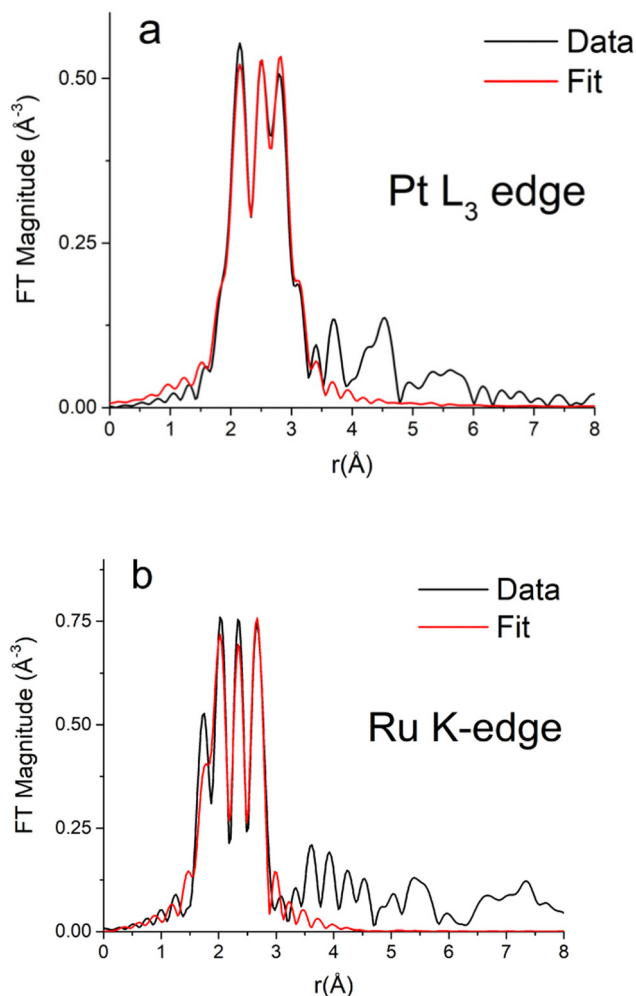


Fig. 6 Fourier transform magnitudes of  $k^2$ -weighted EXAFS data and theoretical fits for the  $\text{Pt}_3\text{Ru}_1$  (generated with CTAB) sample at (A) the  $\text{Pt}$   $L_3$ -edge and (B) the  $\text{Ru}$   $K$ -edge.

Table 1 The values of the total coordination numbers  $N_{\text{Pt}}$  and  $N_{\text{Ru}}$ , the alloy concentrations,  $x_{\text{A}}$  and  $x_{\text{B}}$  and the short-range order parameters,  $\alpha_{\text{PtRu}}$  and  $\alpha_{\text{RuPt}}$

Ultrathin NW – synthesis method	$N_{\text{Pt}}$	$N_{\text{Ru}}$	$x_{\text{Pt}}$	$x_{\text{Ru}}$	$\alpha_{\text{PtRu}}$	$\alpha_{\text{RuPt}}$
$\text{Pt}_1\text{Ru}_1$ -CTAB	6.9(7)	4.3(8)	0.6(4)	0.4(1)	0.5(5)	0.3(3)
$\text{Pt}_3\text{Ru}_1$ -CTAB	6.4(4)	7.1(1.6)	0.9(6)	0.12(4)	0(0)	0.1(1)
$\text{Pt}_1\text{Ru}_1$ -OAm/OA	9.4(1.5)	8.1(1.4)	0.5(3)	0.5(2)	0.5(5)	0.4(4)
$\text{Pt}_1\text{Ru}_9$ -OAm/OA	9.7(2.0)	4.6(6)	0.3(1)	0.7(5)	0.7(8)	0.4(5)

will bind (or not) onto exposed facets. In particular, CTAB can be characterized as a cationic surfactant, containing a negatively charged bromide ion coupled with a positively charged organic component. Table S3 (ESI<sup>†</sup>) gives the chemical structure of the various surfactants used herein.

In a typical CTAB-based method, a ‘soft’ template consisting of worm-like micellar networks is produced in the presence of CTAB within a two-phase water-chloroform system. The

