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# Synthesis and elucidation of local structure in phase-controlled colloidal tin phosphide nanocrystals from aminophosphines†

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The chemical versatility and rich phase behavior of tin phosphides has led to interest in their use for a wide range of applications including optoelectronics, thermoelectrics, and electrocatalysis. However, researchers have identified few viable routes to high-quality, phase-pure, and phase-controlled tin phosphides. An outstanding issue is the small library of phosphorus precursors available for synthesis of metal phosphides. We demonstrated that inexpensive, commercially available, and environmentally benign aminophosphines can generate various phases of colloidal tin phosphides. We manipulated solvent concentrations, precursor identities, and growth conditions to obtain Sn<sub>3</sub>P<sub>4</sub>, SnP, and Sn<sub>4</sub>P<sub>3</sub> nanocrystals. We performed a combination of X-ray diffraction and transmission electron microscopy to determine the phase purity of our samples. X-ray absorption spectroscopy provided detailed analyses of the local structures of the tin phosphides.

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## 1. Introduction

Interest in nanoscale tin-based pnictides has recently grown, owing to the rich phase chemistry available from the multiple oxidation states demonstrated by tin (0, 2+, 4+).<sup>1,2</sup> Tin phosphides (SnP<sub>3</sub>, Sn<sub>3</sub>P<sub>4</sub>, SnP, and Sn<sub>4</sub>P<sub>3</sub>) in particular have been investigated for thermoelectrics,<sup>3</sup> photovoltaics,<sup>4</sup> superconductance,<sup>5</sup> catalysis,<sup>6–8</sup> and as anodes for Na-ion and Li-ion batteries.<sup>9–37</sup> Still, synthesis of phase-pure tin phosphides remains as a

barrier for widespread use in such applications. Tin phosphides exhibit various stoichiometries, and polymorphism has been observed for fixed stoichiometries, *e.g.* in SnP and Sn<sub>3</sub>P<sub>4</sub>.<sup>1</sup> Thus, developing facile and controlled syntheses of phase-pure tin phosphide nanostructures is essential to understanding and optimizing their use.

To date, few syntheses of nanoscale tin phosphides have been reported.<sup>17,26,33–35,39–41</sup> Researchers currently have a small library of expensive and dangerous phosphorus precursors available to them. Sn<sub>3</sub>P<sub>4</sub>, SnP, and Sn<sub>4</sub>P<sub>3</sub> nanocrystals, for example, have been synthesized from the common phosphorus precursor tris(trimethylsilyl)phosphine.<sup>39</sup> Hexamethylphosphoramide has provided phase-pure SnP nanocrystals, but other phase-pure stoichiometries were inaccessible.<sup>40</sup> We present a new route to tin phosphide nanocrystals *via* tris(diethyl)aminophosphine. Aminophosphines are a class of inexpensive, environmentally benign phosphorus precursors that are safe to use in ambient conditions.<sup>42–48</sup> Since their introduction, aminophosphines have been used to synthesize In, Co, Ni, Fe and Cd, and Cu-phosphides.<sup>43–47</sup> Unlike tris(trimethylsilyl)phosphine, which is a P(-III) source ready for reaction, aminophosphines first undergo a series of redox reactions to form an active phosphorus precursor. In previous work, combining aminophosphines with different metal halide precursors altered reaction kinetics. We explored a similar approach to synthesize Sn<sub>4</sub>P<sub>3</sub>, SnP, and Sn<sub>3</sub>P<sub>4</sub> nanocrystals.

We present structural and chemical characterization of our materials using X-ray diffraction (XRD), transmission electron

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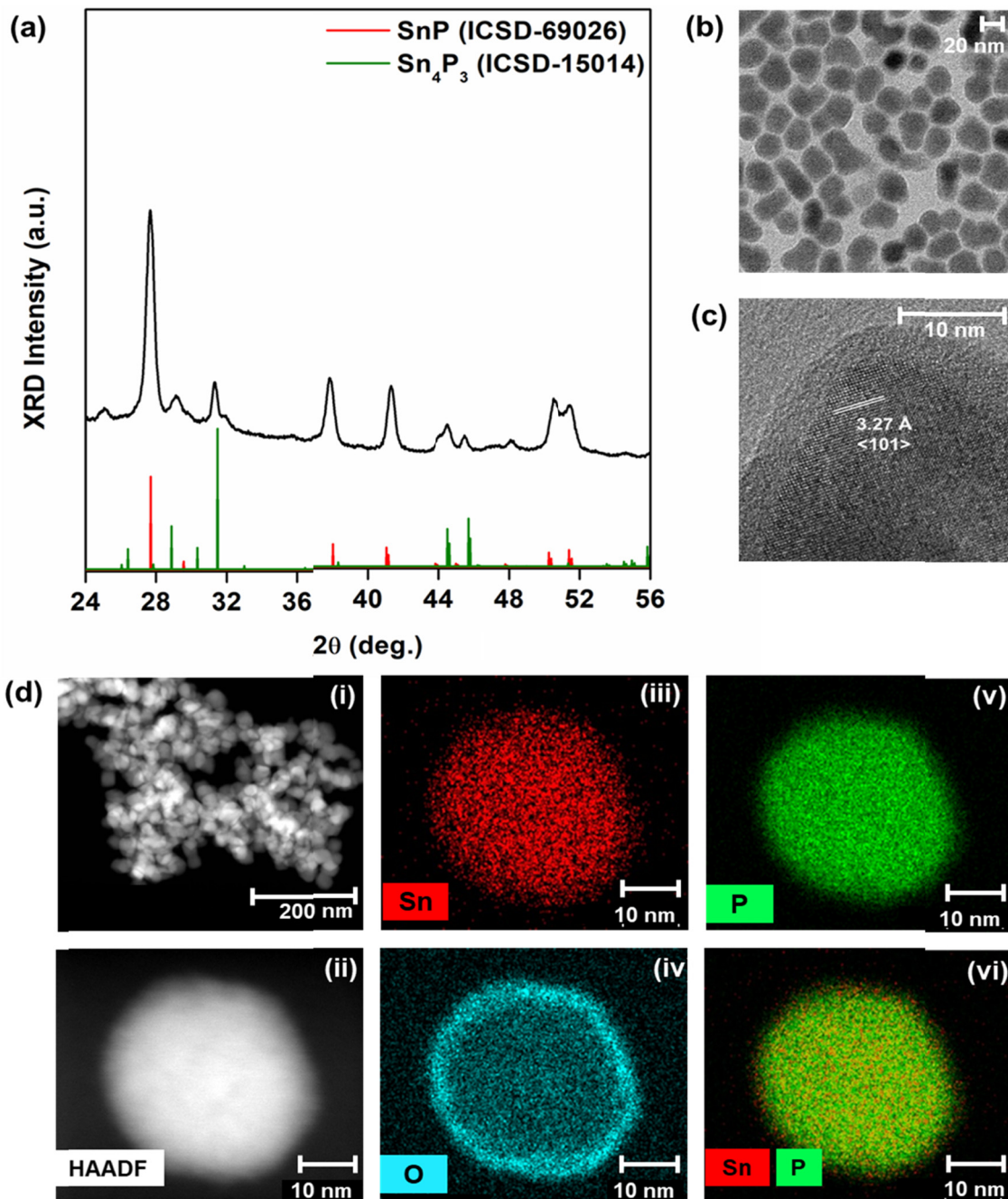
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**Fig. 1** Phase and morphology of tin phosphide nanocrystals. The nanoparticles were grown for 2 min following the injection of tris(diethyl)aminophosphine ((PNEt<sub>2</sub>)<sub>3</sub>) at 250 °C. (a) X-ray diffraction (XRD) patterns for the synthesized nanocrystals (black) were indexed to SnP (red, ICSD-69026), with trace amounts of Sn<sub>4</sub>P<sub>3</sub> present (green, ICSD-15014). (b) Bright field transmission electron microscopy (TEM) images of synthesized nanocrystals revealed that they were 25 ± 3.9 nm in diameter. (c) High resolution transmission electron microscopy (HRTEM) showed *d*-spacings of 3.27 Å, corresponding to the characteristic <101> peak of SnP. We observed oxidation at the surface of the nanocrystals due to exposure to air prior to the HRTEM measurement. (d) Scanning transmission electron microscopy (STEM) and high-angle annular dark field (HAADF) images for nanocrystals show elemental mapping for Sn, O, and P. The overlay of Sn and P maps provided an approximate Sn : P ratio of 1 : 1.

