

Synthesis and Photoluminescence Characterization of Dendrimer-Encapsulated CdS

Quantum Dots

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Abstract

We synthesized CdS quantum dots (QDs) encapsulated within n th-generation hydroxyl-terminated poly(amidoamine) (PAMAM) dendrimers (Gn -OH) by coordination of Cd^{2+} ions and subsequent reaction with Na_2S . We systematically examined the effects of Cd^{2+} ion/dendrimer ratio (x), generations of dendrimers, and temperatures on the resultant CdS QDs to elucidate the formation mechanism and to optimize the conditions for the encapsulation of QDs. The resultant QDs were analyzed by UV-vis spectroscopy, photoluminescence spectra spectroscopy, and transmission electron microscopy (TEM). Through a detailed analysis, we concluded that the first requisite for stable dispersions of CdS QDs with Gn -OH is to set x to be less than 2^n because a Cd^{2+} ion is supposed to bind to two adjacent coordination sites of 2^{n+1} outermost tertiary amines of a Gn -OH dendrimer. Further, we clarified that higher-generation dendrimers can confine smaller QDs because of denser surfaces. Finally a lower temperature is found to promote the coordination of Cd^{2+} ions with dendrimers, and accordingly suppress the nucleation outside the dendrimers which produces larger QDs.

1. Introduction

Semiconductor nanoparticles ranging from one to tens of nanometers in diameter, also known as quantum dots (QDs), exhibit unique size-tunable optical and electronic properties owing to the quantum size effect [1], unlike traditional fluorochromes [2] and fluorescent proteins [3]. QDs are of great interest for various applications, e.g., in optical devices, quantum information processing, and biotechnology[4-9]. CdS is one of the most researched semiconductor materials because bulk CdS has a band gap of 2.53 eV, which lies in the visible wavelength region. Many research groups have attempted to synthesize CdS QDs by various methods [10] since the successful preparation by Brus et al. in the early 1980s [11]. The synthetic methods should meet the following three requirements to fully exhibit the properties of the QDs. First, suspensions are preferred over aerosols or solid sols for better handling. Synthesizing QD suspensions is a popular topic and has been realized using various solvents over the last two decades. Second, the QDs should be strictly size-controlled and monodispersed. A size difference of a few angstroms can lead to a drastic change in their properties [12]. Third, the surfaces of the QDs should be modified to have varied functionality, which includes hydrophilicity, low-toxicity, and electric charge. However, a synthetic method that completely satisfies the three requirements has not yet been developed.

A dendrimer, which is a spherical hyperbranched polymer, is a possible candidate for use in QD synthesis. The structure of a dendrimer is such that it grows in radial directions from a core

with branches (the number of repeated branching cycles is called its “generation”) and is terminated by a surface functional group [13]. Dendrimers are commonly represented as G_n-X according to their generation (n th generation) and surface functional group (X), both of which can be tuned. Poly(amidoamine) (PAMAM) dendrimers with an ethylenediamine core can serve as templates of metal nanoparticles. Crooks et al. and Balogh and Tomalia independently succeeded in encapsulating monodispersed Cu nanoparticles in PAMAM dendrimers [14, 15]. Since then, various types of metal nanoparticles have been prepared by encapsulation [16]. Nanoparticles encapsulated within PAMAM dendrimers can be synthesized by the incorporation of metal ions into dendrimers by coordinate bond formations with tertiary amine groups that are present at branching sites of the dendrimers, and by the subsequent particle formation with reductants such as NaBH_4 . The main advantage of this method is that resultant nanoparticles are monodispersed with controllable particle size.

This encapsulation method can be used for the synthesis of CdS QDs by using sulfides such as Na_2S instead of reductants in the particle formation process after the coordination of Cd^{2+} ions with dendrimers. Several research groups have prepared CdS QDs encapsulated within PAMAM dendrimers, and most of them have used amine-terminated dendrimers ($G_n\text{-NH}_2$) in their preparation [17-22]. However, a major limitation of this method is that the resultant CdS QDs are less stable than other dendrimer-encapsulated metal nanoparticles [20], which can be because Cd^{2+}

ions bound to terminal primary amine ligands cause nucleation outside the dendrimers. For the synthesis of dendrimer-encapsulated Pt nanoparticles, hydroxyl-terminated dendrimers ($Gn-OH$) are preferred over $Gn-NH_2$ dendrimers because Pt^{2+} ions do not have a specific interaction with hydroxyl groups, and they can be transferred inside the dendrimers through the dendrimer surface. Moreover, we have demonstrated that a high coordination ratio of Pt^{2+} ions, which is defined as the ratio of metal ions coordinated with dendrimers to the total amount of metal ions, results in stable dendrimer-encapsulated Pt nanoparticles, whereas a low coordination ratio of Pt^{2+} ions produces particles outside the dendrimers [23]. Thus, the main reason for the aggregation of CdS QDs may be attributed to insufficient Cd^{2+} coordination. Some research groups have also synthesized dendrimer-encapsulated CdS QDs by using $Gn-OH$ dendrimers [24-26]. However, the relationship between the Cd^{2+} coordination and the resultant CdS nanoparticles has not yet been completely understood because the coordination cannot be quantified by using UV-vis measurement. The spectra of the Cd^{2+} ions remained unchanged during coordination, unlike other metal ions [27]. This resulted in a difficulty in elucidating the Cd^{2+} coordination mechanism.

Recently, Bard et al. succeeded in quantifying the Cd^{2+} coordination with cyclic voltammetry [28]. They demonstrated that the Cd^{2+} coordination with G2-OH in water was accomplished in 30 min, which is much shorter than several days for Pt^{2+} , Ru^{3+} , or Ir^{3+} ions [29-31]. This rapid Cd^{2+} coordination possibly results from a fast H_2O -ligand-exchange (aquation) rate

because we have observed in our previous study that a slower aquation rate of metal ions, such as Pt^{2+} ions, results in a longer coordination period and vice versa [32]. This is why we investigated the dynamics of the coordination process in Pt^{2+} ions in that study. On the other hand, the key factor for coordination of Cd^{2+} ions would be thermodynamic equilibrium rather than kinetics because Cd^{2+} ions are predicted to have a lower coordination ratio than that of the solution using Pt^{2+} ions. We calculated the coordination ratio of a 1 mM $[\text{G2-OH} + 4 \text{ Cd}^{2+}]$ solution to be 70% at equilibrium by using the coordination equilibrium constants obtained by Bard et al. (refer to Supporting Information for calculation of the coordination ratio of a solution containing Cd^{2+} ions at equilibrium) [28]. Thus, enhancing the coordination ratio at equilibrium is our strategy to produce stable CdS QDs encapsulated within dendrimers without aggregations.