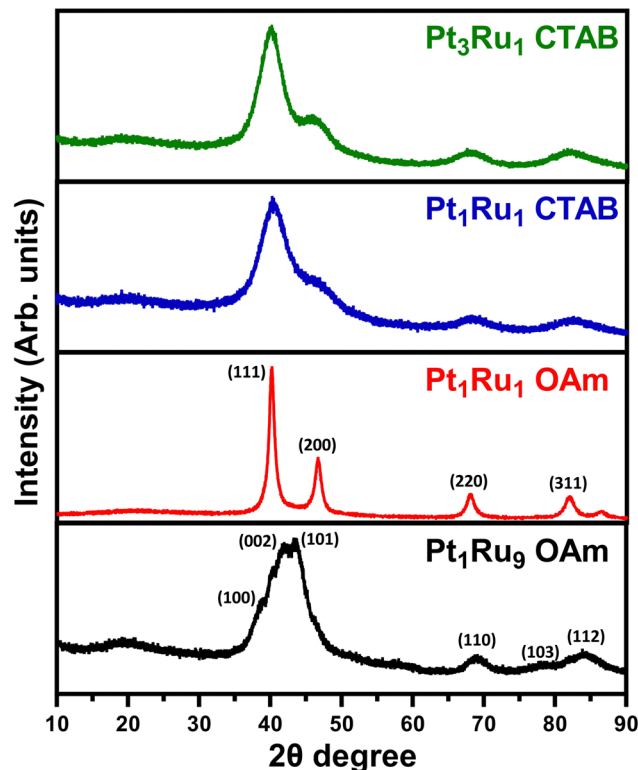


Fig. 7 XRD data for all of the alloyed samples generated.

charged  $\text{Pt}$  and  $\text{Ru}$  precursor ions, added to the mixture, are spatially confined to and subsequently trapped within the ‘micellar pores’, *i.e.*, channels created by the packing and alignment of the highly polar heads of individual CTAB molecules. Hence, porous, high surface area networks of interconnected, ‘worm-like’, and homogeneous metallic nanowires, with average diameters of as small as 1.9 nm, are able to form upon the reduction of these metal precursors with a reducing agent, such as sodium borohydride.<sup>62</sup>

By contrast, as previously mentioned, with the OAm/OA technique, oleic acid and oleylamine function as both surfactants and solvents. In this procedure, unlike with the CTAB-based method, the surfactants are charged but not necessarily closely coupled with a separate counterion, as is the case with the bromide anion present within CTAB. Hence, for our purposes, these smaller surfactants are considered to be charged but non-ionic. In this mixture, both the OA and OAm molecules will form coordination complexes with the  $\text{Pt}$  and  $\text{Ru}$  ions.<sup>39</sup> It is likely that the negatively charged oleic acid will coordinate more strongly with the positively charged  $\text{Pt}^{4+}$  and  $\text{Ru}^{3+}$  ions. Nevertheless, both oleic acid and oleylamine contain similar carbon chains that are 18 carbon atoms long. This renders these surfactants as highly miscible with each other, which likely leads to a more even distribution of precursor elements within the entire solution, thereby enabling the formation of more homogeneous, particulate-like structures as opposed to the worm-like micellar motifs commonly observed with the CTAB method.



With respect to the reducing mechanism, in the CTAB procedure, the precursors are reduced very rapidly, due to the presence of a strong reducing agent, namely  $\text{NaBH}_4$ . By contrast, with the OAm/OA procedure, OAm embraces multiple, simultaneous roles as a solvent, surfactant, and reducing agent. Moreover, because OAm as a reducing agent is a much “weaker” reducing agent than that of  $\text{NaBH}_4$ , it is not surprising that higher temperatures and longer reaction times are needed to facilitate the formation of uniform particles with the OAm/OA process.

From the EXAFS perspective, we observed that the as-synthesized  $\text{Pt}_1\text{Ru}_1$  samples possessed an evident degree of heterogeneity with the Pt and Ru atoms preferentially segregated. This level of segregation is consistent with past results for similarly formed types of NWs, such as  $\text{Ru}_2\text{Co}$ .<sup>39</sup> It is also worth noting that hydrothermally generated PtRu was similarly created preferentially in the form of segregated ultrathin nanowires.

#### 4.3. Probing the effects of varying chemical composition

To confirm the chemical composition of all samples, we looked at both the concentrations calculated from the EXAFS measurements and the atomic % values gathered from the EDS measurements (see Table 2). All of these samples consistently highlight slightly more Pt along with less Ru content than the initial molar ratios mixed during the reactions. For example, the PtRu 1:1 sample generated through the CTAB method yielded a chemical composition closer to that of 6:4 case, based on both the EDS and EXAFS results. Moreover, the PtRu 1:9 OAm sample was characterized by a measured composition of about 3:7. This higher-than-expected amount of Pt may be ascribed to the more positive reduction potential of Pt, which implies a faster reduction process, and consequently, the observed hcp/fcc crystal structure mismatch between Ru and Pt respectively.

As earlier noted, for the CTAB-based method, both  $\text{Pt}_3\text{Ru}_1$  and  $\text{Pt}_1\text{Ru}_1$  were successfully synthesized as ultrathin NWs. For the sake of completeness, we also attempted to synthesize  $\text{Pt}_1\text{Ru}_9$ . However, this latter attempt resulted in the formation of NPs with an average size of 1–3 nm and not ultrathin nanowires (see ESI,† Fig. S8A). This observation implies that a sufficiently high enough relative concentration of Pt is

necessary to enable wire formation using this particular protocol.

