Crystallization of nanoparticles induced by precipitation of trace polymeric additives

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Orthogonal to guided growth of nanoparticle (NP) crystals using DNA or supramolecules, a trace amount of polymeric impurities (<0.1 wt.%) leads to reproducible, rapid growth of 3D NP crystals in solution and on patterned substrates with high yield. When polymers preferentially precipitate on the NP surfaces, small NP clusters form and serve as nuclei for NP crystal growth in dilute solutions. This precipitation-induced NP crystallization process is applicable for a range of polymers, and the resultant 3-D NP crystals are tunable by varying polymeric additives loading, solvent evaporation rate, and NP size. The present study elucidates how to balance cohesive energy density and NP diffusivity to simultaneously favor nuclei formation energetically and kinetic growth in dilute solutions to rapidly crystallize NPs over multiple length scales. Furthermore, the amount of impurities needed to grow NP crystals (<0.1%) reminds us the importance of fine details to interpret experimental observations in nanoscience.
Cryotization is a ubiquitous process seen in most, if not all, classes of matter. Nanoparticles (NPs) are ideal systems for visualizing and understanding cryotization processes and can serve as building blocks for new classes of materials. A dazzling array of NP crystals has been achieved by engineering complimentary interactions, e.g., by attaching DNA ligands or proteins. To achieve highly ordered NP crystals, adequate system mobility must be maintained during the assembly process. While the initial stages of the nucleation process are still open to debate, it is commonly accepted that the critical nucleus size depends on the balance between the NP/solvent interfacial interactions and the NP/NP cohesive energy stored within the nucleus. Nuclei are energetically stable at elevated NP size, and the strength of the ligand pair interactions. Different strategies have been employed to expand the processing window for NP cryotization, including controlled solvent evaporation, interface-mediated assembly, emulsion-based assembly, and slow cooling of concentrated supersaturated solutions. However, it remains challenging to control the cryotization kinetics so that high-quality NP crystals can be rapidly and reliably fabricated.

Precipitating agents such as salts, solvents, and polymers have been used to drive and accelerate the cryotization of small molecules and proteins. These additives can alter the molecular interactions to either stabilize the intermediate phase or to reduce the solubility of the crystallizing species. Adapting this concept for NP cryotization may lead to control over the assembly kinetics and pathways, but has not yet been experimentally explored.

Here, we hypothesize that by introducing polymers with poor solubility, NP surfaces may be the preferred precipitating sites and the adsorbed polymers gradually change the NP surface chemistry. To minimize unfavorable NP-solvent interactions, NPs form small clusters, which subsequently act as nuclei for the formation of NP crystals. The broad and the sharp diffraction peaks were deconvoluted to probe the PGNP crystal growth, as detailed in Supplementary Methods (Supplementary Fig. 6). A series of polymers were employed as precipitants with nonfavorable interactions with toluene and the absence of 3D NP crystals; only 2D superlattices were produced (Supplementary Fig. 6), consistent with previous results. Interestingly, we found that polymers dissolved from a polyolefin centrifuge tube were the best precipitants, leading to rapid and reliable growth of high-quality 3D PGNP crystals. The polymer additive, hereafter called PP/PE, was chemically identified as a low molecular weight, noncrystalline polypropylene (PP) with ~10% of polyethylene (PE) as detailed in Supplementary Methods (Supplementary Fig. 6), consistent with previous results. For homopolymers to act as precipitants, their molecular weight needs to be fairly small, typically less than ~8 KDa. See detailed information in Supplementary Table 1. Considering the polymer chain conformation in solution during precipitation and inter-NP distance at dilute concentrations, the bridging mechanism where a polymer chain acts as a linker to bring NPs together is highly unlikely. Rather, the results are consistent with the hypothesized mechanism shown schematically in Fig. 1a, where the adsorption footprint of the polymeric precipitant remains small enough to gradually modulate the NP-solvent interactions.