microscopy (TEM), X-ray photoelectron spectroscopy (XPS) and X-ray absorption spectroscopy (XAS). XRD and TEM are conventional techniques that provided the morphology and crystal structures of our samples. XPS and XAS allowed us to investigate oxidation states and corresponding local environments of tin in tin phosphides. Characterization of the

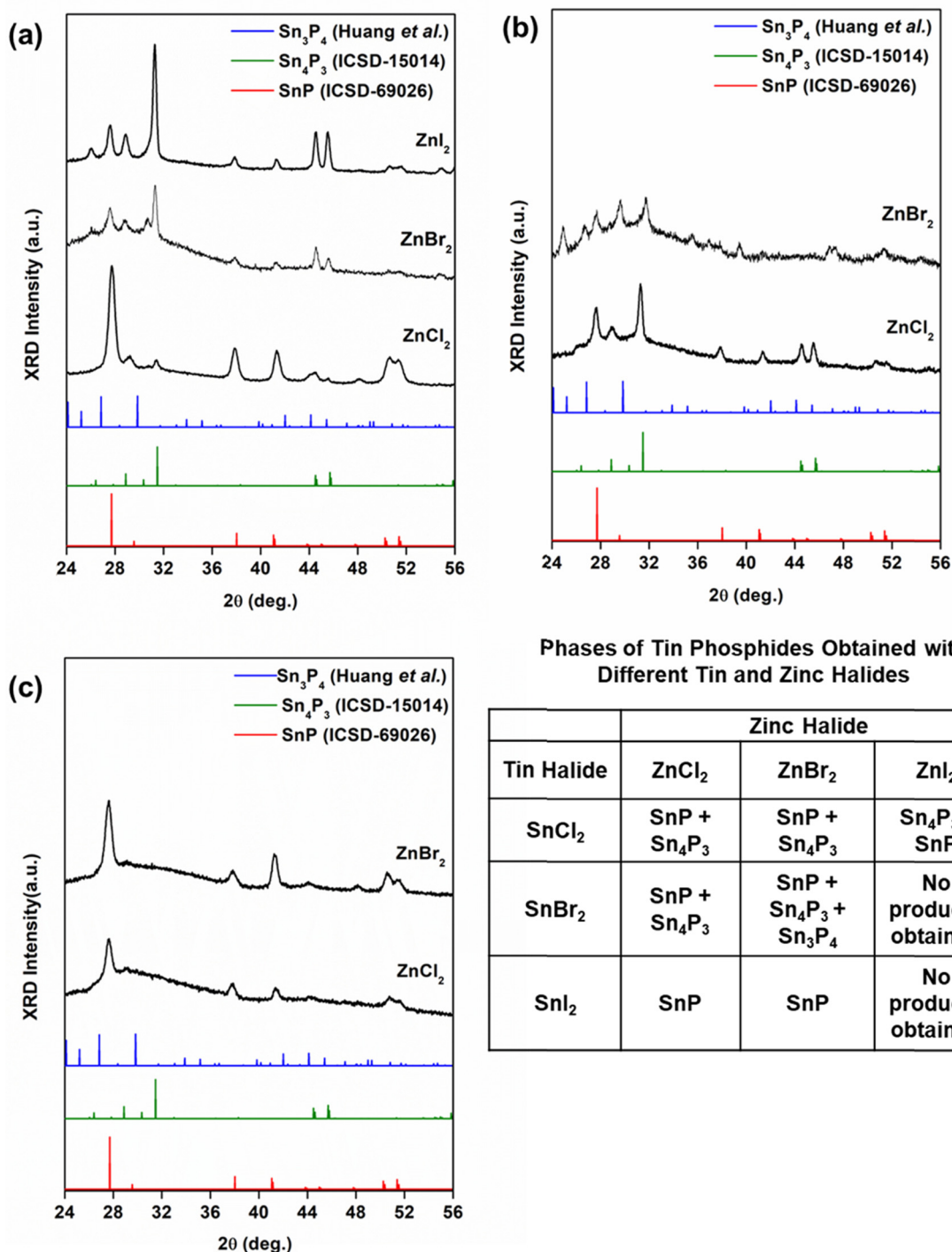
**Table 1** Phases of tin phosphides obtained with different tin and zinc halides

Tin halide	Zinc halide		
	ZnCl <sub>2</sub>	ZnBr <sub>2</sub>	ZnI <sub>2</sub>
SnCl <sub>2</sub>	SnP + Sn <sub>4</sub> P <sub>3</sub>	SnP + Sn <sub>4</sub> P <sub>3</sub>	Sn <sub>4</sub> P <sub>3</sub> + SnP
SnBr <sub>2</sub>	SnP + Sn <sub>4</sub> P <sub>3</sub>	SnP + Sn <sub>4</sub> P <sub>3</sub> + Sn <sub>3</sub> P <sub>4</sub>	No products obtained
SnI <sub>2</sub>	SnP	SnP	No products obtained



oxidation states of various tin phosphides remain a current topic of interest. Results have been mixed, with multiple oxidation states observed in both SnP and Sn<sub>4</sub>P<sub>3</sub>. Even less is

known about Sn<sub>3</sub>P<sub>4</sub>. Our characterization methodology aims to provide insight into the coordination environments of tin in nanostructured tin phosphides.



**Fig. 2** X-ray diffraction (XRD) patterns (obtained for tin phosphide nanocrystals grown with various combinations of SnX<sub>2</sub> and ZnX<sub>2</sub> (X = Cl<sup>-</sup>, Br<sup>-</sup>, and I<sup>-</sup>) precursors. Nanocrystals were grown for 2 min following injection of tris(diethyl)aminophosphine (P(NEt<sub>2</sub>)<sub>3</sub>) at 250 °C. Data is plotted against reference data for SnP (red, ICSD-69026), Sn<sub>4</sub>P<sub>3</sub> (green, ICSD-15014) and Sn<sub>3</sub>P<sub>4</sub> (blue, Huang *et al.*). In each experiment, a 1 : 1 ratio of Zn : Sn was used. (a) Holding SnCl<sub>2</sub> constant, phase mixtures of SnP and Sn<sub>4</sub>P<sub>3</sub> using ZnX<sub>2</sub>. (b) Similarly, holding SnBr<sub>2</sub> constant, phase mixtures of SnP, Sn<sub>4</sub>P<sub>3</sub>, and Sn<sub>3</sub>P<sub>4</sub> were obtained from experiments with ZnCl<sub>2</sub> and ZnBr<sub>2</sub>. Experiments with ZnI<sub>2</sub> were unsuccessful. (c) Holding SnI<sub>2</sub> constant, phase-pure SnP nanocrystals were obtained with ZnCl<sub>2</sub> and ZnBr<sub>2</sub>. Experiments with ZnI<sub>2</sub> were unsuccessful.