With regards to the OAm/OA method, we successfully generated both  $\text{Pt}_1\text{Ru}_1$  and  $\text{Pt}_1\text{Ru}_9$  compositions. However, with  $\text{Pt}_3\text{Ru}_1$ , the resulting product consisted of irregular nanoparticles and aggregates. Fig. S8B (ESI†) highlights TEM images of the  $\text{Pt}_3\text{Ru}_1$  OAm sample. The implication was that excess Pt degraded the observed morphology, when using the OAm/OA method. For the CTAB samples, our EXAFS analysis demonstrated that  $\text{Pt}_3\text{Ru}_1$  could be described as a random alloy, whereas  $\text{Pt}_1\text{Ru}_1$  yielded a degree of segregation between the elements. We note that the  $\text{Pt}^{4+} \rightarrow \text{Pt}^0$  reduction process is characterized by a reduction potential of 0.69 V and that the corresponding  $\text{Ru}^{3+} \rightarrow \text{Ru}^0$  reduction process has an analogous reduction potential of 0.60 V. Based on these facts, both metals should reduce under similar reaction conditions. However, the preferred crystal structure of  $\text{Pt}^0$  is fcc, whereas that of  $\text{Ru}^0$  is hexagonal closed packed (hcp). This structural difference can help to explain why Ru and Pt appear to segregate from one another at higher relative concentrations; that is, the mismatch between the crystal structures of these two elements renders homogeneous alloy formation difficult but not outright impossible. Additionally, we observed that the wire-like morphology is only achievable with large amounts of Pt, whereas Ru tends to preferentially form NPs.

Concerning the OAm method, the differences we observe may be ascribed in part to the differing preferred crystal structures of Pt and Ru, previously discussed, and the corresponding slow reduction process. Although the reduction potentials of  $\text{Pt}^{4+}$  and  $\text{Ru}^{3+}$  are similar and as such, both species should reduce essentially simultaneously, that of Pt is slightly more positive. In practice, in a sample with a comparatively higher concentration of Pt, relatively more Pt may form first and the Ru may either generate more slowly or not alloy as effectively at all. Therefore, as higher and higher amounts of Pt are introduced, these will preferentially form Pt particles first and will not necessarily alloy with the slower-forming  $\text{Ru}^0$  species. EXAFS analysis appeared to indicate that both compositions created from the OAm process gave rise to perceptible elemental segregation.

## 5. MOR

Our ultimate goal was to understand the effects of not only the synthesis method but also the inherent chemical composition of these samples on the observed electrochemical performance of the PtRu alloys. To do this, in separate runs, we deposited the ultrathin wires and spoke-like, dendritic clusters onto Vulcan carbon XC-72R with a nominal loading of about 20%. Since drying the samples can lead to significant aggregation, we confirmed the actual loading quantities using TGA, after the alloyed samples had been combined with the carbon support. Fig. S9 (ESI†) provides for TGA mass loss curves for the loading amounts of either Pt or PtRu onto the carbon support. These

**Table 2** Chemical composition data for all samples, extracted from the EXAFS and EDS results

Sample	Chemical concentration derived from XAS	Chemical composition derived from EDS (atomic%)
$\text{Pt}_1\text{Ru}_1$ -CTAB	Pt: 0.6(4)	Pt: $64 \pm 8.2\%$
	Ru: 0.4(1)	Ru: $36 \pm 8.2\%$
$\text{Pt}_3\text{Ru}_1$ -CTAB	Pt: 0.9(6)	Pt: $88 \pm 3.0\%$
	Ru: 0.12(4)	Ru: $12 \pm 3.0\%$
$\text{Pt}_1\text{Ru}_1$ -OAm/OA	Pt: 0.5(3)	Pt: $77 \pm 2.9\%$
	Ru: 0.5(2)	Ru: $23 \pm 2.9\%$
$\text{Pt}_1\text{Ru}_9$ -OAm/OA	Pt: 0.3(1)	Pt: $25 \pm 3.4\%$
	Ru: 0.7(5)	Ru: $75 \pm 3.4\%$



values were subsequently used to calculate the specific surface area (SSA) for each sample, as shown in ESI,† Table S4.

Representative CV curves for the samples are given in ESI,† Fig. S10. All show the expected hydrogen adsorption and desorption peaks situated at around 0–0.2 V, which is characteristic of platinum. We detected little to no Pt–O reduction peaks present, likely due to their being suppressed by the incorporation of Ru. MOR LSVs and the associated specific activities (SA) are shown in Fig. 8.

All of the SA values were acquired at 0.7 V vs. RHE. Specifically, Pt<sub>3</sub>Ru<sub>1</sub> and Pt<sub>1</sub>Ru<sub>1</sub> derived from the CTAB method yielded specific activities of 1.22 mA cm<sup>-2</sup> and 0.55 mA cm<sup>-2</sup>, respectively. By contrast, Pt<sub>1</sub>Ru<sub>1</sub> and Pt<sub>1</sub>Ru<sub>9</sub> created from the OAm/OA protocol gave rise to corresponding activity values of 0.46 mA cm<sup>-2</sup> and 0.07 mA cm<sup>-2</sup>, respectively. We compared these results with those of pure Pt wires generated using the CTAB method. In effect, as a control sample, pure Pt NWs were characterized by an SA of 0.17 mA cm<sup>-2</sup>, suggesting that the best performing PtRu alloys exhibited an almost 7-fold increase in activity as compared with Pt alone. Approximately the same increase in SA was observed when comparing the behavior of as-synthesized PtRu alloys with those of commercial Pt/C nanoparticles.