In the present study, we systematically examined the effects of the Cd^{2+} ion/dendrimer ratio (x), generations of dendrimers, and temperatures—all of which affect the coordination ratio at equilibrium—on the resultant CdS QDs encapsulated within hydroxyl-terminated PAMAM dendrimers (Gn-OH); our aim was to elucidate the formation mechanism of dendrimer-encapsulated CdS QDs and to determine the optimal conditions required for their encapsulation.

2. Experimental Section

2. 1. Chemicals

The chemicals used in this study, which were *n*th-generation hydroxyl-terminated PAMAM dendrimers (*Gn*-OH in methanol solution, *n* = 4, 6, and 7), sodium polyphosphate (SPP) [(NaPO₃)₆, 96%], cadmium nitrate tetrahydrate [Cd(NO₃)₂·4H₂O, 99.999%], and sodium sulfide (Na₂S), were purchased from Aldrich Chemical Co. Methanol (99.8%, Kishida Chemical Corp.) was used as solvent to prepare the dendrimer-encapsulated CdS QD suspensions. A control experiment was carried out using deionized (DI) water (18 MΩcm, Millipore) as solvent to prepare SPP-protected CdS QD suspensions [21, 33]. All chemicals were used without further purification.

2.2. Preparation of CdS QDs.

Gn-OH solutions (500 μM) were prepared first by being diluted with methanol, and the resulting solution was then mixed with a Cd(NO₃)₂ solution to obtain [*Gn*-OH + *x* Cd²⁺] solutions (*x* = 5, 20, 50, 100, 200, or 400). The concentration of Cd²⁺ ions was adjusted to 526 μM. The solutions were vigorously stirred for a few minutes and aged for 1 h at room temperature and 50 °C, or for 1 day at 4 °C, −18 °C, and −78 °C to achieve equilibrium in Cd²⁺-tertiary amine coordination. Next, 0.25 ml of Na₂S solution (10 mM) was added to 4.75 ml of [*Gn*-OH + *x* Cd²⁺] solution. In our experiments, the concentration of both Cd and S in the CdS QD suspensions was set to be 500 μM, whereas the concentration of *Gn*-OH varied with *x* (500/*x* μM).

SPP-protected CdS QDs were prepared by a procedure similar to the above mentioned one, except that SPP replaced *Gn*-OH as a protective agent and DI water replaced methanol as solvent.

The concentration of SPP was set to be 500 μM .

2. 3. Instruments and Analysis.

UV-vis absorbance spectra of the resultant CdS QD suspensions were measured with a spectrophotometer (UV-1700, Shimadzu) by using quartz cells (path length: 10 mm). Room-temperature photoluminescence (PL) spectra were obtained using a spectrofluorometer (FP-6500, JASCO Corp.) and fluorometer cells (path length: 10 mm). The extinction wavelength (λ_{ex}) was set to be 374 nm for all samples, which is around the wavelength of maximum absorbance. The resultant CdS particles were observed by transmission electron microscopy (TEM) (JEM-1010, JEOL). Samples for TEM observation were prepared by placing a drop of the CdS QD suspensions on a carbon-coated copper grid (Okenshoji Co.), which was then dried in vacuum. Freshly prepared samples were used for all measurements and observations.

3. Results and Discussion

3. 1. Effects of x on resultant CdS QDs.

The G6-OH CdS QD suspensions were colorless ($x = 5, 20, 50$, and 100) or faint yellow ($x = 200$ and 400). However, SPP-protected CdS QD suspensions were yellow, which reflects the color of bulk CdS. This demonstrates that smaller particles were produced in the case of dendrimers than those with SPP. Fig. 1 shows the UV-vis absorbance spectra of G6-OH CdS QDs

for various values of x ($x = 5, 20, 50, 100, 200$, and 400) prepared at room temperature. The experimental diameters of resultant CdS nanocrystals were determined from the absorbance peak by using the empirical equation proposed by Peng et al. [34], which are summarized in the third column of Table 1. Theoretical values of these average sizes were also calculated by assuming spherical-shape nanocrystals that consisted of x atoms of Cd and S, respectively, by using the approximation method proposed by Lippens and Lannoo [35]. The experimental diameters of the CdS nanocrystals decrease as x decreases, and are slightly larger than the theoretical diameters as shown in Table 1. One of the main reasons for the difference between experimental and theoretical diameters is that a CdS particle confined in the limited space in a dendrimer molecule would not be spherical in shape; it would be instead subspherical and resemble a hemisphere, which is similar to other metal nanoparticles encapsulated within dendrimers [36, 37]. Moreover, uncoordinated Cd^{2+} ions cause nucleation and particle growth outside dendrimers, which results in particles with larger diameters because of coalescence. Fig. 2 shows PL spectra of the CdS QD suspensions for various values of x . These spectra were normalized by setting the peak intensity to be 1.0. The PL peak shifted toward shorter wavelengths (blue shift) for smaller values of x , which is similar to the UV-vis absorbance spectra in Fig. 1, because smaller CdS QDs have wider band gaps. Moreover, the

