Probing the Separation Distance between Biological Nanoparticles and Cell Membrane Mimics Using Neutron Reflectometry with Sub-Nanometer Accuracy

Antonius Armanious,* Yuri Gerelli,* Samantha Micciulla, Hudson P. Pace, Rebecca J. L. Welbourn, Mattias Sjöberg, Björn Agnarsson, and Fredrik Höök*

ABSTRACT: Nanoparticle interactions with cellular membranes are controlled by molecular recognition reactions and regulate a multitude of biological processes, including virus infections, biological nanoparticle-mediated cellular communication, and drug delivery applications. Aided by the design of various supported cell membrane mimics, multiple methods have been employed to investigate these types of interactions, revealing information on nanoparticle coverage, interaction kinetics, as well as binding strength; however, precise quantification of the separation distance across which these delicate interactions occur remains elusive. Here, we demonstrate that carefully designed neutron reflectometry (NR) experiments followed by an attentive selection and application of suitable theoretical models offer a means to quantify the distance separating biological nanoparticles from a supported lipid bilayer (SLB) with sub-nanometer precision. The distance between the nanoparticles and SLBs was tuned by exploiting either direct adsorption or specific binding using DNA tethers with different conformations, revealing separation distances of around 1, 3, and 7 nm with nanometric accuracy. We also show that NR provides precise information on nanoparticle coverage, size distribution, material composition, and potential structural changes in the underlying planar SLB induced upon nanoparticle binding. The precision with which these parameters could be quantified should pave an attractive path for investigations of the interactions between nanoparticles and interfaces at length scales and resolutions that were previously inaccessible. This thus makes it possible to, for example, gain an in-depth understanding of the molecular recognition reactions of inorganic and biological nanoparticles with cellular membranes.

INTRODUCTION

Characterization of nanoparticle interactions that occur within nanometer distances to cellular membranes is crucial for an in-depth understanding of a multitude of biological processes, including neuronal signaling, viral infection, exosome-mediated intracellular communication, nanoparticle-assisted drug and vaccine delivery, as well as nanotoxicological effects. Methods to quantify the molecular length scales over which these types of interactions occur in cellular systems are few and essentially restricted to cryogenic transmission electron microscopy (cryo-TEM),1−4 fluorescence resonance energy transfer (FRET),5−6 and super-resolution optical microscopy approaches.7−11 However, the complex biomolecular content and topological architecture of cells complicate the analysis, particularly the determination of the absolute distances at which biomolecular interactions occur. This, in turn, motivated intensive efforts devoted to the design of reductionistic mimics of cell membranes.12−28

Besides offering the opportunity to deliberately define the molecular (lipid, protein, and glycan) composition of the membrane,20,29−33 surface-supported artificial cell membranes have also enabled a broader arsenal of advanced analytical tools to be employed. In particular, the application of tools such as surface plasmon resonance (SPR),24,35 quartz crystal microbalance (QCM),36−42 scanning probe microscopy (SPM),43,44 total internal reflection fluorescence (TIRF) microscopy,38,45−48 and surface-sensitive optical scattering techniques,39,50 advanced the understanding of how nanoparticles interact with cellular membranes, offering key insights on, for example, interaction kinetics,35,37,38,48,51 the dependence of lateral nanoparticle diffusivity on the number of molecular contact points,47,48,52 as well as membrane fusion of DNA-
tethered lipid vesicles, viruses, and SNARE-associated synaptic vesicles. These methods, alone or in combination, can also provide quantitative information on the mass and number of bound nanoparticles, and in the case of, for example, quartz crystal microbalance with dissipation monitoring (QCM-D) or multiparametric SPR, film thickness. However, with these methods, it is difficult to precisely quantify the distance across which the biomolecular interactions between the nanoparticles and an SLB occur, i.e., the actual distance separating the bound nanoparticles from the SLB.

One alternative method that is particularly well suited to quantify biomolecular film thickness and coverage is neutron reflectometry (NR). In particular, aided by the possibility to vary the scattering length density (SLD) contrast between the substrate, the adsorbed film, and the background buffer by exchanging protium (\(^{1}H\)) with deuterium (\(^{2}H\)), NR has been shown to offer absolute thickness determinations with sub-nanometer precision of biomolecular films and their internal structure, even for multilayered systems. In the case of SLBs, NR was not only successfully used to quantify minute differences in the thickness of hydrophobic and hydrophilic portions of SLBs made from different lipid compositions but also structural alterations induced by changes in environmental conditions and biomolecular interactions. However, while thickness determination is relatively straightforward, separation distance determination is particularly challenging using NR, as NR requires strong SLD contrasts in the direction orthogonal to the interface and a well-defined layered structure, so far only realized for homogeneous flat layers. Thus, due to the geometry of spherically shaped nanoparticles, NR has been rarely used for in-depth characterization of adsorbed nanoparticles. In most of the cases reported in the literature, the investigations focused on the effect of NPs on the structure of the underlying layers, e.g., SLBs, and on the quantification of the adsorbed amount without addressing the inherent possibility to characterize the structure, morphology, and separation distance between the substrate and the NP layer.