It has been postulated that the MOR enhancement of PtRu alloys is due to not only (i) their enabling of the direct conversion of methanol to CO<sub>2</sub> (without any intermediate CO through a non-CO pathway) but also (ii) the stimulation of CH<sub>3</sub>OH adsorption and concomitant formation of active OH species from H<sub>2</sub>O which can oxidize intermediate (and poisoning)<sup>63</sup> CO bound to Pt sites.<sup>64</sup> Moreover, the strain effect<sup>65</sup> resulting from the lattice mismatch between Pt and Ru can decrease the binding energy of reaction intermediates, thereby favoring high CO tolerance.

To confirm that these apparent differences could not be ascribed to the degradation of either the wires or dendritic particulate clusters, we obtained TEM images of the samples, both before and after the MOR electrochemical process. These microscopy data are provided in ESI,† Fig. S11 for the sample synthesized by the OAm/OA method and in ESI,† Fig. S12 for that generated from the corresponding CTAB protocol. The corresponding TEM images of the analogous pure Pt/C NW system are given in ESI,† Fig. S13. We found little to no sample degradation, and in fact, there was no obvious change in the degree of sample aggregation after running MOR for our PtRu alloys. By contrast, in the case of the pure Pt NW controls, we observed the presence of an obvious anisotropic wire

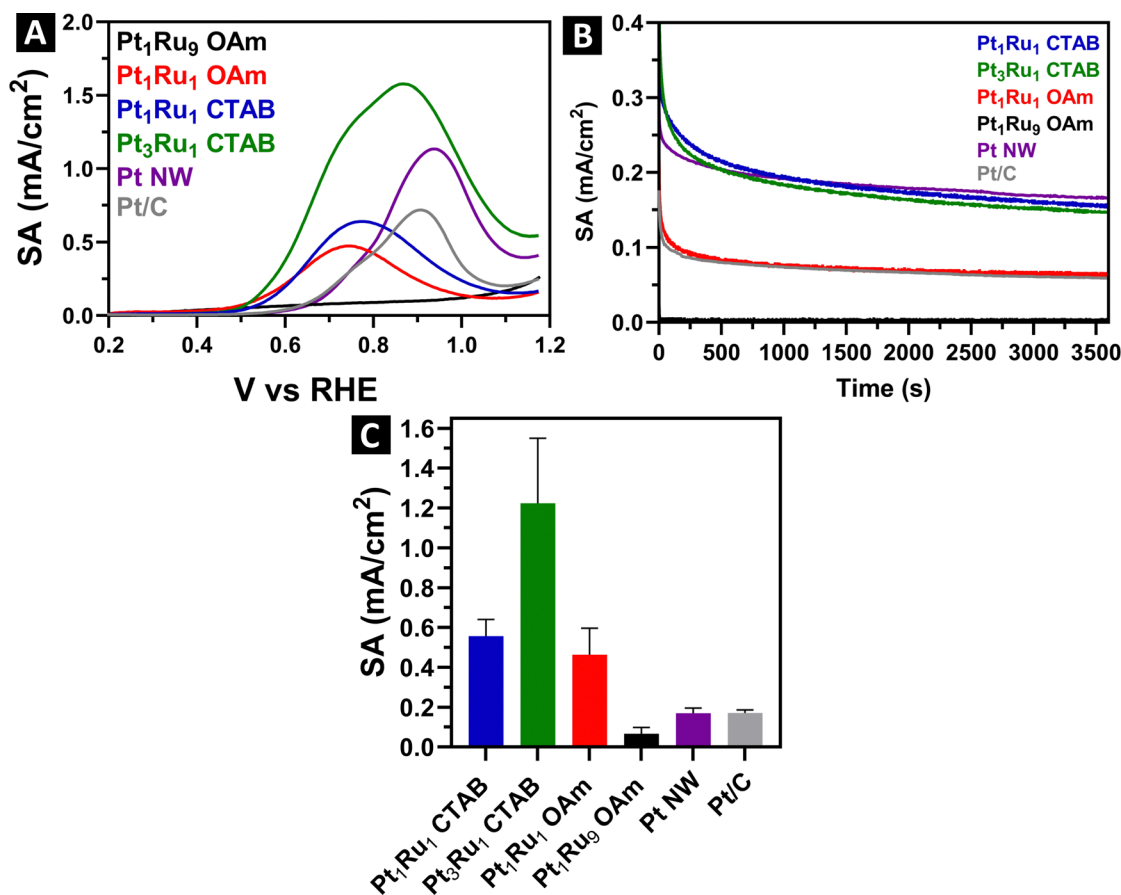


Fig. 8 MOR results for all PtRu alloyed ultrathin wires as well as particulate clusters, in addition to the Pt nanowire and Pt/C commercial control samples, with their corresponding synthesis methods and surfactants. Data shown herein include (A) MOR LSVs, (B) chronoamperometry plots, and (C) bar graphs of the specific activity measurements of the various nanoscale motifs.



morphology prior to MOR but after MOR, there was noticeably more sample aggregation. All of these results collectively indicate that the incorporation of Ru into our samples yielded tangibly better structural and operational stability towards MOR in an acidic environment as compared with pure monometallic Pt NWs.