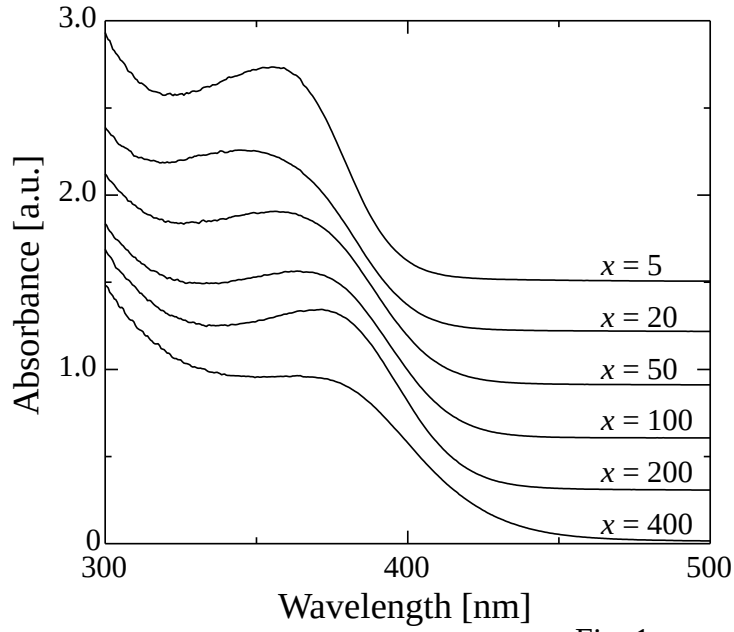


Fig. 1.

x	Absorbance peak [nm]	Experimental diameter [nm] ³¹	Theoretical diameter [nm] ³²
5	355	2.2	0.6
20	344	2.0	1.0
50	356	2.2	1.3
100	364	2.4	1.7
200	372	2.6	2.1
400	375	2.7	2.7

Table 1.

dendrimer-protected QD suspensions have a peak at shorter wavelengths and a significantly higher PL intensity than SPP-protected QD suspensions, which is shown in Fig. S1 in Supporting Information. The shape of PL spectra is multimodal and is attributed to S-related defects (at shorter wavelengths) and Cd-related defects (at longer wavelengths) [38]. The subspherical shape of dendrimer-encapsulated CdS QDs may also lead to multimodal-shaped PL spectra, in contrast to CdS QDs synthesized by other methods [39-41]. According to the PL spectra, the suspensions exhibited different photoluminescence colors varying from blue to orange when illuminated with a

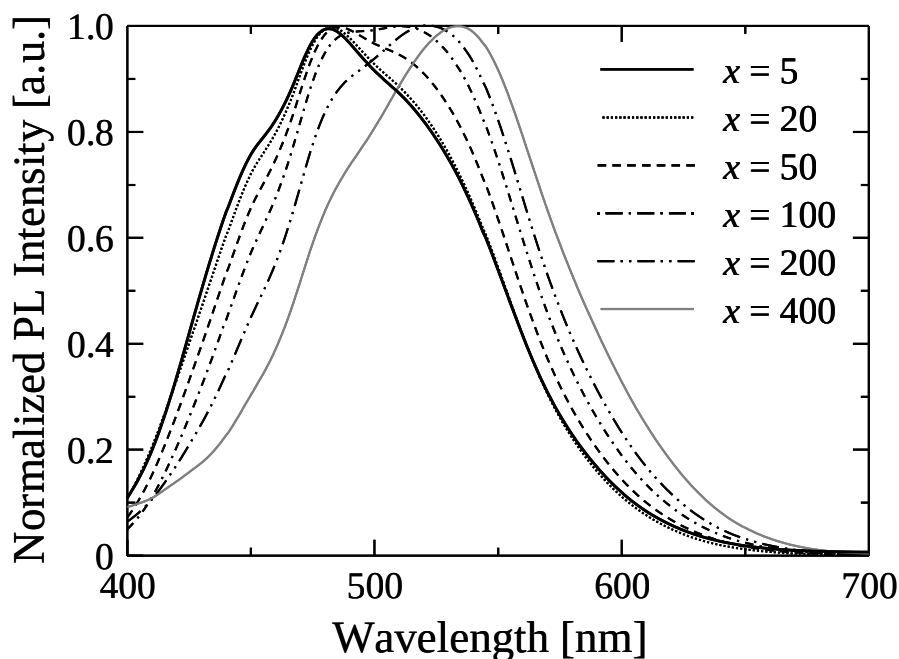


Fig. 2.

UV lamp ($\lambda_{\text{ex}} = 356 \text{ nm}$), whereas a SPP-protected CdS QD suspension turned red (refer to Supporting Information, Fig. S2).

We observed the resultant CdS QDs by TEM to examine the stability in the suspensions (Fig. 3). Note that TEM images can be used to judge the dispersion state (dispersed or aggregated), but are not clear enough to discriminate size and shape of individual particles because of the high transmission nature of the CdS particles. In contrast to well-dispersed particles with $x = 20$ and 50 (Fig. 3a, b), those with $x = 200$ were highly aggregated (Fig. 3c). As in the case of Pt nanoparticles where aggregates are typically observed under a low coordination ratio condition [23], the aggregate formation with $x = 200$ would be attributed to the low coordination ratio of Cd^{2+} ions.

According to our calculations, coordination ratios for $x = 20$ and 200 are estimated to be 93% and 19%, respectively (Fig. S3). When the