Au PGNPs (diameter = 3.8 ± 0.4 nm), denoted by (5.3k) Au-PS0.60 NPs (with PS number-average molecular weight Mn = 5300 g mol⁻¹, grafting density Σ = 0.60 chains nm⁻²), were dispersed in toluene at ΩNP = 0.1 vol%. PP/PE was added at ΩPP/PE = 0.08vol%. As the toluene evaporated, small-angle X-ray scattering (SAXS) profiles showed multiple well-defined diffraction peaks with peak positions at q_p/qp = 1: √2: √3: 2: √5: √7, consistent with a body-centered-cubic (bcc) structure (Fig. 1c). In situ SAXS was carried out to monitor the PGNP assembly during solvent evaporation (Fig. 1d and Supplementary Fig. 10). The solution contained Au PGNPs, denoted by (1.7k) Au-PS0.70 NPs (with PS Mn = 1680 g mol⁻¹ and Σ = 0.70 chains nm⁻²) and PP/PE at volume fraction ΩPP/PE = 0.08%. At ΩNP ∼ 23% (t = 11 min), the scattering profile showed a broad peak, corresponding to an average interparticle distance of 6.8 nm. This peak originated from the poorly ordered PGNP clusters since the average PGNP separation distance is ~7.8 nm at this ΩNP. At t = 15 min, an additional sharp diffraction peak appeared at q_p = 0.10 Å⁻¹. The peak intensified with a second-order peak appearing at q_p/qp = √2, confirming the formation of well-ordered PGNP assemblies. The intensity of the diffraction peaks increased and shifted to higher q positions as the solvent evaporated. The final interparticle distance was 6.1 nm and the estimated domain size, derived from the peak width, was ~450 nm. Thus, the in situ solution SAXS studies show a reduction in the average inter-NP distance from 7.8 nm in solution to 6.8 nm in poorly ordered NP clusters, and 6.1 nm in NP crystals. The broad and sharp diffraction peaks were deconvoluted to probe the PGNP crystal growth, as detailed in “Methods.” As the solvent evaporated, an increasing fraction of PGNPs assembled into ordered structures. NP crystals grew rapidly first and then growth slowed down as shown in Fig. 1e. This can be attributed to the depletion of PGNPs and the decrease in the PGNP diffusivity as the solvent evaporated and again highlights the importance of crystal growth within dilute solutions.

By using a small amount of solution (<5 µL), the self-assembly process was quenched at an early stage. Both NP assemblies with...
high crystalline order and poor order were seen using TEM. Image analysis confirmed that NP assemblies less than 30 nm in diameter consistently showed poor structural ordering (Fig. 1f), while larger NP clusters exhibited crystalline structures. Thus, the poorly ordered NP clusters likely still contain polyolefin precipitants, but nevertheless can act as nuclei for subsequent NP crystallization. Recent work showed that an amorphous, liquid-like intermediate phase exists within the crystallization process, in agreement with our experimental observation. Further SAXS and TEM analysis confirmed that the inter-NP distance remained the
same between the final PGNP assemblies with and without added polyolefins. In conjunction with the fact that the ligand–ligand interaction is much more favorable than the ligand-polymer interaction, it is reasonable to conclude that polyolefins were excluded during subsequent crystallization process.

**Effects of polymer precipitates and solvent evaporation on crystal growth.** The amount of polyolefin added affects the PGNP crystal size. Figure 2a shows representative TEM images of the crystals at different PP/PE loading ($\varnothing_{PP/PE}$). At low $\varnothing_{PP/PE}$ (<0.01%), layered 2D superlattices with small grain sizes were seen. 3D PGNP superlattices formed when $\varnothing_{PP/PE}$ reached 0.05%. At $\varnothing_{PP/PE}$ ~ 0.5%, PGNPs assembled into larger crystals with multiple grains. As more polyolefin was added, there was an increase in the average crystal diameter and size distribution (Fig. 2b and Supplementary Fig. 11). Systems of larger assemblies have a smaller total surface area, reducing the nonfavorable interactions between the PGNP surface and polyolefins. The fact that more polyolefins led to larger crystals suggests that after the polyolefins are excluded from the bulk crystal, they interact with the crystal surface and control the final crystal size. This is consistent with the unchanged interparticle distance with polymeric additives.

