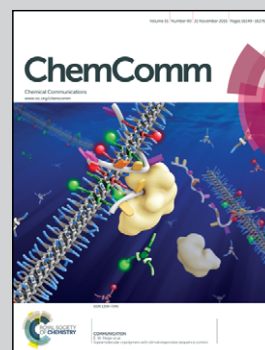


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Visualizing virus particle mobility in liquid at the nanoscale

The first time-resolved movies of virus particles in liquid were recorded at the nanoscale using *in situ* transmission electron microscopy.

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## Visualizing virus particle mobility in liquid at the nanoscale†

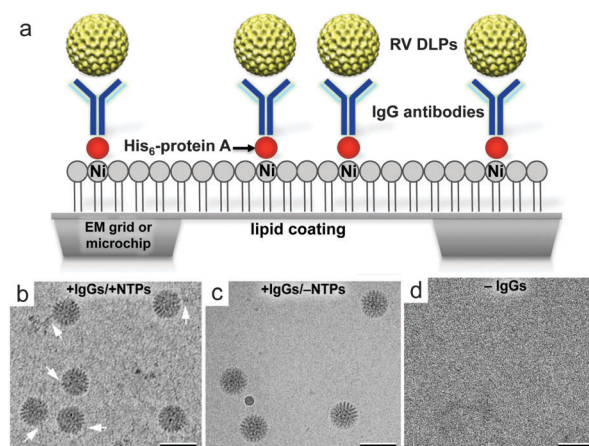
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**Currently, there remains a critical need to develop real-time imaging resources for life sciences. Here, we demonstrate the use of high resolution *in situ* imaging to observe biological complexes in liquid at the nanoscale. Using a model virus system, we produced the first time-resolved videos of individual biological complexes moving in solution within an electron microscope.**

Recent advances in the design and manufacture of nanometer-thick materials has spurred a new era in the imaging field.<sup>1,2</sup> When used in combination with microfluidic devices,<sup>3</sup> it is now possible to observe electrochemical processes,<sup>4</sup> the growth of nanoparticles,<sup>5</sup> drug delivery systems,<sup>6</sup> and live cellular targeting events<sup>7</sup> at unprecedented resolution. We refer to this new innovative mode of imaging as “*in situ*” microscopy as experiments can now be directly observed “*inside*” a transmission electron microscope (TEM). Although *in situ* imaging has gained a great deal of momentum in the materials science field, its application to biological systems is not as widespread. To address this issue, we developed a unique platform to directly observe movements in biological complexes using *in situ* TEM. Here, we present the first time-resolved movies and mathematical analysis of a model virus system moving in solution at the nanoscale. Specifically, we used purified rotavirus (RV) double-layered particles (DLPs) in our experiments since DLPs can be enzymatically activated *in vitro* by the addition of nucleotide triphosphates (NTPs) to initiate viral mRNA synthesis.<sup>8,9</sup>

To capture enzymatically active RV DLPs for downstream *in situ* image recordings, we employed a well-defined immunocapture tethering system. This system promotes the attachment of individual DLPs to surfaces coated with nickel–nitrilotriacetic acid (Ni–NTA)-containing lipid monolayers and adaptor proteins.<sup>10,11</sup>

The hydrophobic carbon chains of the lipid layers adhere readily to the naturally hydrophobic surface of carbon-coated EM grids or SiN microchips (Fig. 1a). We used lipid monolayers comprised of 25% (v/v) Ni–NTA lipids and 75% (v/v) 1,2-dilaurylphosphatidylcholine (DLPC) filler lipids (Avanti Polar Lipids). Adaptor proteins including His-tagged (His<sub>6</sub>) protein A (Abcam) and IgG antibodies against the viral protein (VP6) were added sequentially to the Ni–NTA-coated microchips. Aliquots (3  $\mu$ l of 0.01 mg ml<sup>-1</sup>) of His-tagged protein A in buffer solution containing 50 mM HEPES, pH 7.5, 150 mM NaCl, 10 mM MgCl<sub>2</sub> and 10 mM CaCl<sub>2</sub> were incubated for 1 minute on each EM grid or microchip prior to removing the excess solution by blotting with filter paper. Next, we added 3  $\mu$ l aliquots of VP6-specific guinea pig polyclonal antisera (#53963; 0.01 mg ml<sup>-1</sup>) prepared



