Single-enzyme biomineralization of cadmium sulfide nanocrystals with controlled optical properties

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Biological systems have evolved a diverse array of mechanisms to synthesize inorganic materials from aqueous solutions under ambient conditions. This inherent control over material properties has created interest in using these biological routes to synthesize materials (1–3) such as biosilica from sponges and diatoms (4–8), biogenic CaCO$_3$ from mollusks (9–13), and magnetic particles from magnetotactic bacteria (13–15). Designing a biomineralization strategy requires control of both the material composition and structure; in nature, this control is typically achieved through the assembly of a multiprotein complex, including both structure-directing proteins and proteins responsible for mineralization of a specific composition. In the current work, we demonstrate the reduction of this complexity to its simplest form: a single enzyme capable of both catalyzing CdS mineralization and controlling particle size within the quantum confined size range to form functional biomineralized CdS quantum dots.

Two of the most studied biomineralization proteins are perlucin and silicatelin. Perlucin (16) has been shown to mineralize crystalline forms of CaCO$_3$, a common structural material that constitutes the shell of many marine organisms, in the form of organic–inorganic composites. The role of the nacre protein perlucin in crystallite templating has been elucidated through experiments demonstrating crystallite formation in the presence of purified perlucin, and perlucin selectively being removed from solution during crystallite formation in the presence of a mixture of water-soluble, nacre-associated pro-teins (17). Native silicatelin harvested from sea sponge or engineered forms produced recombinantly are active for biomineralization of silica and titania into structures that are amorphous or crystalline (7, 18, 19). In particular, biomineralization confined within an inverse micelle using an engineered silicatelin has been demonstrated to form crystalline nanomaterials (20).

With regard to semiconducting metal sulfides, the majority of prior reports have used biological components to template structure during synthesis from reactive sulfur precursors, typically Na$_2$S (21–23). For example, peptides displayed on the surface of bacteriophage have been used to produce intricately ordered CdS and ZnS nanowires (24), whereas engineered viral particles (25, 26) along with recombinant, self-assembled cage-like proteins (27) have served as nucleation cavities for CdS nanocrystals. Spoerke and Voigt (21) engineered a library of CdS-capping peptides capable of templating CdS nanocrystal growth from cadmium acetate and Na$_2$S, with the resulting optical properties dependent on the chemical structure of the capping peptide. Singh et al. (28) engineered bifunctional peptides with CdSe- and ZnS-binding domains for the synthesis of CdSe/ZnS core shell nanoparticles by the re-action of CdCl$_2$ with NaHSe followed by the addition of ZnCl$_2$ and Na$_2$S. Harvested natural phytochelatins (22) or engineered cell-based methods to produce phytochelatins (29) have also been developed for the synthesis of CdS nanoparticles. Another intriguing approach is the use of a self-assembled ribbon-like template to control synthesis of CdS nanohelices (30). To the best of our knowledge, the only example of enzymatic production of a reactive sulfur species is that of a sulfite reductase (31) reducing Na$_2$SO$_3$ to generate larger, 5–20 nm, CdS nanoparticles by using additional templating peptides. That is, the enzyme does not template the crystal growth.

We previously identified a putative cystathionine γ-lyase (smCSE) associated with the extracellular synthesis of CdS quantum dot nanocrystals by Stenotrophomonas maltophilia strain SMCD1 (32). Cystathionine γ-lyases (CSEs) are a class of enzymes that catalyze the production of H$_2$S, NH$_3$ and pyruvate from γ-cysteine, and the overexpression of which has been shown to precipitate cadmium sulfide (33). In this study, we demonstrate the putative smCSE from S. maltophilia SMCD1 is capable of reactive H$_2$S generation.