The polymeric ligands on PGNPs are critical for the nucleation and growth of PGNP crystals. In control experiments using oleylamine (OAm) stabilized Au NPs, no ordered 3D crystals were observed (Supplementary Fig. 12). Coassembly of PGNPs and OAm-NPs resulted in crystallization of only the PGNPs, with the OAm-stabilized NPs excluded (Supplementary Fig. 13). The spontaneous phase separation between NPs with two different ligand chemistries confirmed that the PS ligands play an important role in the PGNP crystal growth process. These results support our hypothesis that the polyolefin preferentially precipitates on NPs with polymeric ligands, and thus modulates PGNP nucleation and crystallization.

PGNP crystal growth depends on the solvent evaporation process. Grazing incidence/transmission SAXS (GI(T)SAXS) was performed to monitor the growth of PGNPs on a Si substrate during solvent evaporation. As a transmission technique, GI(T) SAXS probes entire assemblies. The initial $\varnothing_{PP/PE}$ was set to 0.15%. The final scattering patterns of the dried samples are shown (Fig. 2c). At a solvent evaporation rate of ~1 μL min$^{-1}$, diffraction spots appeared at a beam incidence angle of $\alpha = 0.8^\circ$. The peaks are assigned using the diffraction patterns of a bcc crystal lattice with zone axes of [111], [100], [011], and [311] (Supplementary Fig. 14). The presence of defined peaks indicates that, throughout the whole sample, the bcc crystals formed with (110) planes aligned parallel to the substrate (Fig. 2d). Thus, the crystals formed during solvent evaporation are strongly biased by the substrates due to the low cohesive energy of PGNP clusters. There was no diffraction spot when $\alpha = 0.14^\circ$ (Supplementary Fig. 15), meaning PGNP crystals were not formed directly at the Si substrate/PGNP interfaces, again differentiating with the previously reported NP interfacial assembly mechanism.

When the solvent evaporation rate was increased to 2.5 μL min$^{-1}$, very few ordered assemblies formed, indicating that the PGNPs did not have enough mobility to organize into crystals. On the other hand, when the solvent evaporation rate was decreased to 0.7 μL min$^{-1}$, a combination of substrate-oriented crystals, 2D layers, and randomly oriented crystals formed (Supplementary Fig. 16), and the resulting scattering patterns were a combination of diffraction spots and vertical lines. Layered superlattices are typically found when the PGNPs assemble at interfaces without polyolefins. They are formed due to capillary forces and preferential lateral diffusion. As the PGNP flux was low at low solvent evaporation rates, PGNPs close to the solvent drying front were able to stack at the droplet-substrate interfaces to form 2D layers, while others...
continued to grow into 3D PGNP crystals. As the PP/PE loading was increased, however, the precipitant overwhelmed the substrate effect, and randomly oriented crystals formed. A phase diagram is plotted to map out the morphologies formed with varying amounts of polymeric additives and solvent evaporation rates (Fig. 2e).

With a controlled PP/PE amount and solvent evaporation rate, single-crystalline superlattice clusters with preferred orientation formed exclusively upon solvent evaporation (<20 min). Single crystals of (5.3k) Au-PS$_{60}$ NPs formed at $\phi$$_{PP/PE}$ ~ 0.1% with a solvent evaporation rate of 0.8 $\mu$L min$^{-1}$. Multiple TEM images are provided to demonstrate the high-yield crystal formation. Each of the crystals has distinctive rod-like shape and clear bcc packing, supported by the diffraction spots observed in fast Fourier transforms of the TEM images (Supplementary Fig. 17).