**Fig. 1** Tethering DLPs to functionalized surfaces. (a) Schematic to illustrate the immunocapture procedure used to tether asynchronously transcribing RV DLPs to antibody (IgG)-decorated surfaces *via* protein A adaptors. (b) Transcribing DLPs tethered to EM grids or SiN microchips in the presence of VP6-specific IgGs showed varying lengths of associated mRNA transcripts (white arrows) in cryo-EM images. (c) DLPs prepared in the absence of nucleotides needed for transcription do not show associated mRNA in cryo-EM images. (e) EM specimens prepared in the absence of IgGs generally failed to recruit DLPs. Scale bar is 100 nm.

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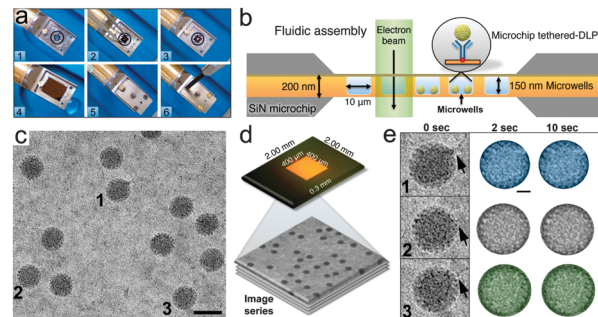
in buffer solution containing 50 mM HEPES (pH 7.5), 150 mM NaCl, 10 mM MgCl<sub>2</sub> and 10 mM CaCl<sub>2</sub>. Following a 1 minute incubation step, the excess solution was removed using a Hamilton syringe (Fig. 1a).

In parallel as negative controls, we also prepared (1) DLP samples lacking ATP, a nucleotide needed for enzymatic activation, and (2) protein A-coated microchips lacking IgG antibodies against VP6. All samples were imaged using cryo-EM to assess the specificity of the method. We found that within a 2 minute incubation period, the antibody-decorated microchips could recruit DLPs diluted to 0.1 mg ml<sup>-1</sup> in 50 mM HEPES buffer solution (pH 7.5) containing 150 mM NaCl, 20 mM CaCl<sub>2</sub> and 20 mM MgCl<sub>2</sub>, and NTPs (Fig. 1b). DLPs lacking nucleotides could still bind to antibody-decorated surfaces but viral mRNA transcripts were not present in the images (Fig. 1c). Microchips prepared without antibodies did not recruit appreciable quantities of DLPs (Fig. 1d).

To biochemically verify transcriptional competence of the viral particles, the purified DLPs were subjected to *in vitro* mRNA synthesis in Eppendorf tubes in the presence of [<sup>32</sup>P]-UTP for 30 minutes at 37 °C. Each reaction contained DLPs, NTPs, and [<sup>32</sup>P]-UTP (Fig. S1a and b +ATP; ESI†). Negative control reactions contained all transcription cocktail components except ATP (Fig. S1a, -ATP; ESI†). Radiolabeled mRNA products were only detected in the reactions containing complete transcription cocktail; no radiolabeled products were seen in the reaction lacking ATP. Therefore, our functional analysis confirmed that purified DLPs could be enzymatically activated. These results were consistent with previous experiments<sup>11</sup> and suggested that the system could be used to recruit and image active DLPs in solution. Therefore, we proceeded to record images of the virus particles using *in situ* TEM.