## 2. Materials and methods

### 2.1 Materials

Ethanol (anhydrous, 200 proof,  $\geq 99.5\%$ ), oleic acid (technical grade, 90%), oleylamine (technical grade, 70%), tin(II) bromide (anhydrous, 99.999% trace metal basis), tin(II) iodide (anhydrous, powder, 99.999%, trace metals basis), toluene (anhydrous, 99.8%), trioctylamine (98%), tris(diethylamino)phosphine (97%), trioctylphosphine (97%), zinc(II) bromide (anhydrous, 99.999% trace metal basis), zinc(II) chloride (anhydrous, 99.999% trace metal basis), and zinc(II) iodide (anhydrous, 99.999% trace metal basis), were purchased from Sigma Aldrich. Tin(II) chloride (anhydrous, 98%) was purchased from Alfa Aesar. All chemicals were used without further purification after their purchase. Anhydrous chemicals were purchased and stored in the glovebox. Extra care was used when handling tin halides and zinc halides, which are hygroscopic.

Please note that quenching of the following reactions requires injection of toluene at 90 °C. We encourage caution when replicating our work, as injection of toluene above its boiling point (110 °C) can cause harm.

### 2.2 Synthesis of trigonal Sn<sub>3</sub>P<sub>4</sub> nanocrystals

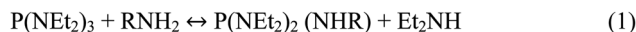
Tin(II) chloride (SnCl<sub>2</sub>) (256.0 mg, 1.35 mmol), zinc(II) chloride (ZnCl<sub>2</sub>) (184.0 mg, 1.35 mmol), oleylamine (20 mL), and oleic acid (1 mL) were mixed in a 100 mL three-neck flask. The reaction mixture was degassed under vacuum at 120 °C, producing a clear, yellow solution that was then heated under nitrogen to 250 °C. At 250 °C, P(NEt<sub>2</sub>)<sub>3</sub> (1.6 mL, 4 mmol) was rapidly injected into the flask, producing a black dispersion that was left for 2 min. After 2 min, the reaction was cooled slowly to room temperature. The reaction mixture was then transferred to a nitrogen-filled glovebox for extraction and cleaning.

In a typical purification process, Sn<sub>3</sub>P<sub>4</sub> nanocrystals were precipitated from ethanol *via* centrifugation at 6000 rpm for 5 min. The precipitated nanocrystals, a black powder, were then washed with toluene (5 mL) by centrifugation at 2000 rpm for 5 min to precipitate out extra surfactants and unreacted precursors. The supernatant was extracted, the dispersed particles were crashed out with ethanol, and finally the nanoparticles were dried under vacuum and stored as a dry powder in the glovebox.

### 2.3 Synthesis of trigonal SnP nanocrystals

SnCl<sub>2</sub> (256.0 mg, 1.35 mmol), ZnCl<sub>2</sub> (184.0 mg, 1.35 mmol), oleylamine (40 mL) and oleic acid (1 mL) were mixed in a 100 mL three-neck flask. The reaction mixture was degassed under vacuum at 120 °C, producing a clear, yellow solution that was then heated under nitrogen to 250 °C. At 250 °C, P(NEt<sub>2</sub>)<sub>3</sub> (1.6 mL, 4 mmol) was rapidly injected into the flask, producing a black dispersion that was left for 2 min. The reaction was then quenched in three steps. First, a heat gun on the “cool” air setting was used until the reaction temperature fell to 200 °C. Then, a room temperature water bath was introduced to further cool the reaction. Finally, toluene (20 mL) was injected at 90 °C to complete the quenching procedure.

The reaction mixture was transferred to a nitrogen-filled glovebox for purification and extraction. In a typical purification



Scheme 1 Transamination of tris(aminophosphines) with primary amines.

process, SnP nanocrystals were precipitated from ethanol *via* centrifugation at 6000 rpm for 5 min. The precipitated nanocrystals, a black powder, were then washed with toluene (5 mL) by centrifugation at 2000 rpm for 5 min to precipitate out extra surfactants and unreacted precursors. The supernatant was extracted, the dispersed particles were crashed out with ethanol, and finally the nanoparticles were dried under vacuum and stored as a dry powder in the glovebox.

### 2.4 Synthesis of rhombohedral Sn<sub>4</sub>P<sub>3</sub> nanocrystals

SnCl<sub>2</sub> (256.0 mg, 1.35 mmol), ZnCl<sub>2</sub> (184.0 mg, 1.35 mmol), oleylamine (15 mL), trioctylamine (5 mL), and oleic acid (1 mL) were mixed in a 100 mL three-neck flask. The reaction mixture was degassed under vacuum at 120 °C, producing a clear, yellow solution that was then heated under nitrogen to 250 °C. At 250 °C, P(NEt<sub>2</sub>)<sub>3</sub> (1.6 mL, 4 mmol) was rapidly injected into the flask, producing a black dispersion that was left for 2 min. The reaction was then quenched in three steps. First, a heat gun on the “cool” air setting was used until the reaction temperature fell to 200 °C. Then, a room temperature water bath was introduced to further cool the reaction. Finally, toluene (20 mL) was injected at 90 °C to complete the quenching procedure.

The reaction mixture was transferred to a nitrogen-filled glovebox for purification and extraction. In a typical purification process, Sn<sub>4</sub>P<sub>3</sub> nanocrystals were precipitated from ethanol *via* centrifugation at 6000 rpm for 5 min. The precipitated nanocrystals, a black powder, were then washed with chloroform (5 mL) by centrifugation at 2000 rpm for 5 min to precipitate out extra surfactants and unreacted precursors. The supernatant was extracted, and the dispersed particles were crashed out with ethanol. Finally, the nanoparticles were dried under vacuum and stored as a dry powder in the glovebox.

### 2.5 Characterization

TEM and XRD measurements were used to study the morphology and phase of the synthesized nanocrystals. Energy dispersive X-ray spectroscopy (EDX) determined relative stoichiometry. XPS and XAS measurements provided the chemical composition and local environment of our materials. Details of sample preparation,

Table 2 Phases of tin phosphides obtained with varying amounts of oleylamine

Volume (ml)	Oleylamine : P(NEt <sub>2</sub> ) <sub>3</sub>	Phase
10	7.5 : 1	Unidentified phase mixture
20	15 : 1	SnP + Sn <sub>4</sub> P <sub>3</sub>
40	30 : 1	SnP



experimental conditions, and analysis procedures can be found in the ESI.†

### 3. Results and discussion

#### 3.1 Synthesis of tin phosphide nanocrystals

$\text{Sn}_3\text{P}_4$ , SnP, or  $\text{Sn}_4\text{P}_3$  nanocrystals were synthesized *via* hot injection of  $\text{P}(\text{NET}_2)_3$  into a mixture of amine solvents, oleic acid, zinc halides, and tin halides (Table S1, ESI†). We introduced zinc halides into the reaction mixture following the example of InP syntheses, where zinc halide choice influenced particle size.<sup>45</sup> Zinc halides have been found to promote the formation and stabilization of activated precursors for growth

Table 3 Phases of tin phosphides obtained with addition of cosolvents

Cosolvent	Volume (mL)	Phase
Triethylamine	5	$\text{Sn}_4\text{P}_3$
	10	$\text{SnP} + \text{Sn}_4\text{P}_3 + \text{Sn}_3\text{P}_4$
	15	$\text{SnP} + \text{Sn}_4\text{P}_3 + \text{Sn}_3\text{P}_4$
Triethylphosphine	0.15	$\text{Sn}_3\text{P}_4 + \text{SnP}$
	0.6	$\text{Sn}_3\text{P}_4 + \text{SnP}$
	3	$\text{Sn}_3\text{P}_4 + \text{SnP}$

*via* Zn–N–P intermediates.<sup>49</sup> We thus began our work by investigating the role of zinc in aminophosphine-based routes to tin phosphides.