**Electron tomography and reconstruction on a single PGNP crystal.** Scanning transmission electron microscopy (STEM) tomography was performed to characterize the 3D spatial organization of the PGNPs in a single crystal formed on a carbon substrate$^{24}$. The reconstruction method and additional details can be found in the “Methods” section. The reconstructed cluster shows a 3D disk shape composed of stacked PGNP layers. A slight decrease in the number of PGNPs in each layer, from the bottom to the top, is observed. Within the reconstructed crystal, the layers closer to the top and the bottom (substrate) planes have more defects and grain boundaries (Fig. 3a). Figure 3b shows the two base plane layers plotted in different colors, demonstrating a locally distorted hexagonal lattice in each plane. The registration between the layers is predominantly bridge sites, and thus is consistent with bcc stacking and the base plane is (110). The in-plane orientation, i.e., the (001) direction, is mostly horizontal as demonstrated, but varies close to the crystal edge. As shown in the enlarged area in Fig. 3b, the orientation is rotated by roughly 60° or ~60°. Similar registration is also seen in base planes near the top, where a hollow site stacking can be observed near the edge (Supplementary Fig. 18). PGNP organization in these regions thus probably reflects the structural transition from hexagonal packing to a bcc lattice during crystal growth. The fact that such organization is more prominent at edges and top layers indicates that the crystal grows from the center to the edges. Out-of-plane structure is examined by analyzing PGNP stacking in the vertical direction. Figure 3c is a slice cut that lies vertically in the crystal with two consecutive PGNP layers depicted in different colors. The figure on the right shows the slice cut from a top-down view. The layers are identified as bcc (001) planes to reflect the majority orientation. Their horizontal are the [−110] direction and their vertical are the [110] direction. The image in the middle depicts a projection of one of the layers, which clearly shows the square lattice of the bcc (001) plane. Based on the structural characterization, we can confirm that the structure of crystals formed on carbon substrates is bcc, and thus identical to the structure of crystals formed in solution. The crystal orientation is biased by the surface yielding (110) base planes. The defects present on the crystal’s surface and its rounded shape reveal that the crystal growth is limited by PGNP diffusion at the late stage of the solvent evaporation. This is not unexpected, considering the drastic viscosity change associated with increasing $\phi$$_{PP/PE}$. However, it opens up the possibilities to produce faceted crystals rapidly since cohesive energy density and crystal growth kinetics can be tuned independently.

**Pattern PGNP crystals on substrates.** PGNP crystal growth is most readily controlled via the solvent evaporation rate, but crystallization kinetics can also be modulated by changing the NP size. When 25 nm PGNPs were used, individual small clusters were found at $\phi$$_{PP/PE}$ ~ 0.05% (Fig. 4a). Decreased cluster size is attributed to the decreased diffusivity of larger particles according to the Stokes–Einstein equation: $D = \frac{k_B T}{6 \pi \eta r}$ (1), where $\eta$ is the dynamic viscosity and $r$ is the radius of a spherical particle. Dimers, trimers, tetramers, pentamers, and hexamers were all observed (Supplementary Fig. 19). Such geometries of metal nanostructures have received great attention in the past years because of their unique applications in photocatalysis, surface-enhanced spectroscopies, and nonlinear optics$^{35}$. Previously, specific clustering of two to six PGNPs has been a major challenge and was only realized through elaborate templating strategies$^{36−39}$. It is worthwhile to further explore the possibilities of separating these nanostructures with high purity.

Low-magnification TEM images demonstrate that the growth of 3D crystals is not a rare event and is highly reproducible. At $\phi$$_{PP/PE}$ ~ 0.05%, the sizes of the crystals can be fitted to a normal distribution with an average diameter of ~165 nm. The crystals are approximated as disks, and their sizes are described by diameter $d = 2\sqrt{area/\pi}$ (2). TEM images were further analyzed by Voronoi tessellation to visualize the corresponding occupied area of each cluster. Figure 4b shows the 2D map of Voronoi cells, which are color coded based on their area. The nucleation sites were randomly distributed. Fitting the linear model to the Voronoi cell area vs. the size of the corresponding crystals gives a slope of (6.8 ± 1.9) and an intercept of (6.3 ± 1.4) $\times 10^4$ nm$^2$ (Fig. 4d). The linear correlation between the two

