Organic ligand and solvent kinetics during the assembly of CdSe nanocrystal arrays using infrared attenuated total reflection

Bosang S. Kim, a) Luis Avila, b) Louis E. Brus, b) and Irving P. Herman, b) c)

Materials Research Science and Engineering Center and the Columbia Radiation Laboratory, Columbia University, New York, New York 10027

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The self-assembly of amorphous three-dimensional arrays of CdSe nanocrystals is probed in real time using multiple-reflection, infrared attenuated total reflection spectroscopy by following the solvent and the organic ligands that passivate the nanocrystal surface. During the self-assembly of a 250 ML array from pyridine-capped CdSe nanocrystals in pyridine solvent, the solvent molecules evaporate in ~30–40 min and the pyridine-capping molecules leave the array very slowly, apparently limited by diffusion, with ~30±8% remaining after three days. © 2000 American Institute of Physics. [S0003-6951(00)00325-9]

Semiconductor nanocrystals have created great interest as possible components of complex materials because of their unusual properties, including their linear and nonlinear optical properties.1 The organic molecules sometimes used to “cap” the surface of semiconductor nanocrystals can passivate surface states, stabilize dispersions of the crystallites, and form an insulating layer that electrically isolates the particles.2 Three-dimensional arrays of CdSe nanocrystals have been self-assembled from solutions of these nanocrystals.3 The kinetics of array formation depends on the interactions of particles with each other, the solvent, and the surface. It also depends on solvent evaporation, which is very different for monolayered arrays and very thick three-dimensional (3D) arrays; this has been little studied. Many unanswered issues also remain about the stability of organic ligands that bind weakly to the nanocrystals, such as pyridine, particularly during self-assembly.

In this letter, the density of surface ligands and solvent molecules are followed during the self-assembly of CdSe nanocrystals passivated by pyridine and tri-n-octylphosphine oxide (TOPO), designated as CdSe/pyridine and CdSe/TOPO, respectively, to form amorphous arrays by using multiple-reflection, horizontal attenuated total reflection (ATR) spectroscopy. ATR permits probing of (1) the loss of solvent molecules during the early stages of drying, (2) the exchange between the ligands capping the CdSe with solvent molecules, and (3) the possible desorption of these capping ligands during the later stages of drying. Since pyridine binds only weakly to the CdSe nanoparticle, mostly at surface Cd,4,5 it has been thought that all of the pyridine rapidly desorbs during the formation of an array from CdSe/pyridine, leaving bare nanocrystals, at least for monolayer arrays at 150 °C.6 It is seen here that in thicker films the nanocrystals are still at least partially capped by the weakly binding pyridine, even after several days of drying.

CdSe nanocrystals 3.0 nm in diameter (with a standard deviation of 5%) and passivated with tri-n-octylphosphine oxide (TOPO, CdSe/TOPO) were synthesized according to the method of Murray et al.1 In many runs, these TOPO surface ligands were replaced by pyridine (CdSe/pyridine) by refluxing the nanocrystals in pyridine three times, each time for 3–5 h at 60 °C,7 prior to self-assembly.

A concentrated 0.1 cc drop of capped CdSe nanocrystals in solution was placed on the top surface (7 mm×68 mm) of the ZnSe prism, initially producing a ~210-μm-thick layer. ATR spectra were taken periodically during drying, after which a disordered, amorphous array of ~0.85±0.20 μm thickness formed—as determined by profilometry. This corresponds to about 250 ML of CdSe nanoparticles. These measurements were performed using a ZnSe GRASEBY-SPECAC horizontal ATR accessory mounted on a PE-1000 Paragon Fourier-transform infrared (FTIR) spectrometer. Each spectrum was averaged over four scans in 16 s, and taken with a resolution of 4 cm−1 and weak apodization. Peak heights were determined from Gaussian fits. Unless otherwise noted, all drying experiments were conducted at ambient temperature.

Figure 1 shows the ATR spectra near the 1445.2 cm−1 peak of pyridine bound to a CdSe nanoparticle, for (1) arrays...
of CdSe/pyridine nanocrystals initially in pyridine solvent dried for 2, 15, or 60 min, (2) a dried array of CdSe/TOPO nanocrystals initially in hexane, and (3) pure TOPO. This C–C, C–N stretching mode of pyridine bound to the CdSe nanocrystal at 1445.2 cm$^{-1}$ appears at 1436.1 cm$^{-1}$ in the neat pyridine. This upshift for bound pyridine is reasonable given the frequency upshift of pyridine adsorbed on Au.