In terms of chemical composition, we noted a direct correlation between the MOR performance and the amount of Pt incorporated within the alloyed samples. Not surprisingly, Pt<sub>3</sub>Ru<sub>1</sub> containing the highest relative Pt content gave rise to the best performance detected, whereas Pt<sub>1</sub>Ru<sub>9</sub> evinced almost no measurable MOR activity. Structurally, as we have seen from EXAFS, the Pt<sub>3</sub>Ru<sub>1</sub> sample maintains the greatest amount of alloying, as indicated by its SRO parameter of about 0.1. Hence, we postulate that Pt<sub>3</sub>Ru<sub>1</sub> behaves the best for MOR, in part because of a higher degree of coordination between the Pt and Ru atoms, and in fact, per ESI,† Table S2, the Ru–Pt coordination number ( $N_{\text{Ru-Pt}}$ ) of 5.6 is at least 2–3 times higher than of the other samples tested.

We note that the  $N_{\text{Ru-Pt}}$  of Pt<sub>1</sub>Ru<sub>1</sub>-OAm sample was 2.7, which is a higher figure than that of the 1.9 value computed for Pt<sub>1</sub>Ru<sub>1</sub>-CTAB. However, the MOR values associated with these samples were very similar, with the NWs created by the CTAB method yielding a slighter better performance metric. This observation indicates that as a tunable parameter, the degree of alloying, in the samples possessing the same apparent composition, in and of itself, is not sufficient to dramatically affect the MOR performance. Instead, the electrocatalytic performance of these samples would appear to be impacted more significantly by morphological and size differences. In particular, the smaller diameters of the NWs generated using the CTAB method gave rise to better performance than their larger-diameter analogues.

Moreover, both of the Pt<sub>1</sub>Ru<sub>1</sub> samples created with OAm/OA and CTAB procedures possessed very similar SA values. This finding indicates that MOR performance is much more connected to and can be viewed as a function of the actual Pt–Ru composition, which to a large extent is independent of the synthesis method employed. It is worth highlighting that the EXAFS data for both OAm/OA and CTAB Pt<sub>1</sub>Ru<sub>1</sub> samples are also relatively comparable to each other, in that both materials possess almost identical SRO values.

To further probe and understand the differences in performance between the samples, we acquired chronoamperometry measurements, provided in Fig. 8B. We ran these measurements at a steady state voltage of 0.7 V vs. RHE for 3600 s. Pt is easily poisoned by CO, so we can attribute the initial rapid drop to the de-hydrogenation of methanol followed by the poisoning of accessible Pt active sites with CO. Steady state SA measurements indicate that both samples generated using the CTAB method yielded the highest durability and stability, followed by the Pt<sub>1</sub>Ru<sub>1</sub> sample produced using the OAm/OA method. As expected, the Pt<sub>1</sub>Ru<sub>9</sub> sample gave rise to almost no steady state performance, in line with its very low MOR SA. We also observed that the steady state SA values of pure Pt are similar to that of Pt<sub>7</sub>Ru<sub>3</sub> and Pt<sub>1</sub>Ru<sub>1</sub> generated using the CTAB method,

thereby indicating that our PtRu alloys and monometallic Pt controls maintain comparable levels of tolerance to CO.

Interestingly, the chronoamperometry results imply a significant discrepancy between synthesis protocols, that had not been apparent with MOR performance data alone. Specifically, the Pt<sub>1</sub>Ru<sub>1</sub> produced with CTAB gave rise to a steady state SA that was more than 2× higher than that of the identical composition fabricated with the OAm/OA procedure. We assert that this observation is associated with the size and morphology of the ultrathin nanowires, made using the CTAB method, *versus* that of the discrete clusters, produced with the analogous OAm/OA protocol. It is worth noting that we have observed similar analogous improvements in the electrochemical performance of our ultrathin motifs *versus* that of their corresponding particulate morphologies in a number of our prior studies.<sup>25–38</sup>

In general, ultrathin anisotropic nanowires not only are smaller in diameter but also possess lower total coordination numbers than their more isotropic particulate cluster counterparts. Hence, considering that our EXAFS analysis was suggestive of a core–shell-like distribution of elements with Pt preferentially localized in the core and Ru confined to the shell, it is reasonable to hypothesize a greater preponderance and proportion of Pt atoms situated on the outer surfaces of the nanowires, mixed in with the adjacent Ru atoms, thereby leading to potentially higher numbers of available active surface sites, which are conducive factors to enabling an overall improvement in measured MOR activity especially as compared with pure Pt alone.