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**Fig. 3** Electron tomography and reconstruction of a single bcc crystal composed of (5.3k) Au-PS$_{60}$ NPs on the substrate. PP/PE was added at $\phi$$_{PP/PE}$ ~ 0.1%. **a** 3D visualization of the crystal surface (see “Methods”) clearly illustrating its overall shape and the defective top layers. **b** Two base plane layers plotted in green and purple, demonstrating a distorted hexagonal lattice in each plane. The registration between the layers is predominantly bridge sites, as seen in the enlarged view on the right. Scale bars are 100 and 25 nm. **c** Two consecutive planes within a vertical slice plotted in green and purple. PGNPs in one plane sit at the center of the PGPN squares in the other plane. The square lattice of one of the bcc (001) planes is shown in the enlarged view in the middle. The image on the right shows where the cut lies in the top view. Scale bars are 50 and 25 nm.
suggests that a 2D self-assembly model can be applied. Once a nucleus forms, the surrounding area is depleted of PGNPs, and further growth of the crystal relies on the short-range diffusion and local rearrangement of PGNPs.

Furthermore, the capability to control the PGNP nucleation and growth process is promising for producing hierarchical assemblies. Patterned surfaces can be used as templates to control the location of NP arrays, taking advantage of various interactions between the NPs and substrates. As an example, PGNPs can be assembled into a square array of crystals by using templated carbon films (Fig. 4e). The commercially available carbon-coated TEM grids have orthogonally patterned vacuum holes of ~1 μm diameter and ~650 nm separation. The available spaces between the holes are comparable to the size of PGNP crystals formed. Here, the nucleation of PGNPs preferentially took place between the holes, and the crystals were thus organized into a square lattice (Fig. 4g, h). Crystallinity was maintained within individual clusters (Supplementary Fig. 20). At least two periodicities existed: the several nanometer inter-PGNP distance in each cluster, and the micrometer separation between clusters.

Within the patterned substrates, the PGNPs had two possible nucleation sites: either between two adjacent vacancies or at interstitial sites of the hole lattice (see blue dots in Fig. 4f). These two sites produced two distinct separation periodicities and cluster sizes, due to the difference in available space. The average diameters of the crystals were ~600 nm and ~900 nm, respectively.

We demonstrated that a properly chosen polymer precipitate and PGNP surface modification can effectively change NP/solvent interactions, leading to controlled nucleation of PGNPs in dilute solutions. The initial PGNP clustering occurs in solution due to the precipitation of poorly soluble polymers onto PGNP surfaces. The crystallization process can be further modulated by manipulating the solvent evaporation rate, NP size, and nucleation sites. These PGNP systems are ideal for future studies in crystallization at single-building-block level, since the NP assembly’s energetic driving force, kinetics, and pathways can be easily modulated. The growth of NP superlattice crystals within several minutes, as demonstrated here, removes an important roadblock toward NP-based composite materials. Lastly, these PGNP crystals are obtained with the presence of trace amount of impurities from plastic containers that are routinely used. The results shown here clearly underscore the importance of identifying any potential chemical impurities and their effects in the field of nanoscience, even at a trace amount.

Methods

Materials. Hydrogen tetrachloroaurate (III) trihydrate (HAuCl₄·3H₂O, Sigma-Aldrich, 99.995%, trace metals basis), OAam (Sigma-Aldrich, technical grade, 70%), tert-butylamine-borane complex (Aldrich, 97%), 1,2,3,4-tetrahydronaphthalene (tetralin, anhydrous, 99%), styrene (ReagentPlus, 99.995%, trace metals basis), OAm (Sigma-Aldrich, technical grade, 70%), and toluene (Sigma-Aldrich, >97% (HPLC)). Poly(ethylene oxide) (PE) was purchased from Aldrich. Polyisoprene (PI) and polystyrene (PS) were synthesized using standard Schlenk line techniques. 3.8 nm OAm-Au NPs were synthesized according to literature methods. The synthesis was carried out under a nitrogen atmosphere using standard Schlenk line techniques. Thiol-terminated PS (PS-SH) was synthesized using radical addition fragmentation transfer polymerization and end-functionalized by aminolysis. Au NPs grafted with PS-SH were prepared by a ligand exchange process as reported before. The solubility of Au NPs grafted with PS-SH was confirmed using TEM image of (1.7k) Au-PS0.70 NP crystals formed with PP/PE at ~0.05%. Scale bar is 200 nm.