We began our study with  $\text{ZnCl}_2$  and  $\text{SnCl}_2$ , as dichlorides have been most used in previous aminophosphine-based

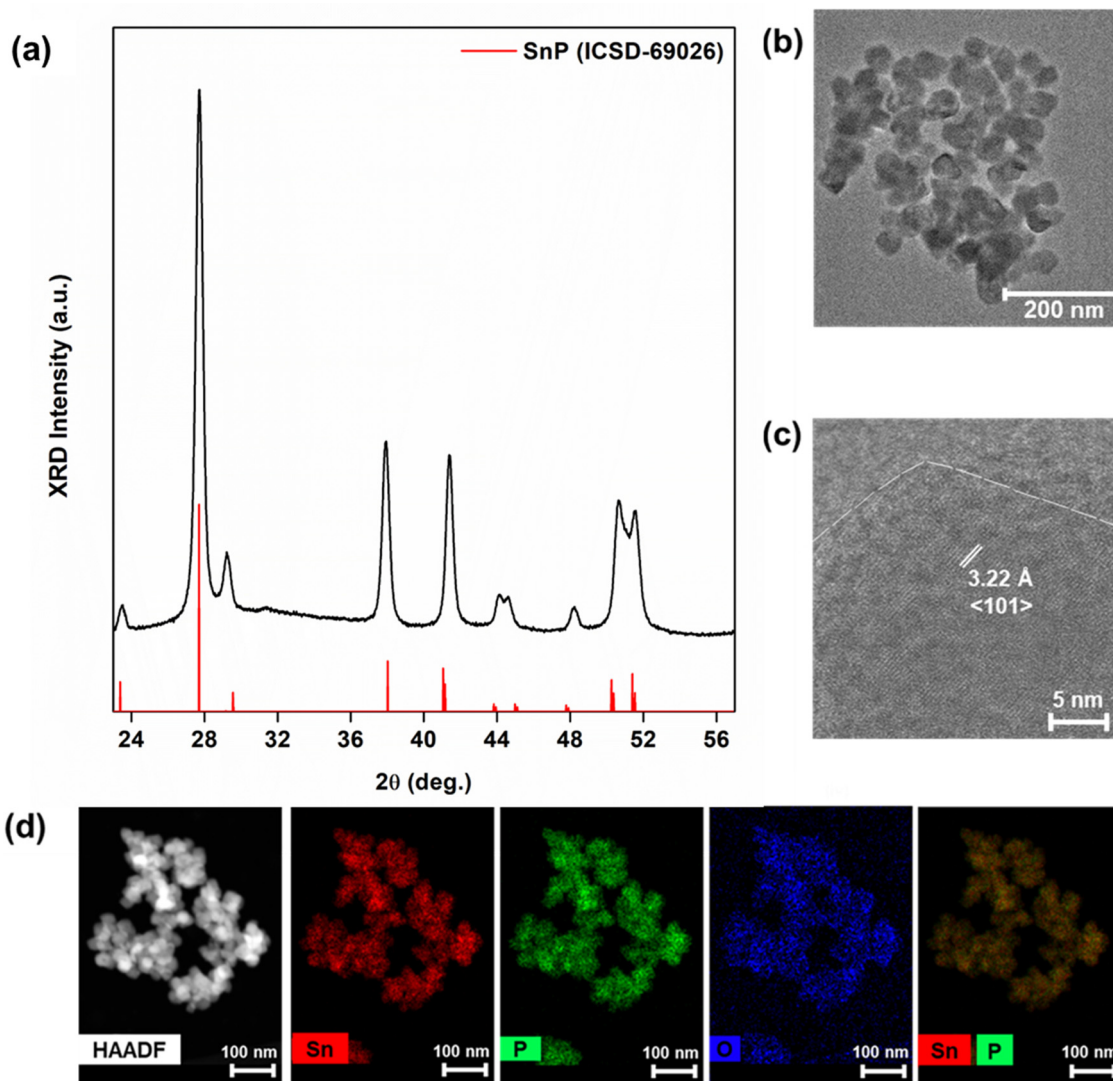


Fig. 3 Phase and morphology information for phase-pure SnP nanocrystals grown for 2 min at 250 °C following injection of tris(diethyl)aminophosphine ( $\text{PNET}_2$ ) into 40 mL of oleylamine and equimolar amounts of  $\text{SnCl}_2$  and  $\text{ZnCl}_2$ . (a) X-ray diffraction data (XRD) obtained for synthesized trigonal SnP nanocrystals plotted against reference data for trigonal SnP (ICSD-69026). (b) Bright field transmission electron microscopy (TEM) revealed that the SnP nanocrystals were  $40 \pm 7.0$  nm in diameter. (c) High resolution TEM (HRTEM) images showed crystalline particles with an amorphous layer on its surface. (d) Scanning transmission electron microscopy High-angle annular dark-field (STEM-HAADF) image and corresponding elemental mapping for Sn, O, and P. The overlay approximated an Sn:P ratio of 1:1.



routes to metal phosphides.<sup>42–48</sup> XRD data were indexed to trigonal SnP with rhombohedral Sn<sub>4</sub>P<sub>3</sub> as an impurity phase (Fig. 1a). We note that in the bulk, SnP is a metastable phase formed only at high temperature and high pressure, while Sn<sub>4</sub>P<sub>3</sub> is the thermally stable phase of tin phosphide.<sup>50</sup> Nanocrystals often stabilize in metastable phases because of their large surface energies.<sup>51</sup> While our data were insufficient to estimate the amount of each phase present, Sn<sub>4</sub>P<sub>3</sub> impurities in SnP have been previously reported.<sup>39</sup>

Bright field TEM images showed that the average diameter of the nanoparticles was  $25 \pm 3.9$  nm (Fig. 1b). Under closer inspection in HRTEM, we observed a *d*-spacing of 3.27 Å, indexed to the characteristic (101) peak of trigonal SnP (Fig. 1c). Assignment of SnP to the structure was consistent with EDX measurements, which approximated an Sn : P ratio of 1 : 1 (Fig. 1d). In HRTEM and EDX, we also observed an amorphous layer of oxidized tin at the surface of the nanocrystals (Fig. 1c and d). We note that no XRD peaks were indexed to tin oxides or tin phosphates in samples that were freshly prepared for use in diffraction studies. Still, metal phosphides readily oxidize. While exposure to air was minimized, sample oxidation may have occurred during transportation to the microscopy facility (from Brooklyn, NY to Berkeley, CA) or during sample loading.

Phase mixtures were previously observed in cobalt phosphides obtained from aminophosphines and cobalt dihalides.<sup>46</sup> Thus, to further investigate these results, we synthesized tin phosphides from several combinations of tin and zinc dihalides (Table 1). Experiments using SnCl<sub>2</sub> and SnBr<sub>2</sub> provided phase mixtures of

SnP, Sn<sub>4</sub>P<sub>3</sub>, and Sn<sub>3</sub>P<sub>4</sub> (Fig. 2a and b). Using SnI<sub>2</sub>, we obtained phase-pure SnP (Fig. 2c).

We attributed our results to the relative dissociation energies of the halides (M–I < M–Br < M–Cl). Based on the dissociation energies of the halides, Sn–P monomers would form most easily from iodides, followed by bromides, then chlorides. Our results obtained from reactions combining SnI<sub>2</sub> with ZnCl<sub>2</sub> or ZnBr<sub>2</sub> supported this reasoning. The reaction occurred as expected, reducing SnI<sub>2</sub> to produce phase-pure SnP. However, combining SnI<sub>2</sub> with ZnI<sub>2</sub> was unsuccessful. In this instance, we noticed a dangerous over pressurization of the reaction flask upon injection of P(NEt<sub>2</sub>)<sub>3</sub>. Attempts to reduce the pressure did not improve the result, and so these experiments were forgone. The over pressurization may have occurred due to the formation of diethylamine made possible by the dissociation of M–I complexes.