## 6. Conclusions

We have successfully synthesized not only ultrathin Pt<sub>1</sub>Ru<sub>1</sub> and Pt<sub>3</sub>Ru<sub>1</sub> nanowires using the CTAB method but also more organized ensembles, consisting of localized spoke-like, dendritic clusters of Pt<sub>1</sub>Ru<sub>1</sub> and Pt<sub>1</sub>Ru<sub>9</sub>, utilizing an effective, in-house, solution-based OAm/OA protocol that we have generalized from mono-metallic to more complex bi-metallic alloy species. HRTEM EDS elemental mapping data indicated that all of the samples gave rise to relatively disordered groupings of atoms to varying degrees, depending on the precise composition. As an example, Pt:Ru 1:1 samples, generated through either the CTAB or OAm/OA-derived protocols, appear to maintain a Pt core/Ru shell structure. This finding was corroborated by our XAS results which were overall suggestive of elemental segregation. Interestingly, Table 1 shows that the  $N_{\text{Pt}}$  and  $N_{\text{Ru}}$  values are significantly different for both Pt<sub>1</sub>Ru<sub>1</sub> samples. Specifically, (i) both values are larger in the CTAB sample as compared with the OAm/OA-derived one and (ii)  $N_{\text{Pt}} > N_{\text{Ru}}$  in both types of samples, suggesting that Pt is localized within the core, while Ru is preferentially arranged at the surface.

Therefore, highlighting the key role of the synthesis methodology in dictating the observed morphology, we postulate that the reducing environment within the CTAB method enables the Ru and Pt precursors to be reduced relatively



quickly at room temperature with  $\text{NaBH}_4$ . By contrast, since OAm is a weaker reducing agent than  $\text{NaBH}_4$ , more highly elevated temperatures are required to enable the reduction process, thereby leading to greater elemental segregation and architectural complexity. With respect to the equally important role of the viability of generating products with specific chemical compositions, we determined that the mismatch in the crystal structures of the fcc Pt vs. the hcp Ru likely leads to the presence of increased elemental segregation at higher loadings of Ru, irrespective of the synthesis method employed.

Finally, MOR measurements indicate a direct correlation and connection between both chemical composition and the degree of alloying versus the electrochemical performance. In particular, our  $\text{Pt}_3\text{Ru}_1$  sample yielded an almost  $2\times$  greater SA value as compared with analogous samples tested. We explain these observations, based on the greater relative Pt content coupled with the high degree of alloying associated with our  $\text{Pt}_3\text{Ru}_1$  sample, as evinced by its small, positive SRO parameter of 0.1. Moreover, the CTAB protocol also produced ultrathin alloyed nanowires that not only appeared to be more tolerant to CO poisoning than nanoscale motifs generated using the OAm/OA fabrication method based on our chronoamperometry data but also gave rise to a  $7\times$  increase in performance as compared with pure Pt NWs alone. A comparison of our PtRu results versus the corresponding MOR data of various PtRu alloys previously reported in the literature emphasizes that the findings discussed herein are certainly within the expected range of prior values previously noted with PtRu (Table S5, ESI†).

## Conflicts of interest

There are no conflicts to declare.