Crystal size distribution plot, fitted with a normal distribution curve. Voronoi cell area vs. crystal size in 2D, fitted with a linear curve. (1.7k) Au-PS0.70 NP crystals characterized in c and d were formed with PP/PE at ~0.08%. Schematic illustration of the templated carbon films with orthogonally arrayed holes mounted on TEM grids. Two possible nucleation sites are highlighted with blue circles in f. (1.7k) Au-PS0.70 NP crystals formed at center sites of the hole lattice. PP/PE was added at 0.15%. Scale bars are 500 nm in f-h.

Fig. 4 Controlled PGNP crystallization on substrates. a PGNP clusters assembled from 25 nm Au NPs (PS = 6k) with PP/PE at 0.05%. Scale bar is 200 nm. b TEM image of (1.7k) Au-PS0.70 NP crystals formed with PP/PE at ~0.08% and Voronoi representation color coded by the Voronoi cell area in 2D. c Crystal size distribution plot, fitted with a normal distribution curve. d Voronoi cell area vs. crystal size in 2D, fitted with a linear curve. (1.7k) Au-PS0.70 NP crystals characterized in c and d were formed with PP/PE at ~0.08%. e Schematic illustration of the templated carbon films with orthogonally arrayed holes mounted on TEM grids. Two possible nucleation sites are highlighted with blue circles in f. (1.7k) Au-PS0.70 NP crystals formed at bridge sites between adjacent holes. h (1.7k) Au-PS0.70 NP crystals formed at center sites of the hole lattice. PP/PE was added at ~0.15%. Scale bars are 500 nm in f-h.

Synthesis of PGNPs. 3.8 nm OAm-Au NPs were synthesized according to literature methods. The synthesis was carried out under a nitrogen atmosphere using standard Schlenk line techniques. Thiol-terminated PS (PS-SH) was synthesized using radical addition fragmentation transfer polymerization and end-functionalized by aminolysis. Au NPs grafted with PS-SH were prepared by a ligand exchange process as reported before. Further details on synthesis and characterization are provided in Supplementary Information.
PGNP crystal growth upon solvent evaporation. Au PGNPs were dispersed in toluene at $\phi_{PGNP} = 0.1\%$. Polymer additives were dissolved in toluene and added to the PGNP solution at $\phi_{polymer}$ ranging from 0.01 to 0.5%. Upon solvent evaporation, samples deposited on Si wafers and TEM grids were used for GISAXS measurements and TEM analysis, respectively. For crystal growth on normal substrates, 50 µL of solution was drop-cast on 200 mesh Cu TEM grids or a Si substrate and dried after ~20 min. For a quick-drying experiment, less than 5 µL solution was drop-cast and the solution dried within 1 min. Solvent evaporation was done under ambient conditions. For crystal growth on templated TEM grids, slow solvent evaporation and relatively high polymer concentration ($\phi_{polymer} = 0.15–0.2\%$) are desired in order to get highly ordered crystals. The TEM grid was laid on a Si substrate and placed in a Telfon well with dimensions of 1.5 cm x 1.5 cm x 1.0 cm, and covered with a glass coverslip to slow down the solvent evaporation. Samples dried completely after 2 h.

Transmission electron microscopy. Au PGNPs and their assemblies were imaged using a FEI Tecnai 12 at an accelerating voltage of 120 kV.

Small-angle X-ray scattering. In situ solution SAXS experiments were performed at Beamline 11-BM (Complex Material Scattering) of National Synchrotron Light Source II (NSLS-II, Brookhaven National Laboratory). The SAXS data were collected on a Dectris 2M detector at a sample-to-detector distance of 2 m, using an X-ray beam with an energy of 13.5 keV (the corresponding wavelength $\lambda = 0.92\ Å$). Au PGNP solution was positioned so that the beam went through the solution. The solution was left dry under ambient environment and were exposed to X-rays every 30 s over a range of 0.001°–0.2°.