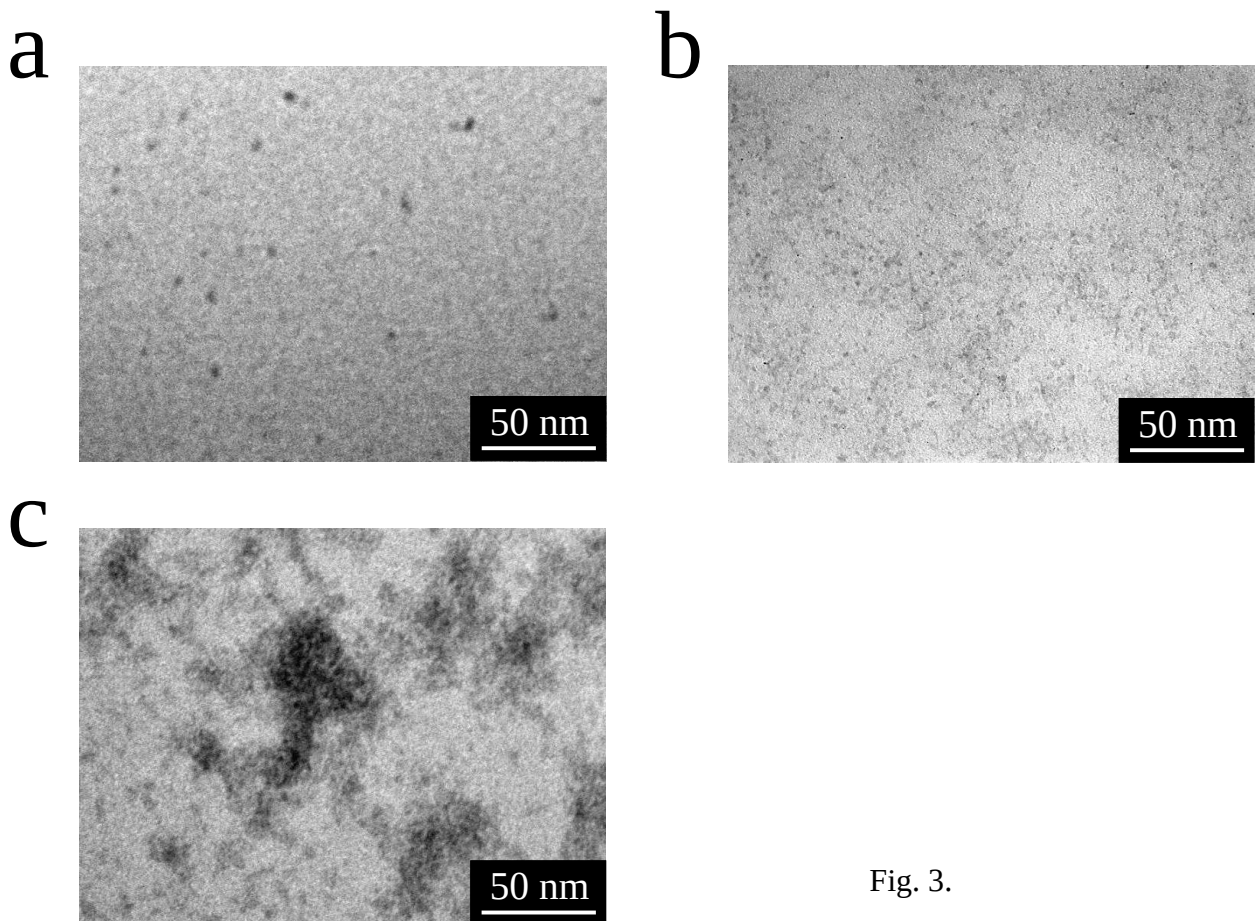


Fig. 3.

coordination ratio is low, nucleation and particle growth occur outside the dendrimers because a number of Cd^{2+} ions remain uncoordinated. Particles not encapsulated in dendrimers aggregate and form subsequent precipitates. In fact, precipitates were observed after a few days for $x = 100, 200$, and 400, whereas no precipitates appeared in several weeks for $x = 5, 20$, and 50. The boundary for the CdS QDs encapsulation thus appears to exist between $x = 50$ and 100. This result agrees with

those reported previously that the coordination behavior of Cd^{2+} ions with dendrimers is similar to that of Cu^{2+} ions, each of which binds to two adjacent coordination sites of the outermost tertiary amines [28, 42]. The maximum coordination number of Cd^{2+} ions within a G6-OH dendrimer would thus be 64 because a G6-OH molecule contains 128 outermost amine sites at the edge. It is thought that stable dispersions of CdS QDs encapsulated within Gn -OH dendrimers are produced by setting the ratio x to be less than 2^n .

Another noticeable observation in Table 1 is that contrary to our expectation, the blue shift of an absorbance peak is not observed for $x = 5$, and the average size ($d = 2.2$ nm) is larger than that in the case of $x = 20$ ($d = 2.0$ nm). This can be attributed to coalescence of CdS clusters outside the dendrimers because the CdS clusters formed in the dendrimers are so small that they can pass through the periphery of dendrimers. Because dendrimers form flocs [43], coalescence occurs more easily inside flocs, which in turn protect resultant nanoparticles for $x = 5$. In fact, precipitations were not observed for $x = 5$, unlike in the case of $x = 400$. Because it was observed that surface density of dendrimers increases with increasing generation [44], we prepared CdS QD suspensions for $x = 20$ with PAMAM dendrimers having various generations (Gn -OH; $n = 4, 6$, and 7) at room temperature (Fig. 4) to examine the effect of the surface density of the dendrimers on the size of resultant CdS QDs. As shown in Fig. 4, the use of a higher-generation dendrimer results in a PL peak at shorter wavelengths, which indicates the formation of smaller QDs. Precipitates were

observed only in G4-OH dendrimers. These results suggest that the size of the smallest QDs that can be encapsulated within dendrimers depends on the generation of the dendrimers. In other words, higher-generation dendrimers having denser surfaces can confine smaller dendrimer-encapsulated CdS QDs.

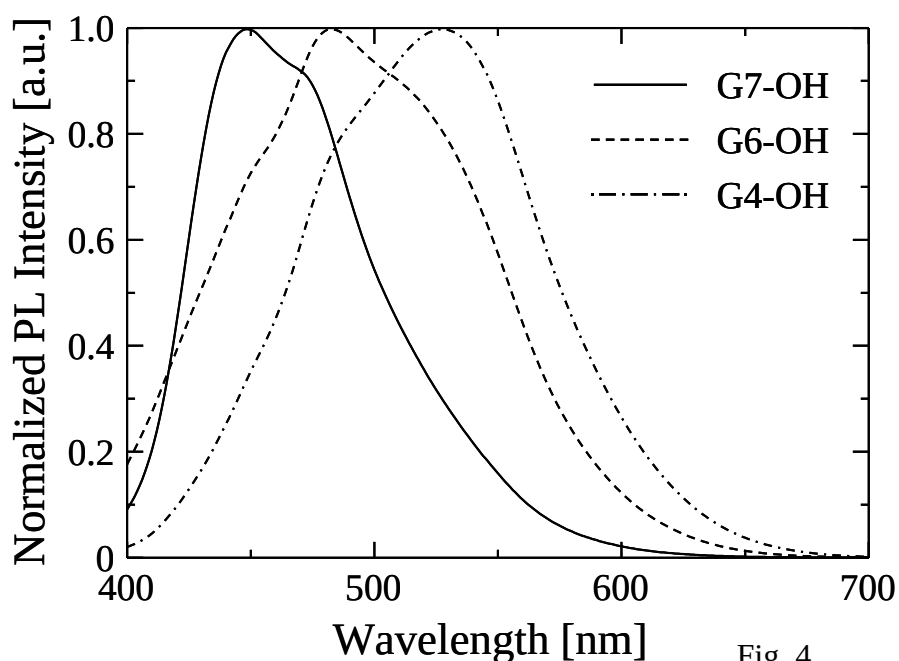


Fig. 4.