For *in situ* imaging experiments, we tethered enzymatically-active DLPs as described in Fig. 1, onto SiN microchips that were loaded into the Poseidon 200 specimen holder (Protochips, Inc.) (Fig. 2a and b). This tethering step served to limit long-range diffusion of the viral particles in solution during imaging. Each microchip contained central imaging windows (~30 nm thick) that are transparent to the electron beam and form an environmental chamber at the tip of the holder. The bottom chip that defined the base of the assembly contained a 400 × 400 micron array of microwells within the central imaging window.<sup>6</sup> Each microwell (10 × 10 microns) was etched down to accommodate a solution thickness of 150 nm. A metal faceplate, held in place by 3 brass screws, was placed on top of the assembled unit. The enclosed DLPs were examined using a FEI Spirit Bio-Twin TEM equipped with a LaB<sub>6</sub> filament and operating at 120 kV. We recorded at least five image series at time intervals up to 10 seconds (4 frames per second) while optimizing output conditions. Images were recorded on an Eagle 2k HS CCD camera employing low-dose conditions (~0.1 electrons per Å<sup>2</sup>) at a 60 000× magnification, yielding a final sampling of 5 Å per pixel. Image series proved reproducible between replicates.

To produce time-resolved movies of the viral particles in liquid, we selected from the image series representative DLPs having different lengths of RNA strands associated with their



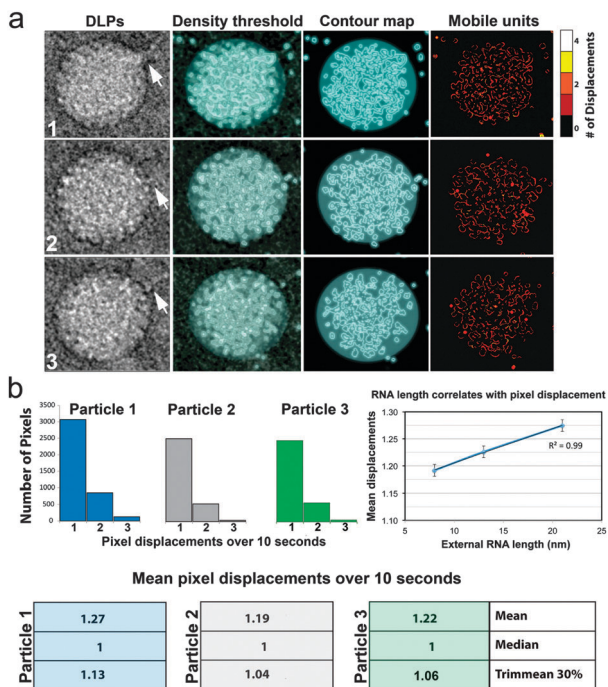
**Fig. 2** Microfluidic system for *in situ* TEM imaging of virus particles. (a) The microfluidic chamber was assembled by placing O-ring fittings in the empty tip of the holder (steps 1–3). A flat chip was placed over top of the wet specimen chip (step 4) and the assembly was covered with a metal face-plate held in place by 3 brass screws (steps 5 and 6). (b) Schematic of a cross-section through the fluidic chamber assembly positioned within the EM beam. Integrated microwells (10 × 10 μm) etched into the chip can each accommodate a solution layer of 150 nm thick. (c) Representative image of DLPs in liquid. (d) The base chip contained a transparent imaging window having array of microwells (400 × 400 μm). DLPs were tethered to the base chips and image series were recorded in the EM. (e) Representative DLPs (1–3) with viral mRNAs transcripts (black arrows). Width of individual panels is ~120 nm. Individual DLPs were contrast-inverted, masked by a diameter of ~80 nm and colored for visualization purposes. Please see Movies S1–S3, ESI.†

exterior shells (Fig. 2c and d). The selected particles were contrast-inverted and arbitrarily colored (blue, gray, and green) for tracking purposes (Fig. 2e). Images sequences of the DLPs within a 10 second time interval were compiled using the iMovie (Apple, Inc.) software package (Fig. S2 and Movies S1–S3, ESI†). The compiled movies revealed real-time movements of the DLPs in the surrounding liquid. We attributed the overall particle movements at the molecular level to Brownian motion, biological activity, beam-induced movement, and beam damage – or some combination of these factors. Another important consideration that can influence protein subunit mobility within individual DLPs is the antibody-tethering step. Although, we did not directly measure the collective forces imposed on the particles by the attached antibodies, we are continuing to investigate these interactions in our on-going experiments.