Static SAXS experiments were performed at beamline 7.3.3 at Advanced Light Source in Lawrence Berkeley National Laboratory. The X-ray wavelength is 1.24 Å. The scattered X-ray intensity distribution was detected using a high-speed detector, Pilatus 2M. Images were plotted as intensities (I vs. q, where $q = (4\pi\lambda/\sin(\theta))$, $\lambda$ is the wavelength of the incident X-ray beam, and $2\theta$ is the scattering angle).

Peak deconvolution. The scattering peaks were deconvoluted using a MATLAB command-line peak fitting program. The area of interest, containing the sharp diffraction peak at $q = 0.1\ Å^{-1}$ from PGNP crystals and the broad peak from PGNP clusters in the solution, was selected to fit with a two-peak Gaussian model. At least ten trial fits were performed and the one with the lowest fitting error was selected. Data points were collected as the integration ratio of the two deconvoluted peaks.

Grazing incidence/transmission SAXS. GIX(S)AXS experiments at an incident angle $\alpha$ of 0.1°–0.8° were conducted at beamline 8-ID-E at the Advanced Photon Source of Argonne National Laboratory. Samples were drop-cast on a 1 cm x 1 cm Si substrate with 100–500 µL solvent reservoir placed in the house-made solvent annealing chamber. The chamber was sealed and $N_2$ flow was purged through to control the solvent evaporation. The scattered X-ray intensity distribution was detected using a high-speed detector, Pilatus 1M. Samples were exposed at 7.35-keV radiation ($\lambda = 1.68\ Å$).

STEM tomography. The tomography experiment was performed using a FEI Titan microscope operated at 300 kV. The HHAADF-STEM images were acquired with a 10 mrad probe semi-convergence angle. The projections of PGNPs were recorded over the tilt range of −70° to 70°, with 2° step size and pixel size of 0.66 nm. The alignment was performed on IMOD (version 4.9.10), making use of fiducial markers and custom python scripts. Reconstructions were performed using inverse Winterbottom constructions of nanoparticle superlattices.

Data availability

The data that support the findings of this study are available within the paper and its Supplementary Information Files.

Code availability

Custom MATLAB codes for Voronoi construction are available from the corresponding author upon request. MATLAB package for peak fitting and deconvolution is an open-source package and available online: https://www.mathworks.com/matlabcentral/fileexchange/23611-peaksfit-m.