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## References

- 1 Y. Shao, G. Yin and Y. Gao, *J. Power Sources*, 2007, **171**, 558–566.
- 2 J. Zhang, K. Sasaki, E. Sutter and R. R. Adzic, *Science*, 2007, **315**, 220–222.
- 3 L. Swette and N. Kackley, *J. Power Sources*, 1990, **29**, 423–436.
- 4 L. Li and S. S. Wong, *ACS Omega*, 2018, **3**, 3294–3313.
- 5 C. Koenigsmann, M. E. Scofield, H. Liu and S. S. Wong, *J. Phys. Chem. Lett.*, 2012, **3**, 3385–3398.
- 6 J. L. Fernandez, D. A. Walsh and A. J. Bard, *J. Am. Chem. Soc.*, 2005, **127**, 357–365.
- 7 J. Greeley and J. K. Nørskov, *J. Phys. Chem. C*, 2009, **113**, 4932–4939.
- 8 G. A. Camara, M. J. Giz, V. A. Paganin and E. A. Ticianelli, *J. Electroanal. Chem.*, 2002, **537**, 21–29.
- 9 H. Liu, C. Song, L. Zhang, J. Zhang, H. Wang and D. P. Wilkinson, *J. Power Sources*, 2006, **155**, 95–110.
- 10 J.-H. Wee and K.-Y. Lee, *J. Power Sources*, 2006, **157**, 128–135.
- 11 X. Zhang and K.-Y. Chan, *Chem. Mater.*, 2002, **15**, 451–459.
- 12 R. Basnayake, Z. Li, S. Katar, W. Zhou, H. Rivera, E. S. Smotkin, D. J. Casadonte and C. Korzeniewski, *Langmuir*, 2006, **22**, 10446–10450.
- 13 S. Sun, H. Xu, S. Tang, J. Guo, H. Li, L. Cao, B. Zhou, Q. Xin and G. Sun, *Chin. J. Catal.*, 2006, **27**, 932–936.
- 14 G.-Y. Zhao, C.-L. Xu, D.-J. Guo, H. Li and H.-L. Li, *J. Power Sources*, 2006, **162**, 492–496.
- 15 B. C. H. Steele and A. Heinzl, *Nature*, 2001, **414**, 345–352.
- 16 S. Garbarino, A. Ponrouch, S. Pronovost and D. Guay, *Electrochem. Commun.*, 2009, **11**, 1449–1452.
- 17 H. A. Gasteiger, N. Markovic, P. N. Ross and E. J. Cairns, *J. Phys. Chem.*, 1993, **97**, 12020–12029.
- 18 P. Liu and J. K. Nørskov, *Fuel Cells*, 2001, **1**, 192–201.
- 19 W. Zhao, D. Huang, Q. Yuan and X. Wang, *Nano Res.*, 2016, **9**, 3066–3074.
- 20 W. Zhao, B. Ni, Q. Yuan, Y. Wang, Q. Zhang and X. Wang, *Langmuir*, 2017, **33**, 8070–8075.
- 21 Z. Shen, M. Yamada and M. Miyake, *Chem. Commun.*, 2007, 245–247.
- 22 N. Jha, A. L. M. Reddy, M. M. Shaijumon, N. Rajalakshmi and S. Ramaprabhu, *Int. J. Hydrogen Energy*, 2008, **33**, 427–433.
- 23 S. Yang, F. Hong, L. Wang, S. Guo, X. Song, B. Ding and Z. Yang, *J. Phys. Chem. C*, 2009, **114**, 203–207.
- 24 H. Liu, C. Koenigsmann, R. R. Adzic and S. S. Wong, *ACS Catal.*, 2014, **4**, 2544–2555.
- 25 C. Koenigsmann, W.-p. Zhou, R. R. Adzic, E. Sutter and S. S. Wong, *Nano Lett.*, 2010, **10**, 2806–2811.