3. 2. Effect of temperature on CdS QDs.

In the experiments shown above, the aging period was set to be 1 h to achieve coordination equilibrium. We confirmed that both the UV-vis absorbance spectra and the PL spectra remained unchanged when the duration of coordination was set to be 1 min or 1 day, indicating immediate coordination of Cd^{2+} ions. However, according to our calculation of the coordination ratio of a $[\text{G6-OH} + x \text{ Cd}^{2+}]$ ($[\text{Cd}^{2+}] = 500 \mu\text{M}$) solution, the equilibrium coordination ratio with G6-OH

would not reach 100% at room temperature, even at $x < 64$ (Fig. S3). Nucleation also appears to occur outside the dendrimers for $x = 50$ because the coordination ratio at room temperature is estimated to be only 63% (Fig. S3).

Bryszewska et al. have demonstrated that the coordination process of Cd^{2+} ions is exothermic [45]. Thus, coordination equilibrium shifts towards the Cd^{2+} -tertiary amine complexes as temperature decreases. Fig. 5 shows the PL spectra of G6-OH CdS QDs for $x = 50$ prepared at various temperatures (50 °C, room temperature, 4 °C, −18 °C, and −78 °C). The peak shifts toward shorter wavelengths with a decrease in the temperature, which indicates that lower temperatures yield smaller CdS QDs. Conversely, in the case of SPP-protected CdS QDs, the PL peak showed no shift at 50 °C, at room temperature, and at 4 °C (see Fig. S4; no data were obtained at −18 °C or −78 °C because water was used as solvent), suggesting that the temperature difference had a negligible effect on the nucleation process. Thus, the results demonstrate that blue shift with dendrimers at lower temperatures cannot be attributed to nucleation and particle growth; instead, it can be attributed to a higher coordination ratio of Cd^{2+} ions. A higher coordination ratio at lower temperatures led to a decrease in the number of uncoordinated Cd^{2+} ions, which caused nucleation outside the dendrimers. Therefore, we conclude that lower temperatures are preferred for encapsulation of CdS QDs. However, it should be noted that lowering the temperature is effective only for metal ions that have high coordination rate, such as Cd^{2+} ions, because the lower

temperature slows the coordination rate. For instance, we have demonstrated that Pt^{2+} coordination ratio at 4 °C is only 50% even after 10 days [23].

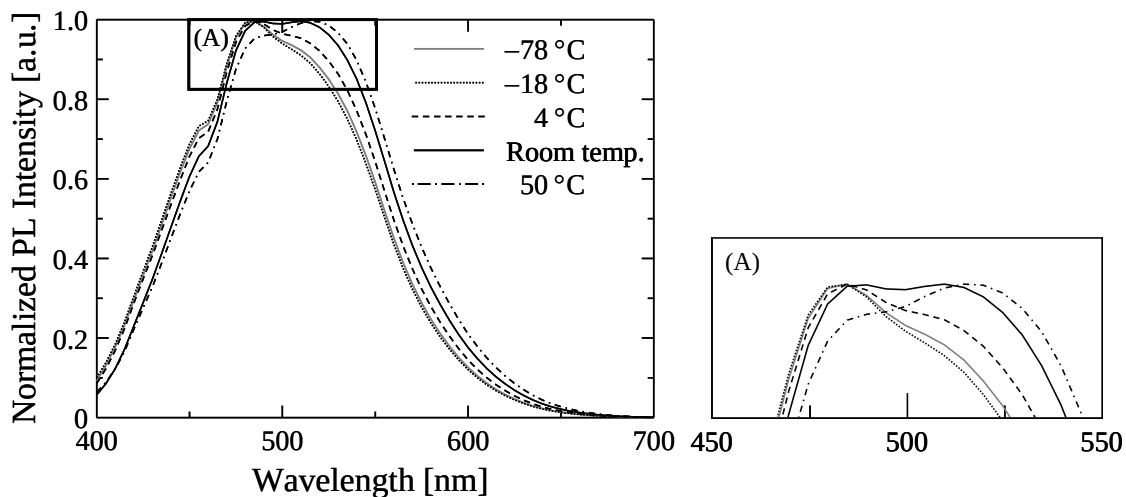


Fig. 5.

3. 3. Formation Mechanism.

From these results, we propose a mechanism of encapsulation of CdS QDs within *Gn*-OH dendrimers (Fig. 6). At a high value of x ($x > 2^n$; x is greater than 64 in the case of G6-OH), most of the nucleation occurs outside the dendrimers and the nanoparticles are not encapsulated but are surrounded by some dendrimer molecules, although precipitates are observed after a few days. At intermediate values of x ($x \sim 2^n$), which means that the value of x is slightly less than 2^n , nucleation occurs not only in the dendrimers but also outside them because the coordination ratio at equilibrium is low at room temperature. A lower temperature is preferred to promote the coordination of Cd^{2+} ions and it leads to the decrease in nucleation outside the dendrimers. At low

values of x ($x < 2^n$), resultant nanoclusters are so small that they cannot be encapsulated within dendrimers. They tend to coalesce into a larger particle. Higher-generation dendrimers are better candidates for templates as they would prevent the clusters from coalescing owing to denser surfaces. Thus, the use of higher-generation dendrimers at lower temperatures is a requisite for the preparation of stable CdS QDs encapsulated within dendrimers.