To better visualize mobile units in the DLPs, we segmented the strongest features in the particles by applying a density threshold filter of 3σ where σ is the standard deviation in the contrast values of each particle image.<sup>12</sup> By focusing on these strongest features, we could attenuate noise in the images due to variations in the liquid flow and to some extent, radiolysis. After applying the density threshold, we were able to visualize the highest electron scattering components within each particle image in the series (Fig. 3a). This density included both protein subunits and viral RNA (> 12 MDa) that constitutes a significant portion of the DLP (~50 MDa). Representative contour maps within a 10 second timeframe were compiled using the iMovie software package (Movies S4–S6, ESI†).

The compiled movies of the contoured regions showed the strongest visual differences in the individual virus particles and their associated mRNAs. This information provided a new





**Fig. 3** Visualizing and quantifying DLP mobility in liquid. (a) Representative DLPs (1–3) with associated viral mRNA strands (white arrows) were selected from an image series and subjected to a density threshold filter. Contour maps were generated based on a  $3\sigma$ -cutoff, highlighting pixel displacements in the particles. Cumulative differences in the mobile units of the viral particles were represented in heat maps using a scale of 0–4 to indicate the number of pixel displacements within a 10 second interval. (b) Histograms that represent the total number of pixels displacements within 10 seconds revealed non-uniform movements between the DLPs. Mean pixel displacement values are also summarized. The plot shows the correlation ( $R^2 = 0.99$ ) between the mean pixel displacements and the lengths of external RNA associated with each particle. Error bars represent the standard error values for each mean pixel displacement. Please see Movies S4–S6, ESI.†

time-resolved view of biological assemblies in liquid at the nanoscale. We also noted the presence of the viral mRNA transcripts ( $\sim 10$  Å in width) associated with the viral particles at the beginning and the end of the image series (Movies S4–S6, ESI†). The conserved presence of these fragile mRNA strands throughout the image series supports the idea that beam damage, or beam induced movements do not impose a stronger influence on the observed particle mobility than Brownian motion or biological activity.

To quantify the pixel movements in the particle images over time, the binary thresholded images were combined into time series. The temporal derivative of each pixel returned values of 1 when a pixel changed from empty to full, and  $-1$  when the pixel changed from full to empty. Summing the absolute value of each pixel's temporal derivative returned a value that represents how frequently each pixel changed value. When applied to the entire time series, we could quantify how much the entire particle moved or changed over the imaging duration (Fig. 3b). This algorithm unlike nearest neighbor, cluster detection, or cross-correlation calculations has no assumptions about whether a particle moved into or out of a feature. Summary data for the

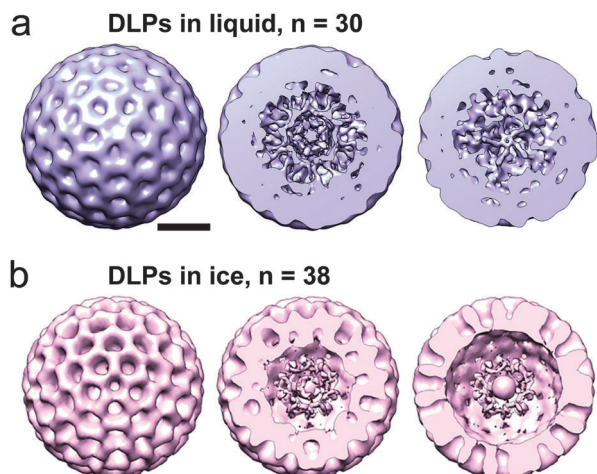
three representative particles demonstrated that Particle 1 (blue) exhibited statistically more pixel displacements than the other particles examined over a 10 second timeframe. Particle 2 (gray) showed the least overall mean pixel displacements, while Particle 3 (green) displayed an intermediate value of mean pixel displacements (Fig. 3b). A heat map summarizing the positions of the cumulative movements for each particle is given in Fig. 3a.

