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Synthesis of highly monodisperse porous manganese oxide spheres using a butyric acid microemulsion

Stanton Ching,* Ian J. Richter, Kathryn A. Tutunjian,* David A. Kriz,*b and Yumi Kovic*a

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Porous manganese oxide spheres form readily from a reaction between KMnO₄ and n-butanol in aqueous butyric acid under ambient conditions. Spheres have uniform size, tuneable diameters and high surface areas. The material is an active catalyst for the oxidation of isopropanol to acetone.

Porous manganese oxides continue to attract considerable interest as promising materials for heterogeneous catalysis, battery and electrochemical applications, and toxic waste remediation. Many crystalline and amorphous forms are known, including those with spherical and hollow spherical structures comprised of aggregated nanoparticles. Such hierarchical morphologies often result in high surface areas that are desirable in applications where interfacial contact is critical.

We previously described the use of butyric acid (BA) as an easily removed soft template to prepare manganese oxide hollow spheres with high surface areas. In aqueous reactions between KMnO₄ and MnSO₄, BA serves as a capping ligand that suppresses manganese oxide particle growth on the nanoscale, while simultaneously forming a microemulsion that enables the resulting nanoparticles to aggregate into hollow spheres. Herein we report further research on BA as a template for hierarchical manganese oxide. Its presence in reactions between KMnO₄ and n-butanol (BuOH) results in highly uniform and porous (but not hollow) spheres. This facile, one-pot reaction occurs under ambient conditions.

Porous manganese oxide spheres form as a brown precipitate within 20 minutes of adding 25 mL of aqueous 0.040 M KMnO₄ to a 25 mL aqueous mixture of 0.48 M BuOH and 1.0 M BA. The reaction is carried out at room temperature under atmospheric pressure, S1. About 80 mg is typically obtained after filtration, washing, and drying at 110 °C. The material is amorphous based on powder XRD, S2. A formula of Kₐ₀.₀₂₇MnO₁₈₈·0.₆₁H₂O was determined by AA (for K and Mn) and TGA (for waters of hydration). From this formula, the isolated yield is about 80% based on Mn. Thiosulfate/iodide redox titration was used to determine an average Mn oxidation state of 3.₅₉. FT-IR shows no evidence of residual BA.

SEM and TEM images reveal highly monodisperse spheres, with average diameters measuring 3₇₉±1₅ nm, Fig. 1. TEM reveals spheres consisting of nanoplatelets, as seen on surfaces in Figure 1B. The distribution of sphere diameters is shown in Fig. 2. The hierarchical structure and morphology are consistent with the high surface area of 2₄₃ m²/g measured by BET analysis. Nitrogen adsorption/desorption shows type-IV isotherm behavior, characteristic of a mesoporous material, S3. Pore size distribution calculated by the BJH method revealed three distinct meso-range distributions, with a major peak at 4.₈₈ nm, followed by peaks at 3.₈₃ nm and 2.₁₈ nm, S4. HRTEM images reveal mesoporosity, as well as short range order that is typical for amorphous manganese oxides, Fig. 3. Platelet size is about 10 nm. Control reactions without BA result in loss of the spherical morphology and formation of larger aggregated particles, Fig. 4. Surface area for the control is 1₃₉ m²/g.

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Fig. 1. (A) SEM and (B) TEM images of porous manganese oxide spheres from KMnO₄ and BuOH in the presence of BA (S1).

Fig. 2. Size distribution for porous manganese oxide spheres (S1) with average diameter of 379 nm.

Fig. 3. HRTEM images of a porous manganese oxide sphere (S1) showing (A) mesoporosity and (B) short range order in the nanoplatelets.
Sphere size is responsive to KMnO₄ concentration, which allows the tuning of sphere diameters between 267 and 633 nm, Fig. 5. High surface areas of 176-256 m²/g are maintained for the different sphere sizes, S5. Larger diameters can be obtained with higher KMnO₄ concentrations, but there is considerable loss of monodispersity, S6. Next, the reaction of KMnO₄ and BuOH without BA (S1 without BA) proposes that BA coordinates weakly enough so that molecule-size clusters are uninhibited from growing into larger material-size particles. Weak coordination also allows easy removal of BA by washing. However, BA coordinates strongly enough to cap manganese oxide growth on the nanoparticle scale. The nanoparticles then aggregate to form a hierarchical structure. Coordinating solvents have been reported to function as ligands by coordinated acetate ligands.