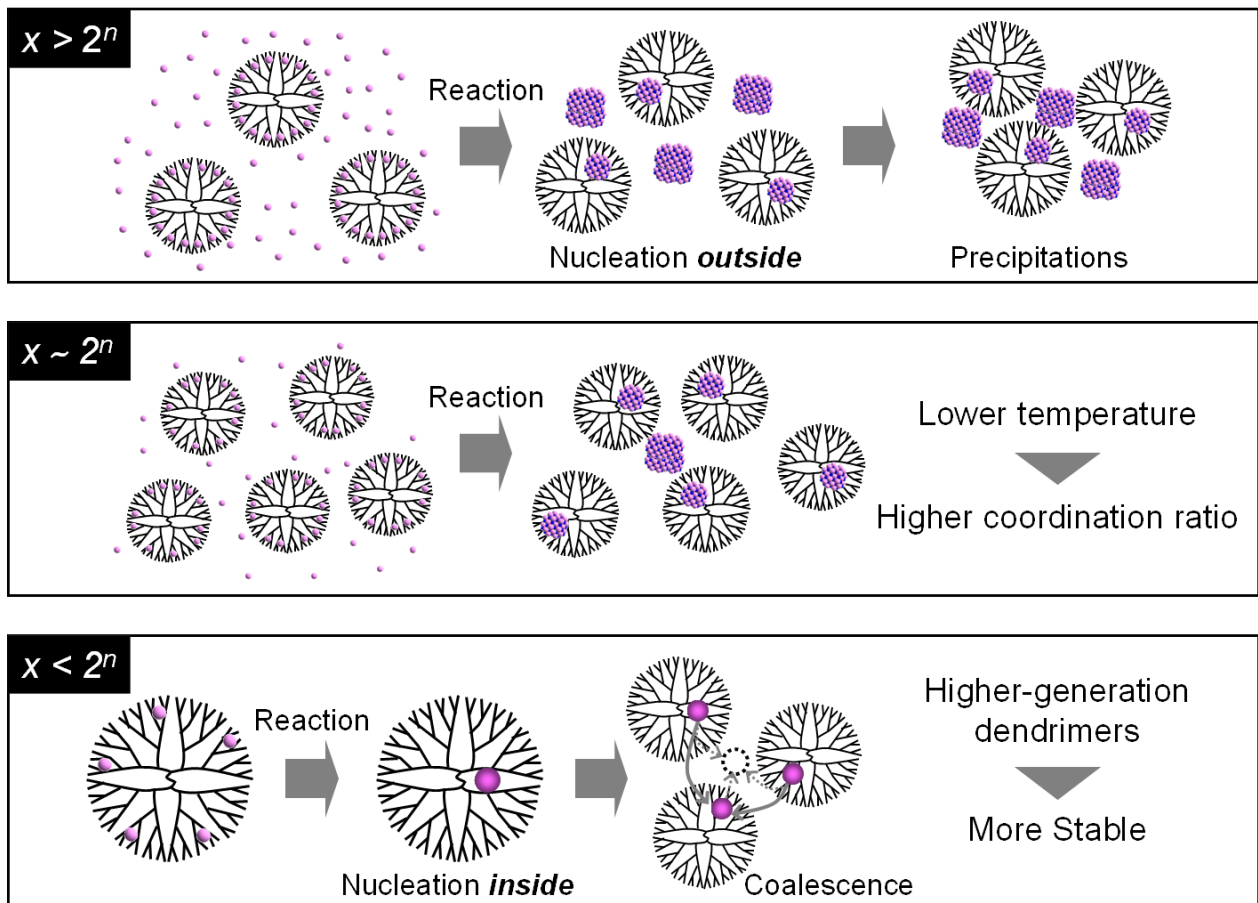


Fig. 6.

Conclusions

We synthesized CdS QDs by using hydroxyl-terminated PAMAM dendrimers as templates

and investigated the effects of Cd^{2+} ion/dendrimer ratio (x), dendrimer generations, and temperatures on the resultant CdS QDs. Through a detailed analysis, we found that setting x to be less than 2^n enables most of the nucleation to occur inside the dendrimers, which leads to stable dendrimer-encapsulated CdS QDs. We also demonstrated that higher-generation dendrimers can confine smaller QDs because they have denser surfaces. Further, lowering the temperature is a simple and effective approach to enhance the coordination ratio at equilibrium and to decrease nucleation outside the dendrimers, although it slows the coordination rate. In this manner, enhancing the coordination ratio of Cd^{2+} ion with dendrimers is the key for synthesizing stable dendrimer-encapsulated CdS QD suspensions, and is possible by changing x , generations of dendrimers, and temperatures to control particle sizes and photoluminescence colors.

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Supporting Information Available: The details of the calculation of the coordination ratio of Cd^{2+} ions at equilibrium are described in Supporting Information. Comparison of PL intensity between

CdS QDs with G6-OH PAMAM dendrimers and those with the SPP-protected dendrimers, PL images of CdS QDs taken with a digital camera (Canon) within G6-OH dendrimers for various values of x illuminated with a UV lamp ($\lambda_{\text{ex}} = 356 \text{ nm}$) in a dark room, coordination ratio of $[\text{G6-OH} + x \text{ Cd}^{2+}]$ solutions, and PL spectra of SPP-protected CdS QDs at various temperatures ($\lambda_{\text{ex}} = 374 \text{ nm}$) are also available free of charge via the Internet at _____.

FIGURE CAPTIONS

Fig. 1. UV-vis absorbance spectra of G6-OH CdS QDs for various values of x at room temperature.

Methanol was used to obtain the background spectrum.

Table 1. Average sizes obtained experimentally and the theoretical sizes of CdS QDs for various values of x . The average sizes were calculated from Fig. 1 using the empirical equation proposed by Peng et al. [34] and the theoretical sizes were calculated by the approximation method proposed by Lippens and Lannoo [35].

Fig. 2. PL spectra of G6-OH CdS QDs for various values of x at room temperature. PL spectra were measured with the excitation wavelength at 374 nm.

Fig. 3. TEM images of resultant CdS particles with G6-OH dendrimers for (a) $x = 20$, (b) 50, and (c) 400.