Significance

Biominalization is a promising, yet complex, route toward the scalable and green biomanufacturing of functional nanomaterials, involving multiple biomolecules acting in unison to control mineralization and crystal structure. Unraveling the interdependent complexity of biomineralization is a barrier to completely realize this approach. We distill this complexity to a single enzyme that both catalyzes the formation of the reactive precursors required for mineralization and templates nanocrystal growth in solution. This is the first report of a single enzyme capable of providing all of the required functionality for active biomineralization from otherwise unreactive solution. This work provides insight into the mechanism of metal sulfide biomineralization and is an example of the elegance in green functional material synthesis achievable through engineered biomineralization.


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consistent with its function as a cystathionine γ-lyase. In addition, the purified smCSE enzyme, by itself, is capable of aqueous phase synthesis of CdS nanocrystals directly from cadmium acetate and l-cysteine. The resulting CdS nanocrystals are within the quantum confined size range and display optoelectronic properties analogous to those previously described for cell-based or chemically synthesized CdS nanocrystals (21–23, 32, 34–37). When the substrate l-cysteine is replaced by the chemical precursor Na2S, smCSE is capable of directing CdS nanocrystal formation in solution. Removal of smCSE results in bulk CdS formation, indicating a role for smCSE not only in H2S generation, but also in templating CdS nanocrystals. Therefore smCSE is capable of synthesizing metal sulfide nanocrystals directly from aqueous solution, opening up a wide range of strategies for engineering the biomineralization of functional materials.

**A Putative smCSE Associated with Biosynthetic Quantum Dots Is Capable of H2S Generation**

Previous studies have shown that overexpressed CSEs are capable of precipitating cadmium sulfide in cell culture (33, 38–42) and suggested that H2S generation from CSE was the primary driver for CdS precipitation. To determine whether the putative CSE identified from S. maltophilia (smCSE) was capable of H2S generation from l-cysteine, smCSE was heterologously overexpressed and purified from Escherichia coli and the intrinsic kinetics of l-cysteine turnover to H2S measured (described in SI Appendix, SI Materials and Methods). Expression and purification using immobilized metal affinity chromatography yielded a single protein of 42 kDa (SI Appendix, Fig. S1), consistent with the expected size of smCSE (NCBI reference no. WP_012509966.1). The purified smCSE also shows a strong absorbance peak at 430 nm (SI Appendix, Fig. S1), which is indicative of a covalently bound pyridoxal 5’-phosphate (PLP) cofactor; PLP is an obligate cofactor required for CSE catalysis (43). The Michaelis–Menten rate parameters for l-cysteine turnover measured for smCSE using l-cysteine as a substrate are consistent with reported values for other well-characterized CSEs (44, 45) (SI Appendix, Fig. S1).

The Quantum Dot-Associated smCSE Regulates CdS Nanocrystal Growth and Size

Having established the ability of smCSE to generate reactive sulfur species, it was hypothesized that the addition of a cadmium salt to the solution would yield CdS. To test this hypothesis, l-cysteine (4 mM) and cadmium acetate (0.5 mM) were added to an aqueous solution of smCSE (0.1 mg/mL in Tris buffer, pH 7.5), and the solution was monitored by using UV-visible and fluorescence spectroscopy as a function of time at ambient temperature and pressure (Fig. 1). After 90 min, a distinct absorbance peak was observed with an absorbance maximum at 333 nm (Fig. 1A). A corresponding fluorescence emission peak at 464 nm was also observed (Fig. 1B). The maxima of both absorbance and fluorescence progressively shift to longer wavelengths over the course of 195 min (Fig. 1A and B) with visible photoluminescence observed in solution for each time point measured when illuminated under UV light (Fig. 1C). It should be noted that the presence of smCSE, cadmium acetate, and l-cysteine were all required; omission of any one of these components produced solutions that displayed no absorbance or fluorescence peaks and no observable photoluminescence when illuminated under UV light (SI Appendix, Fig. S2). Placing the reactant mixture on ice essentially arrests smCSE activity, allowing a specific population of particles with a given set of photoluminescent properties to be collected.