Still, the results obtained from SnCl<sub>2</sub> and SnBr<sub>2</sub> were more complicated. We obtained phase mixtures of tin phosphides, with the dominant phase changing depending on the halide used. We note that the role of zinc in these reactions is still under investigation. In all experiments, the ratio of zinc to tin halide (Zn : Sn) was held constant at 1 : 1. Maintaining a 1 : 1

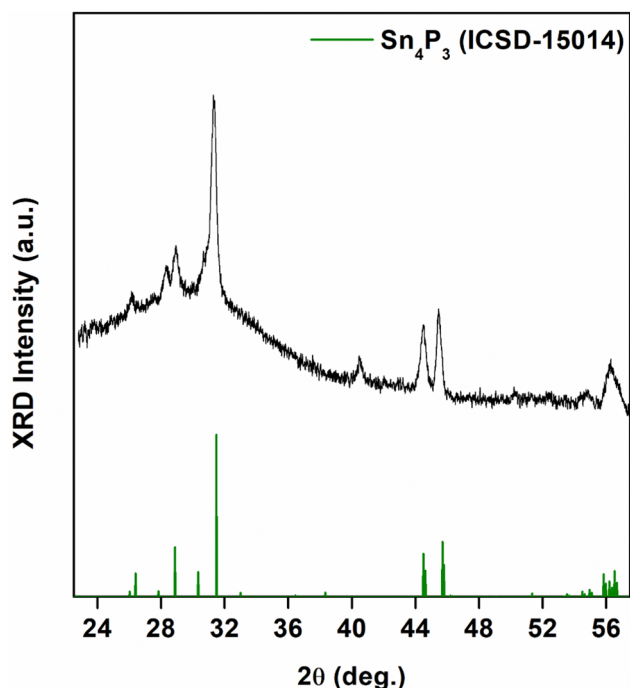


Fig. 4 X-ray diffraction data obtained for synthesized Sn<sub>4</sub>P<sub>3</sub> nanocrystals plotted against reference data for rhombohedral Sn<sub>4</sub>P<sub>3</sub> (ICSD-15014). Nanocrystals were grown for 2 min at 250 °C following injection of tris(diethyl)aminophosphine (P(NEt<sub>2</sub>)<sub>3</sub>) into a 25/75 (v/v) mixture of triocetylamine to oleylamine.

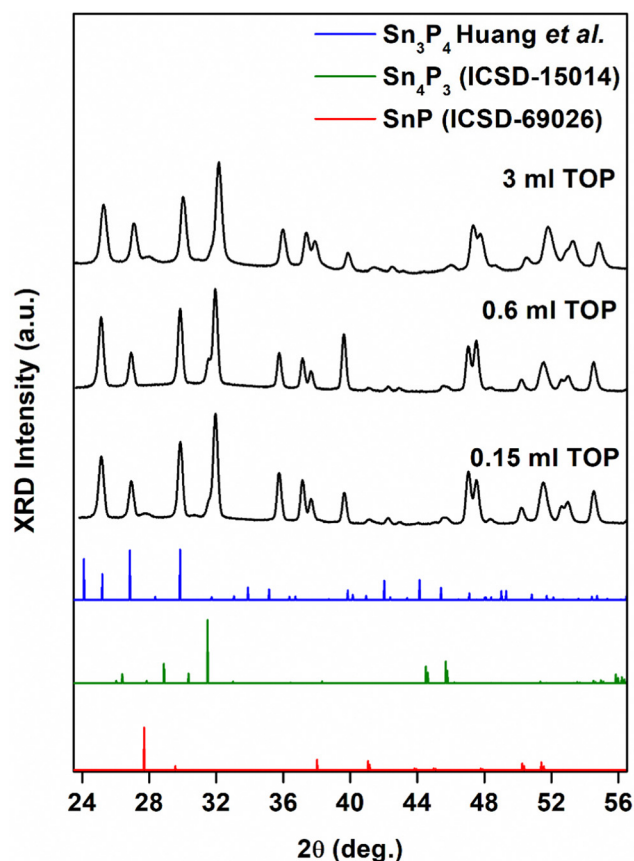


Fig. 5 X-ray diffraction (XRD) data for Sn<sub>3</sub>P<sub>4</sub> nanocrystals grown for 2 min at 250 °C following injection of tris(diethyl)aminophosphine ((PNEt<sub>2</sub>)<sub>3</sub>) in the presence of 0.15 mL, 0.6 mL, and 3 mL TOP. The data was compared to trigonal SnP (ICSD-69026), rhombohedral Sn<sub>4</sub>P<sub>3</sub> (ICSD-15014), and trigonal Sn<sub>3</sub>P<sub>4</sub> (Huang *et al.*<sup>56</sup>).



ratio of Zn:Sn was essential to obtain identifiable products. When we altered the Zn:Sn ratio or removed zinc from the reaction, we could not obtain any phase-pure materials (Fig. S1, ESI<sup>†</sup>). As mentioned previously, zinc halides form Zn–N–P intermediates that promote the formation activated precursors for growth. These intermediates can undergo competing processes that can modify reaction equilibria.<sup>49</sup>

Previous work has shown that reaction kinetics vary with the ratio of P(NEt<sub>2</sub>)<sub>3</sub> to primary amine solvent. This is because the amine plays two roles in the reaction; the solvent activates P(NEt<sub>2</sub>)<sub>3</sub> and complexes the metal halide. In its first role, activation occurs *via* transamination (Scheme 1).

The reaction is driven by the evaporation of diethylamine (Et<sub>2</sub>NH, with a boiling point of 7 °C) at reaction temperature. As the dialkylamine evaporates, the equilibrium shifts the reaction towards production of the final transamination product, P(NHR)<sub>3</sub>. Buffard *et al.* found that ratio of P(NEt<sub>2</sub>)<sub>3</sub> to oleylamine changed the transamination products present.<sup>44</sup> They identified a minimum amount of oleylamine required to produce quality nanocrystals, as we have observed in our work (Table 2).

We investigated the kinetics of this reaction first by tuning solvent volume. We hypothesized that changing the ratio of oleylamine to P(NEt<sub>2</sub>)<sub>3</sub> would drive the reaction towards phase-pure SnP or Sn<sub>4</sub>P<sub>3</sub>. With more oleylamine present to activate P(NEt<sub>2</sub>)<sub>3</sub>, we expected to form of phase-pure SnP; with less, we expected to form of Sn-rich Sn<sub>4</sub>P<sub>3</sub>. In our work, doubling the amount of oleylamine from a 15:1 ratio of oleylamine to P(NEt<sub>2</sub>)<sub>3</sub> to a 30:1 ratio provided phase-pure, trigonal SnP nanocrystals (Fig. 3a). The particles were 40 ± 7.0 nm in size,

with the observed *d*-spacing consistent with that of the ⟨101⟩ peak of SnP (Fig. 3b and c). STEM-EDX measurements confirmed a stoichiometry of 1:1 Sn:P (Fig. 3d). Halving the ratio, however, provided a complicated phase mixture inseparable from remaining reactants in the solution (Fig. S2, ESI<sup>†</sup>). The role of Zn–N–P intermediates in our reaction may have become more influential at this scale, changing disproportionation reactions. Buffard *et al.* and Mundy *et al.* saw similar in their work with other phosphide systems.<sup>44,46</sup>

To confirm the validity of this trend, we performed a control experiment where transamination and nucleation were separated. The “two-pot” protocol was adapted from work by Rachkov *et al.*<sup>47</sup> Oleylamine and P(NEt<sub>2</sub>)<sub>3</sub> were reacted under vacuum for 3 hours, and the resulting transamination product was then injected into the mixture of zinc and tin halides, oleylamine, and oleic acid at 250 °C. In the two-pot case, the results followed the trend observed in the one-pot experiment; increasing the amount of oleylamine present drove the reaction pathway towards SnP, and away from a phase mixture of SnP and Sn<sub>4</sub>P<sub>3</sub> (Fig. S3, ESI<sup>†</sup>).