Solvent evaporation during array formation with CdSe/TOPO nanocrystals dispersed in hexane solvent was too fast to be followed with this method. No changes in the frequency or height of the TOPO peaks were observed after the hexane evaporated, showing that the TOPO remained bound to the surface; this is expected because TOPO has a very low vapor pressure. The small TOPO peaks at 1377.4 and 1466.5 cm$^{-1}$ in the CdSe/pyridine array suggest that even after ligand exchange with pyridine the CdSe is still partially capped by TOPO. Comparing the IR absorbances in CdSe/pyridine and CdSe/TOPO arrays, $\sim$11% of TOPO molecules remain after ligand exchange with pyridine. This is consistent with measurements of $\sim$10–15% of the sites remaining capped by TOPO made using other methods.\(^8\)

The spectra in Fig. 1 show that during the self-assembly of nanoparticles in solution the absorbance peak at 1436.1 cm$^{-1}$ decreases to zero, while the pyridine-on-CdSe peak at 1445.2 cm$^{-1}$ first increases from roughly zero and then decreases. The inset in Fig. 2 follows these peaks in greater detail for CdSe/pyridine and CdSe/TOPO dispersed in pyridine. The removal of solvent in the probed region is indicated both by the decrease in 1436.1 cm$^{-1}$ peak (loss of solvent molecules) and the increase in the 1445.2 cm$^{-1}$ peak (increase of the number of nanoparticles). Complete drying of the solvent occurs in $\sim$30–40 min. The bound pyridine peak initially increases, reaches a maximum at $\sim$15 min, and then slowly decreases. Since the absorbance of bound pyridine in CdSe/TOPO in pyridine traces are the same as those for CdSe/pyridine in pyridine (Fig. 2), surface exchange of TOPO for pyridine apparently occurs very fast on these time scales. (The CdSe/TOPO particles were dispersed in the pyridine solvent within $\sim$2 min of deposition on the ATR crystal and the first FTIR scan.) TOPO peaks still remain; a small fraction of TOPO is still bound to the CdSe nanoparticles and most of it is interstitial and cannot evaporate.

The main part of the Fig. 2 tracks the desorption of bound pyridine to longer times, up to eight days; each trace is normalized to unity at its maximum, accounting for the optical thickness of the sample. Sample (a) was dried for 3 h under nitrogen, in vacuum for 1 h, and then under argon for 3 days—each at ambient temperature; this was followed by drying in vacuum at 50 ± 10°C for an hour. In (b), (c), and (d) the samples were dried in nitrogen for 30 min and then in argon, at ambient temperature. In each case at 20 °C there is a relatively rapid exponential decrease in absorbance with a time constant of $\sim$208 min, followed by a very slow decrease, with a time constant of very roughly 170 h. After three days of drying the absorbance peak decreases to $\sim$30 ± 8% of its peak value, which means that this fraction of capping pyridine molecules remain bound. If this peak absorbance corresponds to complete surface capping by pyridine, after three days of drying $\sim$30±8% of the pyridine sites remain occupied. The rate of drying does not seem to depend on the gas environment, which is expected since there is essentially no pyridine vapor above the layer during drying (to which only $\sim$2% of the total surface of the nanocrystals are directly exposed). Drying at elevated temperature led to a much more rapid desorption of surface-bound pyridine.

The pyridine peaks at 1010, 1036, and 1600 cm$^{-1}$ in CdSe/pyridine and 991, 1031, and 1580 cm$^{-1}$ in pyridine solvent (the symmetric and asymmetric ring breathing, and C–C, C–N stretching modes, respectively\(^8\) inset to Fig. 1) were also followed as a function of drying time. The normalized time dependences of bound and neat pyridine intensity were the same for each of these modes as for those in Fig. 2. All four modes (including that at 1445.2 cm$^{-1}$) were followed initially with a resolution of 4 cm$^{-1}$ and later with 1 cm$^{-1}$ resolution to look for a spectral shift of bound pyridine during drying. None was seen. In contrast, Ref. 11 reported that the $\approx$2850 and 2920 cm$^{-1}$ peaks of the dodecanethiol cap Ag nanocrystals in a dried array are red-shifted significantly from those of the nanoparticles in solution (by 7 cm$^{-1}$ for the latter peak).