**Enzymatically Synthesized CdS Quantum Dots Are Monodisperse and Crystalline**

Crystallites were harvested from solution after 180 min of growth (absorbance maximum at 350 nm) and analyzed via scanning transmission electron microscopy (STEM). Fig. 2A shows the existence of discrete, but irregularly shaped and overlapping, nanocrystals between 2–4 nm in diameter. Elemental analysis...
from this region using X-ray energy-dispersive spectroscopy (XEDS) (Fig. 2B) reveals strong Cd and S signals. The presence of strong Cu and C peaks are artifacts arising from the copper mesh TEM grid coated with a carbon film. High-resolution high-angle annular dark field (HAADF) imaging of individual and isolated particles confirms the size of the nanocrystals and the presence of both wurtzite (Fig. 2C and D) and zinc-blende structured (Fig. 2E and F) nanocrystals. The lattice parameters and interplanar angles of these crystals are consistent with those for CdS (SI Appendix, Tables S1 and S2). These lattice parameters and both crystalline phases have been observed for chemically and biologically synthesized CdS nanocrystals (21, 36).

Powder X-ray diffraction measurements (SI Appendix, Fig. S3) on material harvested after 12 h growth is also consistent with the presence of both structures of CdS.

The irregular shape of the biomineralized nanocrystals prevents a precise comparison with reported literature correlations (46) for the optical properties as a function of particle diameter. However, for all reported growth times, the absorbance maximum (Fig. 1A) remains well below the absorbance maximum corresponding to the band gap of bulk CdS (2.5 eV, 495 nm) (47, 48) and the observed biomineralized CdS nanocrystals size is well within the quantum confined size range for CdS (47). The observed red shift in optical properties with growth time (Fig. 1A and B) is indicative of CdS nanocrystal growth as a function of time within this quantum-confined size range. For reference, our previous work (32) demonstrated that absorption peak maxima of 324, 334, and 344 nm correspond to biomineralized nanocrystallite sizes of 2.75 ± 0.68, 3.04 ± 0.75, and 3.36 ± 0.95 nm, respectively. Additionally, smCSE-produced CdS nanocrystals show similar quantum yield (1.8%) compared with synthesized CdS nanocrystals from S. maltophilia (2%) (32).

CdS Quantum Dot Growth Depends on both H2S Production and Available Capping Agents

Although l-cysteine is a substrate for smCSE to generate H2S, it can also act as an aqueous phase capping agent for CdS (34). To confirm the dual role of l-cysteine as the sulfur source and capping agent, CdS biomineralization was studied as a function of l-cysteine concentration. At a concentration of 4 mM l-cysteine, found to be the practical lower limit for nanocrystal synthesis at 0.1 mg/mL smCSE and 0.5 mM cadmium acetate, the observed optical absorbance maximum increases with time, reaching 370 nm after 4 h of growth (Fig. 3A); longer growth times beyond 4 h led to formation of bulk CdS with an absorbance maximum consistent with bulk CdS (SI Appendix, Fig. S4).

Increasing l-cysteine concentration to 10 and 20 mM red shifts the absorbance maximum and corresponding photoluminescence during growth (Fig. 3A and B). This red shift is indicative of an increase in CdS growth rate with increasing l-cysteine concentration for nanocrystals within the quantum confined size range and is particularly noticeable for the 20 mM l-cysteine data. Interestingly, all of the growth curves reach similar maxima after 4 h: 370, 375, and 380 nm at 4, 10, and 20 mM l-cysteine, respectively. At longer growth times the solutions lose optical clarity, indicating a maximum in the solubility of the presumably l-cysteine–capped nanocrystals; images of solutions illuminated by using UV light at each l-cysteine concentration and growth time are given in Fig. 3B to demonstrate optical clarity at each solution condition.

To further demonstrate that thiol-mediated capping is important for stabilizing the nanocrystals, glutathione was introduced into the enzymatic synthesis mixture. Glutathione is a capping agent derived from l-cysteine and l-glutamine that has been shown previously to stabilize water-soluble CdS nanocrystals (22, 49). However, unlike l-cysteine, glutathione is not a substrate for CSEs (50) and, therefore, acts solely as a capping agent. No nanocrystal formation was observed in the absence of l-cysteine which clearly demonstrates that l-cysteine is the sulfur source for the nano-crystal growth through the enzymatic generation of H2S.