The SnP nanocrystals exhibited poor stability (Fig. 3b). While the particles were visibly colloidal stable, aggregation still occurred (Fig. S4, ESI<sup>†</sup>). We attempted to stabilize the particles using the co-solvents trioctylphosphine (TOP) and trioctylamine (Table 3). Trioctylamine was previously used to stabilize InP nanocrystals synthesized from aminophosphines.<sup>44</sup> TOP was previously used to stabilize tin phosphide nanocrystals.<sup>39</sup>

We found that a mixture of trioctylamine to oleylamine shifted the reaction pathway away from SnP to other tin

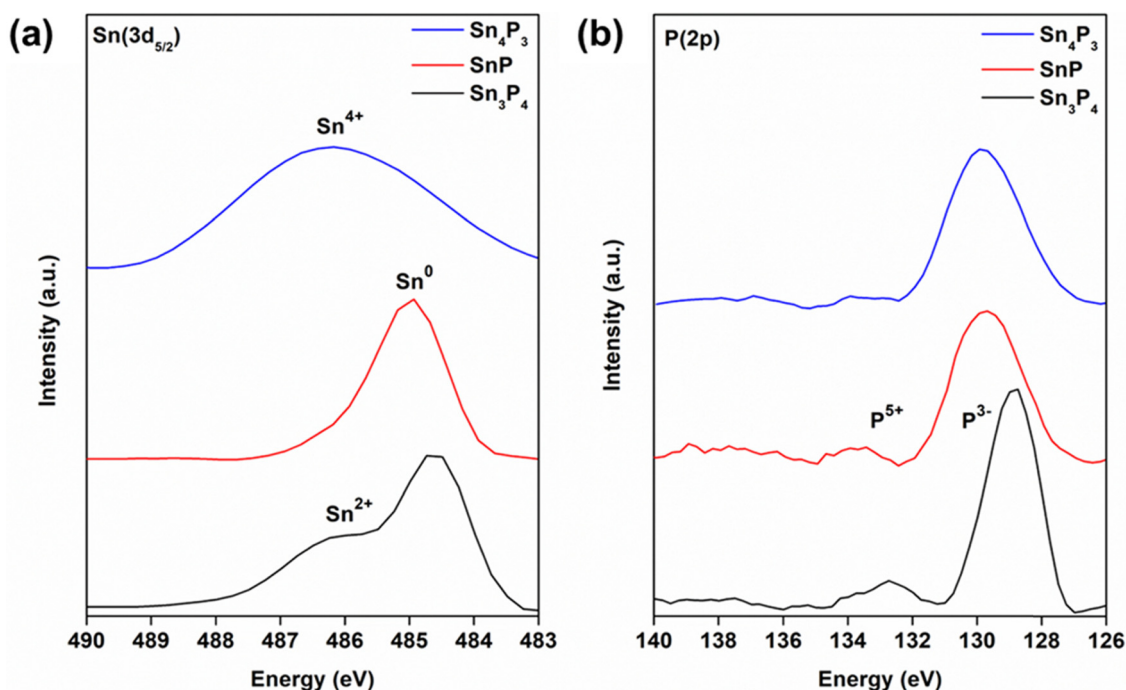


Fig. 6 X-ray photoelectron spectroscopy (XPS) spectra observed at the (a) Sn(3d<sub>5/2</sub>) and (b) P(2p<sub>3/2</sub>) regions for Sn<sub>3</sub>P<sub>4</sub> (bottom, black), SnP (middle, red), Sn<sub>4</sub>P<sub>3</sub> (top, blue) nanocrystals. Sn(3d<sub>5/2</sub>) binding energies revealed the presence of mixed oxidation states in all tin phosphides, which displayed binding energies from 484.6 eV to 486.2 eV.



phosphide phases. With 25 vol% of oleylamine replaced with trioctylamine, the effective ratio of amine to  $P(\text{NEt}_2)_3$  was reduced, producing less  $P(\text{NHR})_3$  and shifting the direction of the reaction towards Sn-rich  $\text{Sn}_4\text{P}_3$  (Fig. 4). An unidentified minor impurity phase was also observed. Scherrer analysis approximated that the  $\text{Sn}_4\text{P}_3$  nanoparticles were approximately 20.2 nm in diameter (ESI†, Section 1.1.2); size could not be confirmed *via* TEM as the particles could not be completely isolated. With increased amounts of trioctylamine, we obtained phase mixtures of  $\text{Sn}_3\text{P}_4$ , SnP, and  $\text{Sn}_4\text{P}_3$  (Fig. S5, ESI†). We attributed the change in reaction pathway to shifts in the equilibrium of transamination depending on the concentration of the amine consumed. If only oleylamine participates in transamination, the reaction occurs between  $P(\text{NEt}_2)_3$  and  $\text{RNH}_2$  to form  $P(\text{NHR})_3$  (R = methyl or ethyl group), which forms a dative bond with  $\text{SnX}_2$ .

When we performed reactions with TOP, we obtained  $\text{Sn}_3\text{P}_4$  nanocrystals with an SnP impurity phase (Fig. 5). We obtained this mixture across the range of temperatures of experiments performed (Fig. S6, ESI†). The successful synthesis of bulk  $\text{Sn}_3\text{P}_4$  has required a high vapor pressure of phosphorus to provide phase-pure crystals.<sup>52</sup> We suspect that the presence of phosphorus byproducts formed during transamination may have analogously elevated the phosphorus activity. Furthermore, colloidal synthesis provides routes to phases unattainable in the bulk.<sup>53</sup> The XRD pattern was comparable to a structure predicted by Huang *et al.*<sup>38</sup> Nanocrystals were  $8.3 \pm 2.8$  nm in diameter. EDX measurements demonstrated an Sn:P ratio of  $0.8 \pm 0.2$ , approximately  $\text{Sn}_3\text{P}_4$  (Fig. S7, ESI†). To the best of our knowledge, this is the first report of crystalline  $\text{Sn}_3\text{P}_4$  nanoparticles.

### 3.2 Chemical characterization of tin phosphide nanocrystals

Understanding the local environment of tin is necessary to understand its use in various applications. The presence of several tin oxidation states ( $\text{Sn}^0$ ,  $\text{Sn}^{2+}$ , and  $\text{Sn}^{4+}$ ) within the same crystal structure has hindered studies to date.<sup>1</sup> We use XPS and XAS to provide insight into the chemical environment of tin in SnP,  $\text{Sn}_3\text{P}_4$ , and  $\text{Sn}_4\text{P}_3$ .

Through XPS, we measured binding energies of  $\text{Sn}(3d_{3/2})$  and  $\text{P}(2p_{3/2})$  in our nanocrystals (Fig. 6 and Fig. S8–S10, ESI†). The experimental  $\text{Sn}(3d_{3/2})$  binding energies were consistent with reported results for tin phosphide nanocrystals.<sup>34</sup> For  $\text{Sn}_3\text{P}_4$ , we observed two peaks, at 484.6 eV and 486.2 eV. We attributed the peak at 484.6 eV to metallic Sn and the peak at 486.2 eV to  $\text{Sn}^{2+}$ .<sup>39,54</sup> For SnP, we observed a peak at 485 eV, suggesting metallic Sn bonding as previously described.<sup>55</sup> For  $\text{Sn}_4\text{P}_3$ , we observed a broad peak at 486.2 eV, suggesting the presence of multiple oxidation states. To date, formal charges for Sn in rhombohedral  $\text{Sn}_4\text{P}_3$  have not been assigned. Evidence of metallic bonding like that of  $\beta$ -Sn has been observed, while XAS data have suggested Sn–P and Sn–P–Sn bonding structure.<sup>10,56,57</sup> Mixed oxidation states may have also been present due to surface oxidation, as observed in TEM. Evidence of oxidation was also observed in experimental  $\text{P}(2p_{3/2})$  binding energies, which showed a peak at 133 eV, typical of metal phosphates. Binding energies below 130 eV, characteristic to metal phosphides, were dominant; the binding energy