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References

1. Manoharan, V. N. Colloidal matter: packing, geometry, and entropy. Science 349, 1235751 (2015).
2. Ou, Z., Wang, Z., Luo, B., Luijten, E. & Chen, Q. Kinetic pathways of crystallization at the nanoscale. Nat. Mater. 19, 450–455 (2020).
3. Grzelczak, M., Vermaas, J., Furst, E. M. & Liz-Marzán, L. M. Directed self-assembly of nanoparticles. ACS Nano 4, 3591–3605 (2010).
4. Feng, M. A., Engel, M. & Talapin, D. V. Self-assembly of colloidal nanocrystals from intricate structures to functional materials. Chem. Rev. 116, 11220–11289 (2016).
5. Alivisatos, A. P. et al. Organization of nanocrystal molecules using DNA. Nature 382, 609–611 (1996).
6. Mirkin, C. A., Letsinger, R. L., Muñoz, R. C. & Storhoff, J. J. A DNA-based method for rationally assembling nanoparticles into macroscopic materials. Nature 382, 607–609 (1996).
7. Nykypanchuk, D., Maye, M. M., Van Der Lelie, D. & Gang, O. DNA-guided crystallization of colloidal nanoparticles. Nature 451, 549–552 (2008).
8. Mullin, J. W. Crystalization (Elsevier, 2001).
9. Hoover, W. G. & Ree, F. H. Melting transition and communal entropy for hard spheres. J. Chem. Phys. 49, 3609–3617 (1968).
10. Dong, A., Chen, J., Vora, P. M., Kikka, J. M. & Murray, C. B. Binary nanocrystal superlattice membranes self-assembled at the liquid-air interface. Nature 466, 474 (2010).
11. Bigioni, T. P. et al. Kinetically driven self assembly of highly ordered nanoparticle monolayers. Nat. Mater. 5, 265 (2006).
12. Lewis, D. J., Zorzoll, L. Z., Carter, D. J. & Macfarlane, R. J. Single-crystal Winterbottom constructions of nanoparticle superlattices. Nat. Mater. 19, 719–724 (2020).
13. Park, J. E. et al. Surfactant-assisted emulsion self-assembly of nanoparticles into hollow vesicle-like structures and 2D plates. Adv. Funct. Mater. 26, 7791–7798 (2016).
14. Zhuang, J., Wu, H., Yang, Y. & Cao, Y. C. Supercrystalline colloidal particles from artificial atoms. J. Am. Chem. Soc. 129, 14166–14167 (2007).
15. Jones, A. G. & Mullin, J. Programmed cooling crystallization of potassium sulphate solutions. Chem. Eng. Sci. 29, 105–118 (1974).
16. Sun, J.-K., Sobolev, Y. I., Zhang, W., Zhuang, Q. & Grzybowski, B. A. Enhancing crystal growth using polyethylene oxide solutions and shear flow. Nature 579, 73–79 (2020).
17. Kulkarni, A. & Zukoski, C. Depletion interactions and protein crystallization. J. Cryst. Growth 232, 156–164 (2001).
18. McPherson, A. Crystalization of macromolecules: general principles. in Crystallization in Enzymology, 112–120 (Elsevier, 1985).
19. Song, R.-Q. & Cölfen, H. Additive controlled crystallization. CrystEngComm 13, 1249–1276 (2011).
20. Sauter, A. et al. On the question of two-step nucleation in protein crystallization. Faraday Discuss. 179, 41–58 (2015).
21. Dunnet, A. C., Snellinger-O’Brien, A. M., Kaler, E. W. & Lenthoff, A. M. Patterns of protein–protein interactions in salt solutions and implications for protein crystallization. Protein Sci. 16, 1867–1877 (2007).
22. Akkora, P. et al. Anisotropic self-assembly of spherical polymer-grafted nanoparticles. Nat. Mater. 8, 354–359 (2009).
23. Kumar, S. K., Jouault, N., Benicewicz, B. & Neely, T. Nanocomposites with polymer-grafted nanoparticles. Macromolecules 46, 3199–3214 (2013).
24. Ye, X. et al. Structural diversity in binary superlattices self-assembled from polymer-grafted nanocrystals. Nat. Commun. 6, 10052 (2015).
25. Fernandes, N. J., Koerner, H., Gianneli, E. P. & Vaia, R. A. Hairpin nanoparticle assemblies as one-component functional polymer nanocomposites: opportunities and challenges. MRS Commun. 3, 13–29 (2013).
26. Choi, J. et al. Effect of polymer-graft modification on the order formation in particle assembly structures. Langmuir 29, 6452–6459 (2013).
27. Choi, J., Dong, H., Matysiaszewski, K. & Rockstall, M. R. Flexible particle array structures by controlling polymer graft architecture. J. Am. Chem. Soc. 132, 12537–12539 (2010).
28. Nie, Z. et al. Self-assembly of metal–polymer analogues of amphiphilic triblock copolymers. Nat. Mater. 6, 609–614 (2007).
29. Choueri, R. M. et al. Surface patterning of nanoparticles with polymer patches. Nature 538, 79–83 (2016).
30. Josten, E. et al. Superlattice growth and rearrangement during evaporation-induced nanoparticle self-assembly. Sci. Rep. 7, 1–9 (2017).
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Author contributions

T.X. conceived the idea and guided the project. Y.Q. performed studies contributed to Figs. 1, 2, and 4. E.Y. performed nanoparticle synthesis. A.d.S. performed STEM tomography in Fig. 3. C.L.A. performed GIWAXS and HT-GPC measurements under the supervision of Y.L. A.d.S., W.T., and P.E. performed tomography analysis.

Competing interests

The authors declare no competing interests.

Additional information

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