In the presence of 4 mM l-cysteine, 1 and 4 mM glutathione addition yielded growth curves and absorbance maxima at 4 h (375 and 385 nm for 1 and 4 mM glutathione, respectively), similar to those with elevated l-cysteine concentration, although the solutions maintain optical clarity after 5 h of growth (Fig. 3C and D). This retention of optical clarity is indicative of increased stabilization of the aqueous nanocrystal solution with the glutathione capping agent versus l-cysteine capping agent. Although the 4 mM glutathione growth curve shows some indication of a decreased growth rate, this phenomenon is clearly observed upon increasing the glutathione concentration to 10 mM. Compared with lower glutathione concentrations, the 10 mM glutathione growth curve shows a blue shift in absorbance maximum at all growth times and yields an optically clear solution with absorbance maximum at 400 nm after 24 h of growth (Fig. 3C and D).

To further demonstrate the role of glutathione as a capping agent, the absorbance maximum and l-cysteine concentration were measured as a function of time (SI Appendix, Fig. S5) in the presence and absence of glutathione. Whereas l-cysteine degradation is not influenced by the presence of glutathione, the absorbance maximum occurs at lower wavelength, indicating a smaller average particle size, for the glutathione containing sample. This reduction in average size is consistent with the capping role envisioned for glutathione because an increased capping agent concentration will lead to smaller average particle size. Where the
\( \text{Absorbance} \)

<table>
<thead>
<tr>
<th>Concentration (mM)</th>
<th>1</th>
<th>4</th>
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<tr>
<td>Time (hours)</td>
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<td>3</td>
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Absorption maxima for the glutathione and L-cysteine capping ligands are sharper, which likely reflects their higher affinity for the particle surface and their smaller size, both of which lead to higher coverage on the particle surface during growth (21, 51). Thus, in all three cases, smCSE, L-cysteine, and glutathione act as structure directing agents to form CdS nanocrystals, rather than bulk CdS, from Na\(_2\)S and cadmium acetate. Therefore smCSE has the intrinsic ability to regulate CdS nanocrystal growth independently from, and in addition to, its role in reactive H\(_2\)S generation (Fig. 4C).

**Discussion**

This work clearly demonstrates the ability of a single enzyme (smCSE) to produce both crystalline CdS (Fig. 2) and regulate growth to form nanocrystals within the quantum confined size range (Fig. 1). Previous studies involving enzymatic or peptide-based biominalization of metal sulfide nanoparticles have demonstrated either CdS formation from enzymatically generated H\(_2\)S without intrinsic size control (31, 33), or have demonstrated nanocrystal size control by adding specific peptides or proteins and use reactive Na\(_2\)S as a sulfur source (21–23). As an example, Wang et al. overexpressed Treponema denticola CSE in E. coli (33) and demonstrated that aqueous cadmium ions were removed under aerobic growth conditions. In this instance, the flux of H\(_2\)S from the cell surface led to the nucleation and growth of CdS precipitates over the course of 48 h without apparent size control. An alternative example used engineered overexpression of cysteine-rich phytochelatin peptides in E. coli (29), which are known to sequester metal ions in solution, to produce 3–4 nm CdS nanocrystals in E. coli cells in the presence of the reactant Na\(_2\)S. In contrast, smCSE acts...
both as the sulfur generating source to mineralize CdS (Fig. 1) and as a structure directing agent to control nanocrystal growth (Fig. 4). Thus, smCSE efficiently combines mineralization and templating into a single enzyme (smCSE; Fig. 4C) that are engineered separately in other biomineralization strategies.