Fig. 4. PL spectra of CdS QDs with PAMAM dendrimers having various generations for $x = 20$ (λ_{ex}

= 374 nm).

Fig. 5. PL spectra of G6-OH CdS QDs for $x = 50$ at various temperatures ($\lambda_{\text{ex}} = 374$ nm).

Fig. 6. Synthetic mechanism for CdS QDs encapsulated within Gn -OH dendrimers.

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Supporting Information for:

Synthesis and Photoluminescence Characterization of Dendrimer-Encapsulated CdS

Quantum Dots

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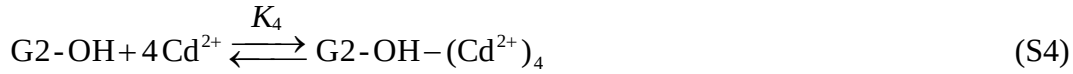
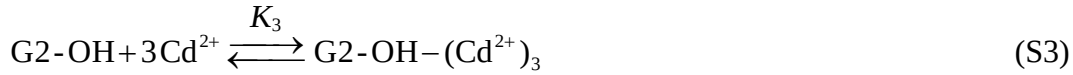
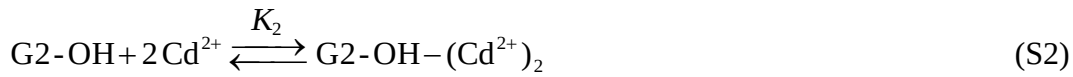
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Calculation of Cd^{2+} coordination ratio at equilibrium.

We calculated Cd^{2+} coordination ratio at equilibrium by using cumulative consecutive constants of Cd^{2+} -G2-OH coordination determined by Bard et al. [1] The reactions of Cd^{2+} complexes with G2-OH at 25 °C are expressed by the following equations:



$$K_1 = \frac{[\text{G2-OH}-\text{Cd}^{2+}]_{\text{eq}}}{[\text{Cd}^{2+}]_{\text{eq}} [\text{G2-OH}]_{\text{eq}}} = 1.3 \times 10^5 \text{ M}^{-1} \quad (\text{S5})$$

$$K_2 = \frac{[\text{G2-OH}-(\text{Cd}^{2+})_2]_{\text{eq}}}{[\text{Cd}^{2+}]_{\text{eq}}^2 [\text{G2-OH}]_{\text{eq}}} = 1.6 \times 10^9 \text{ M}^{-2} \quad (\text{S6})$$

$$K_3 = \frac{[\text{G2-OH}-(\text{Cd}^{2+})_3]_{\text{eq}}}{[\text{Cd}^{2+}]_{\text{eq}}^3 [\text{G2-OH}]_{\text{eq}}} = 3.3 \times 10^{12} \text{ M}^{-3} \quad (\text{S7})$$

$$K_4 = \frac{[\text{G2-OH}-(\text{Cd}^{2+})_4]_{\text{eq}}}{[\text{Cd}^{2+}]_{\text{eq}}^4 [\text{G2-OH}]_{\text{eq}}} = 4.6 \times 10^{14} \text{ M}^{-4} \quad (\text{S8})$$

where, K_i ($i = 1-4$) is the equilibrium constant, the square brackets [] represent the concentration, and the subscript eq denotes the equilibrium condition.

Material balances of Cd^{2+} and G2-OH are described as follows:

$$[\text{Cd}^{2+}]_{\text{initial}} = [\text{Cd}^{2+}]_{\text{eq}} + \sum_{k=1}^4 k \times [\text{G2-OH}-(\text{Cd}^{2+})_k]_{\text{eq}} \quad (\text{S9})$$

$$[\text{G2-OH}]_{\text{initial}} = [\text{G2-OH}]_{\text{eq}} + \sum_{k=1}^4 [\text{G2-OH}-(\text{Cd}^{2+})_k]_{\text{eq}} \quad (\text{S10})$$

where, the subscript initial denotes the initial condition. We solved simultaneous equations (S5–10) using Newton–Raphson method.

The coordination ratio α at equilibrium is obtained by the following equation:

$$\alpha = \frac{[\text{Cd}^{2+}]_{\text{initial}} - [\text{Cd}^{2+}]_{\text{eq}}}{[\text{Cd}^{2+}]_{\text{initial}}}$$

From the calculation result, for example, α at equilibrium of a 1 mM $[\text{G2-OH} + 4 \text{ Cd}^{2+}]$ ($[\text{G2-OH}]_{\text{initial}} = 1 \text{ mM}$, $[\text{Cd}^{2+}]_{\text{initial}} = 4 \text{ mM}$) solution was estimated to be 70%.

Reference

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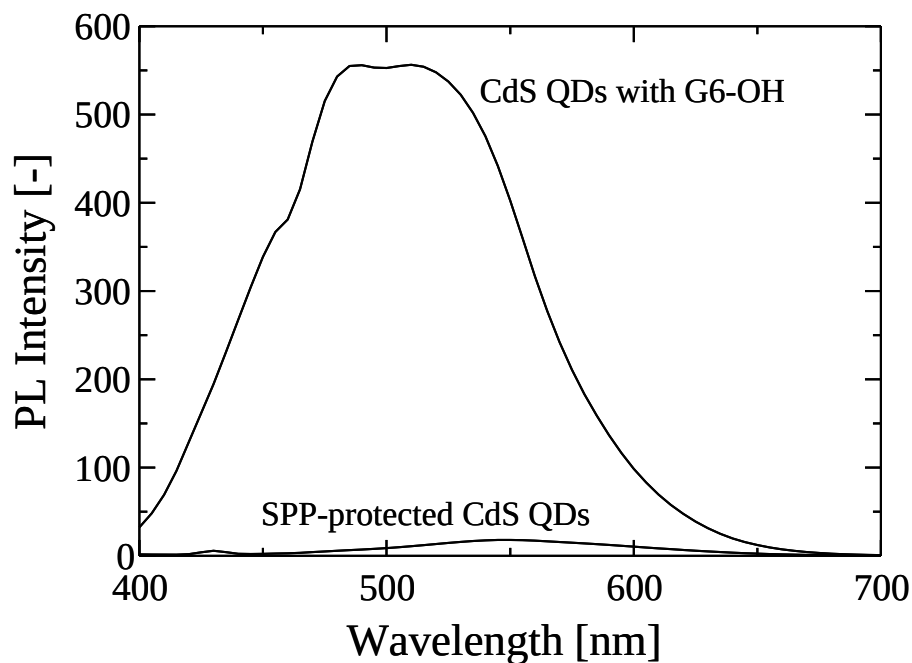


Figure S1. Comparison of PL intensity between CdS QDs with G6-OH PAMAM dendrimers and those with the SPP-protected dendrimers. The suspensions were prepared at room temperature. The concentrations of G6-OH and SPP were set to be 10 and 500 μM , respectively.