This research grows out of our broader efforts to develop synthetic routes to nanoparticle manganese oxides having high surface areas. Our work is inspired by the chemistry of [Mn₂O₃(C₄H₉O)₆(H₂O)₄] clusters and related molecular systems. The clusters have cores of MnO₆ octahedral units, similar to those of manganese oxide materials, but are stabilized by coordinated acetate ligands. In our nanoparticle system, we propose that BA coordinates weakly enough so that molecule-size clusters are uninhibited from growing into larger material-size particles. Weak coordination also allows easy removal of BA by washing. However, BA coordinates strongly enough to cap manganese oxide growth on the nanoparticle scale. The nanoparticles then aggregate to form a hierarchical structure. Coordinating solvents have been reported to function as ligands by coordinated acetate ligands.

A proposed mechanism for the formation of porous spheres is shown in Fig. 7. BA forms an aqueous microemulsion, which is substantiated by an observed laser Tyndall Effect. BuOH is miscible with both BA and water, but more favorably partitions into BA. The redox reaction between BuOH and KMnO₄ thus occurs inside the BA microemulsion, which serves as a receptacle and template for the manganese oxide nanoparticles. BA also plays the role of capping ligand for the nanoplatelets, limiting their growth. The observed dependence of sphere diameter on KMnO₄ concentration (Fig. 5) is a consequence of more nanoplatelets being formed within the microemulsion.

This proposed mechanism is further supported by reactions carried out at higher and lower temperatures. SEM images of porous manganese oxide spheres prepared at 60 °C and 5 °C are shown in Fig. 8. Reactions at 60 °C yield highly monodisperse spheres, similar to observation at room temperature, but with much smaller diameters. Conversely, reactions at 5 °C produce spheres with much larger diameters, and there is noticeable loss of monodispersity. The increase in sphere size with decreasing

Fig. 4. (A) SEM and (B) TEM images manganese oxide from the control reaction of KMnO₄ and BuOH without BA. (S1 without BA)

Fig. 5. Plot of sphere diameter vs. [KMnO₄] used in each reaction.

Porous manganese oxide spheres are stable to prolonged stirring, sonication, and reflux. However, the spheres are crushed under high compression (10 tons). Thermal stability measured by TGA shows gradual weight loss of 12% from 40 °C to 230 °C due to loss of adsorbed water, Fig. 6A. From 230 °C to 475 °C, a more gradual change of 4% is attributed to the loss of more strongly adsorbed water and chemisorbed O₂. The sharp transition at 475 °C corresponds to loss of O₂ and concomitant formation of Mn₂O₃, as confirmed by XRD. Surface area measured after calcination at 500 °C shows dramatic decrease to 32 m²/g. Sintering is observed as the amorphous nanoplatelets convert to crystalline Mn₂O₃, even while the overall spherical motif is retained, Fig. 6B.

Fig. 6. (A) TGA analysis of porous manganese oxide spheres. (B) TEM image after calcination at 500 °C. Samples prepared as in S1.
reaction temperature is consistent with the trend in carboxylic acid miscibility with water. Lower temperature results in lower BA miscibility, resulting in larger-sized emulsion particles and ultimately larger manganese oxide spheres.

The mechanism shown in Fig. 7 for manganese oxide porous spheres is consistent with what we proposed in a previously published communication on the synthesis of manganese oxide hollow spheres. The latter were obtained by reacting MnO$_4^-$ and Mn$^{2+}$ in aqueous BA. The key difference in the reaction leading to hollow structures is that Mn$^{2+}$ serves as the reducing agent, rather than BuOH, as reported here. Thus, both MnO$_4^-$ and Mn$^{2+}$ reside in the aqueous phase, and the resulting nanoplatelets form outside the BA microemulsion, aggregating on the surface to form a shell. This BA template is easily removed from the porous shell by washing, leaving behind the hollow sphere. By contrast, in the BA/BuOH system reported here, BuOH partitions into the BA microemulsion and reacts with KMnO$_4$ on the inside to form porous spheres comprised of aggregated nanoparticles, Fig. 7.