The combination of H\(_2\)S generation and controlled growth afforded by smCSE is remarkable compared with other enzymatic and peptide approaches to nanocrystal synthesis. In the presence of excess capping agents (L-cysteine or glutathione) under conditions where smCSE can generate H\(_2\)S, we find that the CdS nanocrystals can span a range of quantum confined sizes and resulting optical properties (Fig. 3). Reducing the amount or quality of capping agent limits the stability of the resulting colloidal solution (Fig. 3). It is noted that the addition of capping agents L-cysteine or glutathione appear to improve the nanocrystal quality as well through providing capping agents with smaller size and higher affinity for the particle surface (21, 51).

Independent of its role in H\(_2\)S generation, smCSE is also capable of templating CdS nanocrystals from solution when using chemical precursors such as Na\(_2\)S (Fig. 4), to exhibit intrinsic structure directing and capping activity similar to naturally occurring phy- tochelatins (22, 23, 29) or nanocrystal-binding peptides (21, 24, 28).

In a previous study using the protein pepsin as a templating agent (52), the rate of reactive sulfur generation and subsequent nanocrystal synthesis was at least an order of magnitude slower than that of smCSE (days versus hours). This rate is dictated by the slow, spontaneous hydrolysis of thioacetamide to generate reactive sulfur, in contrast to the smCSE-catalyzed H\(_2\)S generation demonstrated in this work.

Given the intrinsic heavy metal resistance of S. maltophilia (41), our results would suggest that smCSE secretion may be one of several independent mechanisms of heavy metal resistance used by microorganisms to both sequester metal ions and convert them into insoluble precipitates outside of the cell. The strategy demonstrated herein distills the potential complexity of biomineralization down to its simplest form: a single enzyme. The ongoing challenge is to develop strategies that enable a wider range of materials synthesis.

In summary, an engineered smCSE is capable of controlled CdS nanocrystal synthesis directly from aqueous solution by using L-cysteine and cadmium acetate as reactants. Furthermore, the ability of smCSE to mineralize CdS and template nanocrystal formation provides a single enzyme route for engineered nanocrystal biomineralization.

Materials and Methods

Expression, Purification, and Kinetics of Recombinant CSE. An E. coli codon-optimized form of the putative S. maltophilia CSE (Smal_0488; Genscript) was subcloned into pET28a (+) and transformed in BL21 cells with expression and purification as described (53). Detection of H\(_2\)S product formation using the substrate L-cysteine to determine Michaelis–Menten rate parameters was performed as described (54).

Cadmium Sulfide Nanocrystal Biosynthesis. smCSE (0.1 mg/mL) was incubated with 0.5 mM cadmium acetate and 4 mM L-cysteine in 100 mM Tris buffer (pH 7.5). CdS nanocrystal formation was directly observed in solution as a function of time by illumination using a UV lamp. Absorbance spectra for CdS nanocrystals were taken at regular time intervals on a Shimadzu UV-Vis 3300 spectrometer, and fluorescence emission spectra were measured on a PTI QuantaMaster fluorimeter with a 350-nm excitation wavelength by using a 5-nm excitation slit width. To demonstrate the ability of CSE to control CdS nanocrystal growth and size independent of H\(_2\)S generation, Na\(_2\)S was added instead of L-cysteine to a final concentration of 4 mM in a solution containing 0.5 mM cadmium acetate and 0.1 mg/mL CSE. Absorbance and fluorescence spectra as well as direct observation of solutions under UV light were used to characterize the optical properties of the CdS nanoparticles in solution.

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Transmission Electron Microscopy. Biosynthetic CdS nanocrystals were grown for 3 h, corresponding to an absorbance maximum at 350 nm, and subsequently dialyzed at 4 °C against a reservoir containing 0.5 mM L-cysteine for 24 h. The dialyzed CdS nanocrystals solutions were placed dropwise onto Ar plasma-treated carbon-coated TEM grids and the liquid evaporated under vacuum. High resolution (HR)-TEM and HAADF-STEM images were taken on a 200-kV aberration corrected JEOL ARM 200CF analytical electron microscope equipped with a Centurio XEDS system.

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