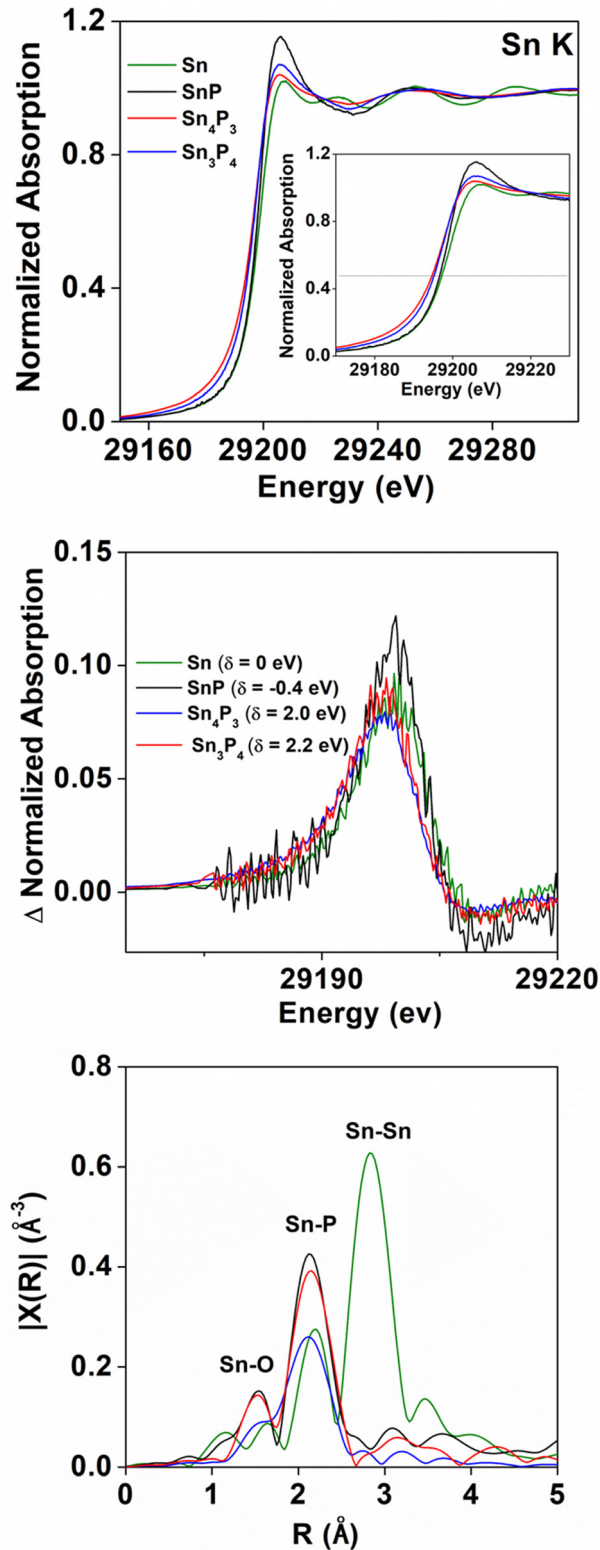
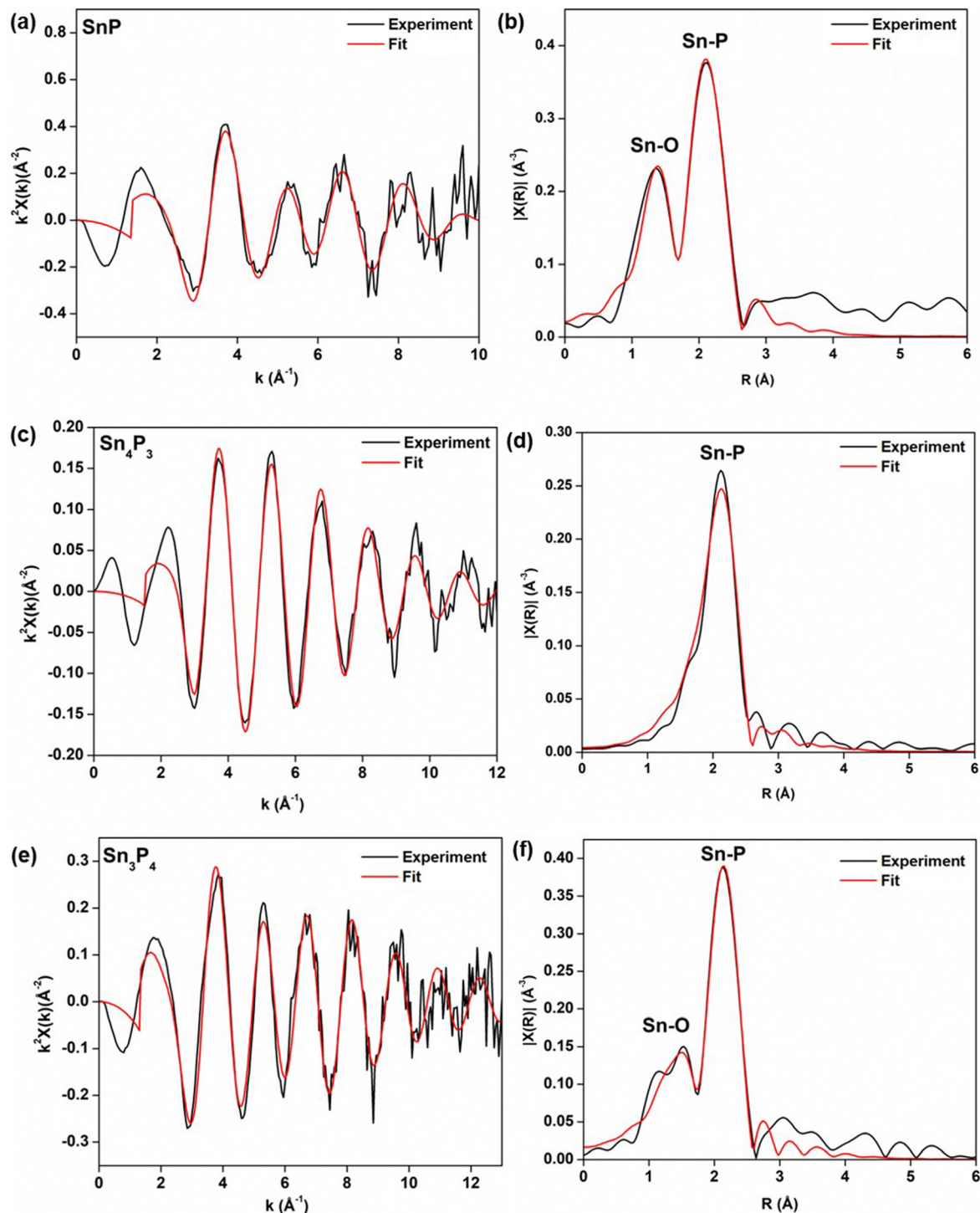


Fig. 7 (a) X-ray absorption near edge structure (XANES) spectra at the Sn K-edge for  $\text{Sn}_3\text{P}_4$  (red),  $\text{Sn}_4\text{P}_3$  (blue), and SnP (black) nanocrystals. The inset shows the normalized absorption spectrum. Data is plotted against Sn foil ( $\text{Sn}^0$ , green). (b) The first derivative of the XANES spectra with relative shifts ( $\delta$ ) from the first derivative point of  $\text{Sn}^0$  and (c) Fourier transform magnitude  $k^2$ -weighted EXAFS spectra (FT-EXAFS). Two first-shell pathways were identified in tin phosphides: Sn–O (1.4 Å) and Sn–P (2.1 Å). Note that peak positions in and the actual bond lengths differ by ca. 0.5 Å because of the photoelectron phase shift. The bond lengths for tin phosphide materials are approximately 2.6 Å, consistent with previous reports.