To build upon these measurements, we also quantified the lengths of the mRNA strands surrounding each particle, starting from the edge of the DLPs and extending to a 150 nm region in all directions. The total length of mRNA that surrounded Particle 1 (blue) was  $\sim 21$  nm. Particle 2 (gray) was surrounded by  $\sim 8$  nm of mRNA, while  $\sim 13$  nm of mRNA surrounded Particle 3 (green). We plotted the mean pixel displacements for each particle versus the relative lengths of associated mRNAs. These values represented a combination of more than 10 000 pixel movements in total. Upon examining a linear regression analysis of these measurements, we found a strong correlation ( $R^2 = 0.99$ ) existed (Fig. 3b). This information suggested that particles having more mobile units were associated with greater quantities of viral mRNAs. Although, the asynchronous nature of the DLPs does not permit us to fully determine the history of each particle, the use of these newly developed tools and analysis can be used to probe more deeply into viral transcription events in future experiments.

To validate the integrity of the DLPs used in our analysis, we calculated a 3D image reconstruction on the initial image in the series. We used the PARTICLE software package to select 30 individual DLPs from the image as previously described.<sup>6</sup> For reference, when imposing icosahedral (60-fold symmetry) during the reconstruction phase, 30 particles equates to 1800 asymmetric particles. The selected particles were then exported into the RELION software package.<sup>13</sup> Using RELION, we calculated a 3D structure of the DLPs while enforcing icosahedral symmetry operators over an angular search space of  $7.5^\circ$  throughout 25 cycles of refinement (Fig. 4a). Refinement parameters in RELION included a DLP reference model<sup>14</sup> low-pass filtered<sup>15</sup> to 50 Å, a pixel size of 5 Å, and a regularization parameter of  $T = 4$ . Consistent with previous 3D structures of enzymatically activated particles,<sup>6</sup> the resulting reconstruction of the DLPs in liquid showed dynamic features in their core interiors at  $\sim 2.8$  nm-resolution (Fig. 4a).

As an additional control, we analysed images of frozen-hydrated active DLPs. We implemented the same computing procedures and calculated a reconstruction using 38 particles from a single EM image while enforcing icosahedral symmetry. For reference, when enforcing icosahedral (60-fold symmetry) during the reconstruction phase, 38 particles equates to 2280 asymmetric particles. We chose to use slightly more particles for the cryo-EM calculations as we anticipated more views of the static DLPs would be needed to equate with the fluid system. The resulting 3D structure was strikingly similar to the DLP reconstruction in liquid and resolved to  $\sim 2.5$  nm. (Fig. 4b). A notable difference between reconstructions calculated for DLPs in ice and liquid is that the liquid structure had a continuum of density throughout the particles while the ice





**Fig. 4** 3D structures of DLPs reveal internal densities in liquid and ice specimens. (a) A 3D reconstruction was calculated for actively transcribing particles in a liquid environment. Cross-sections of the reconstruction are shown at  $\sim 20$  nm intervals to the midsection. The diameter of the reconstruction is  $\sim 80$  nm. (b) A 3D reconstruction was calculated for frozen-hydrated particles using the same image processing steps and while imposing icosahedral symmetry during the refinement procedure. The diameter of the resulting reconstruction is  $\sim 80$  nm. Scale bar is 30 nm.

structure had more rigidly defined segmented features visible in cross-sections through the density map (Fig. 4a and b). These differences may be due to a lack of fluidity in the captured particles resulting from the specimen freezing process. Overall, the 3D structural information in both the liquid and ice preparations complements the time-resolved measurements by ensuring the proper integrity of the activated viral assemblies.

As we continue to make progress toward real-time molecular imaging of biological systems, one item that can elevate our efforts is the use of direct electron detectors. These new detectors can operate in movie-mode at lower magnifications to further minimize issues of electron dose and frame rate. In addition, the use of thinner silicon nitride windows or graphene liquid cells may enable us to gather more information of how single-stranded mRNA transcripts emerge from viral particles. Similarly, reducing the thickness of the liquid cell in general could enable us to view protein–protein interactions or other viral processes. Improving each of these technologies will be important as we move toward examining smaller, active biological macromolecules. Smaller biological entities are weak

phase contrast objects and pose a great challenge to image in liquid with currently available resources. On course with data presented here, molecular imaging of biological assemblies at the nanoscale inspires a new era in microscopy, with the potential to further decode life processes and improve our knowledge of human health worldwide.

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