The penetration depth in ATR is $\lambda/2\pi(\sin^2 \theta - n_2^2/n_1^2)^{1/2}$, where $\theta$ is the incident angle, $n_2 = n_2/n_1$, $\lambda$ = $\lambda_{\text{vacuum}}/n_1$ ($\lambda$ is the wavelength in the prism), $n_1$ is the refractive index of the ZnSe prism (2.4 at 6.9 μm), and $n_2$ is the refractive index of the sample (1.507 at 6.9 μm for neat pyridine). $\theta$ is fixed at 45° by the prism and coupling geometry. At this angle internal reflection occurs when the external media has $n_2 \ll 1.7$. The beam reflects $\sim$6–7 times from the surface covered with the array, and this does not change during drying. Right after the drop is placed on the ZnSe prism, the IR beam probes about 0.2 $\lambda$ or $\sim$1.4 $\mu$m into the solvent, and sees a very low density of nanoparticles (cumulatively $\sim$0.7% of the total). As the solvent evaporates, the density of nanoparticles in the depth probed increases, accounting for the initial increase in the bound pyridine peaks. The absorbance at the maximum of the adsorbed pyridine peak in Fig. 2 approximately equals that of the free pyridine peak, so that at that time there are roughly equal quantities of bound and solvent pyridine being probed if their absorption cross sec-
ations are equal. The refractive index of the array is then \( n_{\text{array}} = 1.79 \) [using effective medium theory with volume averaging of dielectric constants, \( \varepsilon_{\text{CdSe}} = 9.7 \), assuming face-centered-cubic (fcc) close packing of the nanocrystals with 74% packing density, with pyridine filling the interstitial regions]; it is \( n_{\text{array}} = 1.4 \) for dried arrays, with all solvent pyridine removed, 30% of the bound pyridine remaining, and unchanged interdot separations. Therefore, the penetration depth increases beyond 1.4 \( \mu m \) during drying and is always \( \geq \) the maximum thickness of the dried array (\( \sim 0.85 \mu m \)). Since \( n_{\text{array}} > 1 \), the infrared beam likely probes the entire depth of the dried array and is totally internally reflected at its outer surface. The total optical path (summed over all the internal reflections) does not change after the absorbance of the bound pyridine peak reaches its maximum.

Figure 3 follows the absorbance of bound pyridine for the original \( \sim 0.85 \mu m \)-thick film (“thick” array) and for films that were formed with the same volume drop with CdSe/pyridine nanocrystal concentrations that were \( \sim 1/3 \) (“thinner” array, \( \sim 0.35 \mu m \)-thick) and \( 1/10 \) (“thinnest” array, \( \sim 0.08 \mu m \)-thick) as large. The thinner the film, the faster the initial decay of the bound peak pyridine peak; the time constant is 19 min for the thinner array and less than the 2 min instrumental limit for the thinnest array; it was 208 min for the thick samples. Also, the thinner the sample, the lower the long-term fractional level of bound pyridine (\( \sim 0.56, 0.19, 0.06 \times \) peak value, respectively, at 5 h). Both of these observations are consistent with diffusion out of the film with a corresponding time constant \( \tau \propto (\text{thickness})^2 \) and not desorption—as being the rate limiting step.

Figure 3 also shows the ATR during the drying of CdSe/pyridine in pyridine when a drop of CdSe/TOPO nanocrystals in hexane was placed on the thinner sample after the pyridine solvent had evaporated, at 17 min (see arrow). This results in an increased level of bound pyridine, presumably because the fast drying and very stable CdSe/TOPO array acts as a barrier for pyridine diffusion out of the array (as well as making the film thicker). When the diffusion barrier is applied, the rapid decrease in absorbance of bound pyridine stops, while this decrease continues in the corresponding cases with no diffusion barrier. The ATR spectra also showed increased level of TOPO in the film, which proves that the IR beam probes the entire thickness of the films, even for the thick film (not shown).

In conclusion, ATR is seen to be a quantitative real-time probe of several important features of nanocrystal self-assembly. For the self-assembled 250 ML CdSe nanocrystal arrays, the pyridine solvent evaporates in \( \sim 30 - 40 \) min and \( \sim 30\% \) of the labile surface-bound pyridine molecules remain after three days. Diffusion seems to be the rate limiting step.

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13. The IR absorption cross sections for some molecules adsorbed on metal surfaces are larger than those of free molecules, as in: S. Badilescu, P. V. Ashrit, and V.-V. Truong, Appl. Phys. Lett. 52, 1551 (1988).