**Fig. 8** Summary of EXAFS fitting for (a and b) SnP, (c and d)  $\text{Sn}_4\text{P}_3$ , and (e and f)  $\text{Sn}_3\text{P}_4$  shown in both  $k$  and  $r$  space. SnP data were fit with an  $r$  range of 1.1 to 2.8 Å and  $k$  range of 2.0 to 9.0 Å<sup>-1</sup>. Sn-O and Sn-P photoelectron paths were used in the fit to SnP.  $\text{Sn}_4\text{P}_3$  data were fit with an  $r$  range of 1.45 to 3 Å and  $k$  range of 2.3 to 11.45 Å<sup>-1</sup>. Sn-P photoelectron path was used in the fit to  $\text{Sn}_4\text{P}_3$ .  $\text{Sn}_3\text{P}_4$  fits were performed using a  $k$  range of 2.5 to 10.5 Å<sup>-1</sup> and an  $r$  range of 1.1 to 2.7 Å. Sn-O and Sn-P photoelectron paths were used in the fit for  $\text{Sn}_3\text{P}_4$ .

slightly decreased with increasing phosphorus content. While this trend has been observed in other metal phosphides, the exact nature of these shifts has not been determined.<sup>58</sup>

Formal charge(s) of tin in tin phosphides remain under investigation. However,  $\text{Sn}^{2+}$  and  $\text{Sn}^{4+}$  species present in  $\text{Sn}_3\text{P}_4$

and  $\text{Sn}_3\text{P}_4$  may have also appeared due to surface oxidation;  $\text{P}(2p_{3/2})$  binding energies of 133 eV are characteristic of metal phosphates. Still, metal phosphide bonding, characteristic of binding energies below 130 eV, were observed. XAS measurements at the Sn K-edge further showed the complexity of the



local environment around Sn (Fig. 7a). SnP showed metallic behavior as expected; the very small electronegativity difference between tin and phosphorus causes delocalization of the electrons and allows for the layered structure of SnP.<sup>55</sup> Closer inspection of the X-ray absorption near edge structure (XANES) region in derivative plot further showed this small shift in ionization energy for Sn<sub>4</sub>P<sub>3</sub> and Sn<sub>3</sub>P<sub>4</sub>, unexpectedly to binding energies below that of metallic tin, represented by a reference foil (Fig. 7b). Back bonding from phosphorus to metals could be a possible explanation for this behavior.<sup>58</sup> The Sn K-edge XANES spectra reflects the transition energy from the 1s state to the 5p state. If back donation occurred, it could result in an increased shielding effect that would support our observed red shift. However, our XPS data contradict this hypothesis, as back donation implies a positive charge on phosphorus. Further investigation is ongoing.

Despite unusual XANES behavior, the Fourier Transform of the  $k^2$ -weighted Extended X-ray absorption fine structure (EXAFS) data (Fig. 7c) were consistent with Sn–P bonding expected for tin phosphides. The position of the first shell peaks at approximately 2.1 Å for Sn<sub>4</sub>P<sub>3</sub>, SnP, and Sn<sub>3</sub>P<sub>4</sub> were attributed to the Sn–P photoelectron path. Note that peak positions in Fig. 7c and the actual bond lengths differ by *ca.* 0.5 Å because of the photoelectron phase shift. The actual observed bond lengths for our materials were therefore approximately 2.6 Å, consistent with reported Sn–P bond lengths.<sup>5</sup> A small feature around 1.4 Å that was observed in SnP and Sn<sub>3</sub>P<sub>4</sub> was attributed to the Sn–O photoelectron path.

For SnP and Sn<sub>4</sub>P<sub>3</sub>, fitting was performed using their bulk structures. For Sn<sub>3</sub>P<sub>4</sub>, a survey of pathways was explored; though XRD data showed a similar structure to that predicted by Huang *et al.*, some peaks remain unindexed. We thus investigated several models to gain insight to local structure in Sn<sub>3</sub>P<sub>4</sub> (Table S1, ESI†). Aside from Sn–P, the scattering contributions of Sn–O were also included in some cases, as oxidation was observed in TEM measurements.

A summary of the fits can be found in Fig. 8. For SnP, the trigonal phase provided an *R*-factor of 0.4% (Table S2 and Fig. 8a, b, ESI†). The agreement with the trigonal structure was expected as the XRD data for synthesized SnP was indexed to the same phase. The calculated Sn–P coordination number and bond length, however, were larger than expected; we obtained an Sn–P bond length of  $2.69 \pm 0.13$  Å, and a coordination number *N* of  $2.9 \pm 0.7$ . The structure model had a total of three Sn–P bonds: two shorter bonds and one longer bond with a 0.52 Å difference (*i.e.*, 1.96 Å and 2.52 Å). The influence of *k*-weighting on the expected Sn–X coordination numbers, *N*, was investigated and the results indicate overall consistency within the obtained values for *N* (Table S3, ESI†).

Similarly, for Sn<sub>4</sub>P<sub>3</sub> the bulk structure model was rhombohedral, as indexing of the XRD directed (Table S4 and Fig. 8c, d, ESI†). Even though the expected bond lengths were consistent with predicted Sn–P bonds; the apparent change in coordination number from *N* = 3 to *N* = 1 could be attributed to the decrease in cluster size, but a full understanding of structure would require modeling of higher-order shells unobserved in

our nanoparticles.<sup>59</sup> We hypothesize that these differences may come from Sn–Sn bonding from interlayers of SnP (excluded in our model) but have been previously observed in EXAFS.<sup>34</sup>

For Sn<sub>3</sub>P<sub>4</sub>, we expected the structure to match most closely that of trigonal Sn<sub>3</sub>P<sub>4</sub> provided by Huang *et al.*<sup>38</sup> However, the tetragonal phase of SnP provided the best fit with an *R*-factor of 0.3% (Table S5 and Fig. 8e, f, ESI†). While our XRD data was insufficient to provide a weight fraction of SnP present, the data suggest that local structure may be closer to that of SnP. Further investigation of the structure is thus needed. We note that to date, crystalline nanoparticles of Sn<sub>3</sub>P<sub>4</sub> have not yet been reported. Even in the bulk, the structure of Sn<sub>3</sub>P<sub>4</sub> remains unresolved; several phases have been predicted to exist in literature.

While the structures of tin phosphides remain to be solved, our measurements provided insight on the effect of crystallite size and phase on the local coordination in the synthesized tin phosphides. Further understanding of how particle size influences the local environments of Sn in layered tin phosphides will be critical to provide understanding of how intercalation processes in electrochemical systems using these tin phosphides are affected by particle size.

## 4. Conclusions

To the best of our knowledge, we report the first use of aminophosphines to achieve significant phase flexibility in tin phosphide nanocrystals and obtain Sn<sub>3</sub>P<sub>4</sub>, SnP, and Sn<sub>4</sub>P<sub>3</sub>. Understanding the mechanisms responsible for each phase requires further investigation. X-ray spectroscopy measurements provided insight on the local chemical environments of tin, of interest for further study of tin phosphides in applications such as electrochemical reactions.

## Author contributions

All authors contributed to the preparation of the manuscript. A. Sahu conceived and supervised the project. R. Y. and S. C. performed all preliminary syntheses and characterization. I. J. P., S. L., and H. X. optimized syntheses protocols and performed XRD measurements. I. J. P. and S. L. performed bright field TEM imaging. I. J. P., A. M. E., and M. K. performed XAS measurements. A. I. F. supervised XAS measurements and data analysis. I. J. P. and A. M. P. analyzed XRD data for Sn<sub>3</sub>P<sub>4</sub>. S. H. performed STEM imaging and elemental analysis at BNL. A. Singh performed STEM imaging and elemental analysis at LBNL. I. J. P. and A. Sahu analyzed all the data.

## Conflicts of interest

The authors declare no competing financial interest.



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