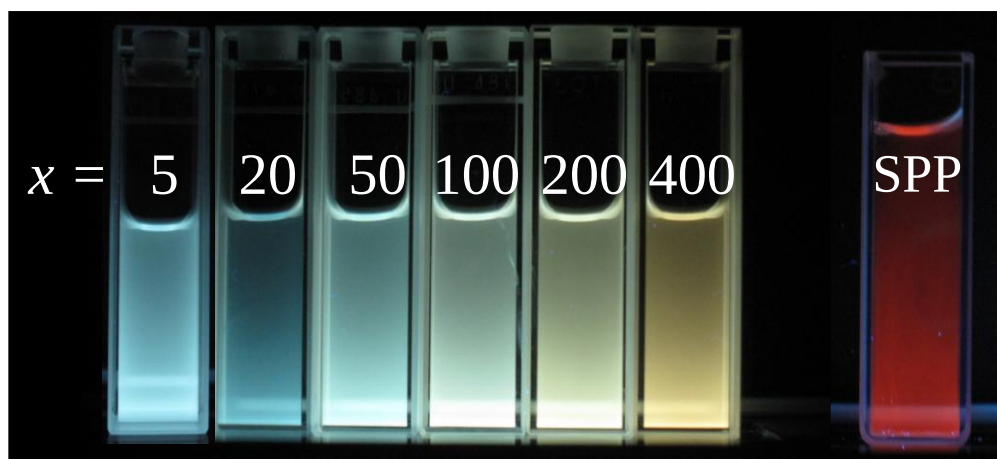


Figure S2. PL images of CdS QDs taken with a digital camera (Canon) within G6-OH dendrimers for various values of x illuminated with a UV lamp ($\lambda_{\text{ex}} = 356 \text{ nm}$) in a dark room.

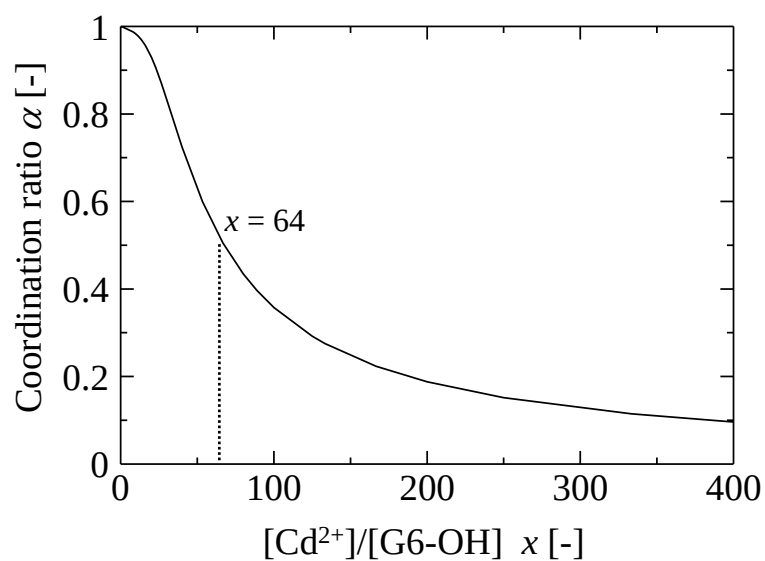


Figure S3. Coordination ratio of $[G6-OH + x Cd^{2+}]$ ($[Cd^{2+}]_{initial} = 0.5 \text{ mM}$) solutions.

The values were estimated according to “Supporting information: Calculation of Cd^{2+} coordination ratio at equilibrium.” The concentrations of G6-OH were converted to those of G2-OH by the following equation: $[G6-OH] = 16 \times [G2-OH]$.

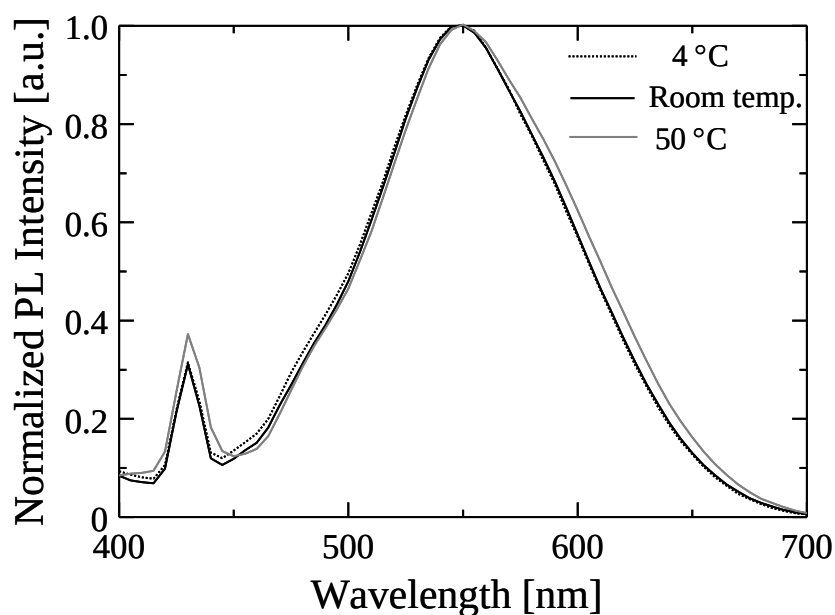


Figure S4. PL spectra of SPP-protected CdS QDs at various temperatures ($\lambda_{ex} = 374 \text{ nm}$).