- 26 C. Koenigsmann and S. S. Wong, *Energy Environ. Sci.*, 2011, **4**, 1161–1176.
- 27 C. Koenigsmann, E. Sutter, R. R. Adzic and S. S. Wong, *J. Phys. Chem. C*, 2012, **116**, 15297–15306.
- 28 N. P. Tarasova, Y. V. Smetannikov and A. A. Zanin, *Russ. Chem. Rev.*, 2010, **79**, 463–477.
- 29 X. Teng, M. Feyngenson, Q. Wang, J. He, W. Du, A. I. Frenkel, W. Han and M. Aronson, *Nano Lett.*, 2009, **9**, 3177–3184.
- 30 H. Zhou, W.-p Zhou, R. R. Adzic and S. S. Wong, *J. Phys. Chem. C*, 2009, **113**, 5460–5466.
- 31 W.-P. Zhou, M. Li, C. Koenigsmann, C. Ma, S. S. Wong and R. R. Adzic, *Electrochim. Acta*, 2011, **56**, 9824–9830.
- 32 M. E. Scofield, Y. Zhou, S. Yue, L. Wang, D. Su, X. Tong, M. B. Vukmirovic, R. R. Adzic and S. S. Wong, *ACS Catal.*, 2016, **6**, 3895–3908.
- 33 M. E. Scofield, C. Koenigsmann, D. Bobb-Semple, J. Tao, X. Tong, L. Wang, C. S. Lewis, M. B. Vukmirovic, Y. Zhu, R. R. Adzic and S. S. Wong, *Catal.: Sci. Technol.*, 2016, **6**, 2435–2450.
- 34 H. Liu, R. R. Adzic and S. S. Wong, *ACS Appl. Mater. Interfaces*, 2015, **7**, 26145–26157.
- 35 C. Koenigsmann, A. C. Santulli, K. Gong, M. B. Vukmirovic, W.-P. Zhou, E. Sutter, S. S. Wong and R. R. Adzic, *J. Am. Chem. Soc.*, 2011, **133**, 9783–9795.
- 36 C. Koenigsmann, D. B. Semple, E. Sutter, S. E. Tobierre and S. S. Wong, *ACS Appl. Mater. Interfaces*, 2013, **5**, 5518–5530.
- 37 M. E. Scofield, C. Koenigsmann, L. Wang, H. Liu and S. S. Wong, *Energy Environ. Sci.*, 2015, **8**, 350–363.
- 38 L. Li, H. Liu, C. Qin, Z. Liang, A. Scida, S. Yue, X. Tong, R. R. Adzic and S. S. Wong, *ACS Appl. Nano Mater.*, 2018, **1**, 1104–1115.
- 39 S. C. McGuire, A. M. Ebrahim, N. Hurley, L. Zhang, A. I. Frenkel and S. S. Wong, *Chem. Sci.*, 2021, **12**, 7158–7173.
- 40 A. Halder and N. Ravishankar, *Adv. Mater.*, 2007, **19**, 1854–1858.
- 41 S. Mourdikoudis and L. M. Liz-Marzan, *Chem. Mater.*, 2013, **25**, 1465–1476.
- 42 Z. Peng, H. You and H. Yang, *ACS Nano*, 2010, **4**, 1501–1510.
- 43 N. Poudyal, G. S. Chaubey, V. Nandwana, C.-B. Rong, K. Yano and J. P. Liu, *Nanotechnology*, 2008, **19**, 355601.
- 44 A. I. Frenkel, *Chem. Soc. Rev.*, 2012, **41**, 8163–8178.
- 45 R. M. Anderson, L. Zhang, J. A. Loussaert, A. I. Frenkel, G. Henkelman and R. M. Crooks, *ACS Nano*, 2013, **7**, 9345–9353.
- 46 J. Timoshenko, C. J. Wrasman, M. Luneau, T. Shirman, M. Cargnello, S. R. Bare, J. Aizenberg, C. M. Friend and A. I. Frenkel, *Nano Lett.*, 2019, **19**, 520–529.
- 47 A. I. Frenkel, V. S. Machavariani, A. Rubshtein, Y. Rosenberg, A. Voronel and E. A. Stern, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2000, **62**, 9364–9371.
- 48 S.-A. Chen, Y.-C. Liang, K.-T. Lu, C.-W. Pao, J.-F. Lee, T.-L. Lin and J.-M. Chen, *Phys. Chem. Chem. Phys.*, 2014, **16**, 3939–3945.
- 49 S. Alayoglu, P. Zavalij, B. Eichhorn, Q. Wang, A. I. Frenkel and P. Chupas, *ACS Nano*, 2009, **3**, 3127–3137.
- 50 S. Stoupin, E.-H. Chung, S. Chattopadhyay, C. U. Segre and E. S. Smotkin, *J. Phys. Chem. B*, 2006, **110**, 9932–9938.
- 51 S. Stoupin, H. Rivera, Z. Li, C. U. Segre, C. Korzeniewski, D. J. Casadonte Jr., H. Inoue and E. S. Smotkin, *Phys. Chem. Chem. Phys.*, 2008, **10**, 6430–6437.
- 52 S.-H. Liu, W.-Y. Yu, C.-H. Chen, A.-Y. Lo, B.-J. Hwang, S.-H. Chien and S.-B. Liu, *Chem. Mater.*, 2008, **20**, 1622–1628.
- 53 B. Ravel and M. Newville, *J. Synchrotron Radiat.*, 2005, **12**, 537–541.
- 54 A. I. Frenkel, Q. Wang, S. I. Sanchez, M. W. Small and R. G. Nuzzo, *J. Chem. Phys.*, 2013, **138**, 064202.
- 55 J. M. Cowley, *Phys. Rev.*, 1965, **138**, 1384–1389.
- 56 S. C. McGuire, A. M. Ebrahim, N. Hurley, L. Zhang, A. I. Frenkel and S. S. Wong, *Chem. Sci.*, 2021, **12**, 7158–7173.
- 57 A. Frenkel, *Z. Kristallogr. – Cryst. Mater.*, 2007, **222**, 605–611.
- 58 D. Glasner and A. I. Frenkel, *AIP Conf. Proc.*, 2007, **882**, 746–748.
- 59 N. Poudyal, G. S. Chaubey, V. Nandwana, C.-B. Rong, K. Yano and J. P. Liu, *Nanotechnology*, 2008, **19**, 355601.
- 60 T. F. Tadros, *An Introduction to Surfactants*, De Gruyter, 2014.
- 61 T. Song, F. Gao, S. Guo, Y. Zhang, S. Li, H. You and Y. Du, *Nanoscale*, 2021, **13**, 3895–3910.
- 62 Y. Song, R. M. Garcia, R. M. Dorin, H. Wang, Y. Qiu, E. N. Coker, W. A. Steen, J. E. Miller and J. A. Shelnutt, *Nano Lett.*, 2007, **7**, 3650–3655.
- 63 Y. Hu, A. Zhu, Q. Zhang and Q. Liu, *Int. J. Hydrogen Energy*, 2016, **41**, 11359–11368.
- 64 L. Zhao, S. Wang, Q. Ding, W. Xu, P. Sang, Y. Chi, X. Lu and W. Guo, *J. Phys. Chem. C*, 2015, **119**, 20389–20400.
- 65 S. Lu, K. Eid, D. Ge, J. Guo, L. Wang, H. Wang and H. Gu, *Nanoscale*, 2017, **9**, 1033–1039.