Oleic acid has previously been reported to react with KMnO$_4$ in an oil-and-water emulsion that generates layered manganese oxide, which in turn aggregates into honeycomb or hollow sphere hierarchical morphologies, depending on KMnO$_4$ concentration. This is a very slow reaction that involves oxidizing the C=C double bond of the water-insoluble oleic acid. There is no evidence of carboxylic acid coordination or other influences on particle size. Reactions between Mn$^{2+}$ with HMnO$_4$ have also been shown to control the formation of $\gamma$-MnO$_2$ as hollow or solid spheres. The type of hierarchical structure is determined by the rate of permanganic acid addition. Small particle size is reported, as well as the highest surface area for $\gamma$-MnO$_2$.

Reactions between KMnO$_4$ and smaller alcohols (MeOH, EtOH, PrOH) in aqueous BA generate materials that have morphologies resembling that of the control reaction between KMnO$_4$ and BuOH in the absence of BA, Fig. 4. This can be attributed to the greater miscibility of smaller alcohols with water, favoring a reaction in the aqueous phase rather than in the BA microemulsion. However, hollow spheres are not observed either, indicating that nanoparticle aggregation occurs differently from the MnO$_4^-$/$\gamma$-MnO$_2$ system, despite both reactions occurring in the aqueous phase. Longer chain alcohols such as various isomers of pentanol and hexanol are poorly soluble in water, but in small amounts they are viable reducing agents in preparing manganese oxide porous spheres. Work in this area is ongoing.

Isobutyric acid also forms aqueous microemulsions and can be utilized to prepare porous manganese oxide spheres, similar to BA. However, other carboxylic acids do not serve as good templates. Acetic acid and propionic acid do not form microemulsions, based on the lack of an observed Tyndall Effect. Their presence in the KMnO$_4$/BuOH reaction gives product similar to the control reaction without carboxylic acid. Longer chain carboxylic acids are poorly soluble in water and have not been successful in preparing monodisperse spheres.

Porous manganese oxide spheres are active catalysts for the oxidation of i-PrOH to acetone. Catalysis is carried out in the gas phase by bubbling a stream of He carrier gas with 1% O$_2$ through liquid i-PrOH. Conversion percentages range from 67% to 84% at 200 °C for different diameter spheres, Fig. 10. By comparison, manganese oxide prepared in a control reaction without BA converted only 40%. The enhanced activity may arise from the smaller particle size and higher surface area, which could translate into more active sites. This is particularly important if the sites are edges on the amorphous nanoplatelets, since these sites would increase considerably as platelet size decreases. The enhancement of oxidation catalysis aligns with results reported earlier for hollow spheres from our KMnO$_4$/MnSO$_4$/BA system and its ability to promote conversion of CO to CO$_2$. Decreases in particle size and increases in surface area have been attributed to greater catalytic activity in the $\gamma$-MnO$_2$ system cited above, which is also synthesized with both solid and hollow sphere morphologies. Additional catalytic studies are ongoing and beyond the scope of this communication.

Fig. 8. SEM images of spheres (SI procedure) at (A) 60 °C and (B) 5 °C. Spheres from the room temperature reaction are shown in Fig. 1A.

Fig. 9. Proposed mechanism for hollow manganese oxide spheres. In this reaction, the reducing agent is Mn$^{2+}$ instead of BuOH. Since both Mn$^{2+}$ and MnO$_4^-$ favor the aqueous phase, the reaction occurs outside the BA microemulsion and the resulting nanoparticles aggregate on the surface to form a shell. Washing removes the BA template.

Fig. 10. Percent catalytic conversion of i-PrOH to acetone for porous manganese oxide spheres of varying diameters.
Conclusions

The reaction between KMnO₄ and BuOH in aqueous butyric acid provides a facile route to monodisperse manganese oxide porous spheres with tuneable diameters and high surface areas. The proposed mechanism has butyric acid serving both as a ligand that promotes nanoplatelet formation and a microemulsion that serves as a soft template for nanoparticle aggregation and sphere growth. Initial studies indicate that this is a promising material for oxidation catalysis.

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Notes and references