In this work, we demonstrate that by carefully designing the experiments and combining different theoretical models, NR can indeed be used to quantify the distance separating an SLB from naked and lipid-coated silica nanoparticles, as well as polydisperse hollow lipid vesicles. The distance between nanoparticles/vesicles and SLBs was varied by exploiting either direct nonspecific adsorption or specific binding. The latter was achieved via hybridization between cholesterol-modified DNA molecules, which were self-incorporated in the planar SLB and in the outer surface of lipid-coated NPs and hollow lipid vesicles. The DNA complexes were designed to adopt (i) a rod configuration to enforce a gap between the nanoparticles and the SLB or (ii) a zipper configuration, previously designed to mimic the SNARE–protein complex. We also show that NR provides an in-depth characterization that goes beyond the determination of separation distances. In fact, it provides precise information on the composition of adsorbed biological nanoparticles, their surface coverage and size distribution, as well as potential structural changes in the planar SLB upon interaction with nanoparticles or vesicles. The results refute a long-standing conviction in the field that NR cannot be used to study polydisperse biological nanoparticles. The unprecedented precision with which these parameters could be quantified should offer new means for an in-depth understanding of interactions between nanoparticles and interfaces occurring at nanometric separation distances, which is particularly relevant for investigations of the molecular recognition reactions of inorganic and biological nanoparticles with cellular membranes.

### RESULTS AND DISCUSSION

To cover different separation distances between planar SLBs and adsorbed nanoparticles as well as different nanoparticle compositions and size distributions, the following five systems formed on planar SLBs were investigated (Figure 1):

(i) direct adsorption of \(\text{SiO}_2\) NPs,

(ii) binding of lipid-coated \(\text{SiO}_2\) NPs (referred to as nanoSLBs) via extended DNA (referred to as ABCD-DNA),

(iii) binding of nanoSLBs via zipper DNA (referred to as XYWZ-DNA),

(iv) binding of hollow lipid vesicles via ABCD-DNA,

(v) binding of hollow lipid vesicles via XYWZ-DNA.

A thorough characterization of the \(\text{SiO}_2\) NPs, nanoSLBs, and lipid vesicles was conducted before running the NR experiments. The spicricity of the \(\text{SiO}_2\) NPs was independently confirmed using scanning electron microscopy and atomic force microscopy (AFM). Their diameter was also independently determined and found to be 143 and 146 ± 2 nm according to AFM and dynamic light scattering (DLS; Figure S1), as determined using DLS. Control experiments to ensure the proper formation of the nanoSLBs (Figure S7), optimize the experimental conditions (Figure S8), and rule out nonspecific binding between the nanoSLBs/vesicles and the planar SLB (Figures S9–S12), were conducted by means of the QCM-D technique.

The NR data were collected and analyzed systematically after each adsorption step to characterize (i) the bare silicon block, (ii) the planar SLBs formed on the surface of the silicon block, (iii) the DNA-decorated planar SLBs (for the relevant experiments), and (iv) the nanoparticle/vesicle layer. To achieve an unambiguous fitting of the data, we applied the contrast variation method, collecting NR data in aqueous buffers with various \(\text{D}_2\text{O}/\text{H}_2\text{O}\) mixing ratios, as explained hereafter. The measured neutron reflectivity depends on the so-called scattering length density (SLD), which is a volumetric average of the nuclear scattering length of the atoms that constitute a certain molecule. The difference between the SLD of different regions within the sample defines the contrast. SLD is also isotope-dependent and is, therefore, different for hydrogen-rich and deuterium-rich molecules. Indeed, SLDs for \(\text{D}_2\text{O}\) and \(\text{H}_2\text{O}\) are very different, namely, \(6.35 \times 10^{-6} \text{ Å}^{-2}\) and \(-0.56 \times 10^{-6} \text{ Å}^{-2}\), respectively (at 25 °C). Thus, by varying the SLD of the aqueous buffers, the contrast of the adsorbed layers with respect to the background buffer was varied, allowing to reveal different structural features of the sample and to enhance the robustness of the data modeling. In addition to \(\text{D}_2\text{O}\) and \(\text{H}_2\text{O}\), NR data were collected in \(\text{D}_2\text{O}/\text{H}_2\text{O}\) mixtures with nominal SLDs of \(3.47 \times 10^{-6} \text{ Å}^{-2}\) and \(2.07 \times 10^{-6} \text{ Å}^{-2}\). The former mixture was chosen to match the SLD of \(\text{SiO}_2\) (null contrast between the buffer and \(\text{SiO}_2\)), here referred to as \(\text{SiO}_2\)-matched water (\(\text{SiO}_2\text{-MW}\)), while the latter mixture was selected to match the SLD of Si (null contrast between the buffer and Si) and is therefore referred to here as Si-matched water (\(\text{Si-MW}\)). A detailed account of the experiments and all solutions used are provided in the Materials and Methods section, together with...
Figure 1. Schematics representing the five systems investigated in this work. (i) SiO$_2$ NPs: directly adsorbed SiO$_2$ NPs to a planar SLB; (ii) ABCD-nanoSLBs: lipid-coated SiO$_2$ NPs (referred to as nanoSLBs) attached to a planar SLB via extended DNA (referred to as ABCD-DNA); (iii) XYWZ-nanoSLBs: nanoSLBs attached to a planar SLB via zipper DNA (referred to as XYWZ-DNA); (iv) ABCD-vesicles: lipid vesicles attached to a planar SLB via ABCD-DNA; (v) XYWZ-vesicles: lipid vesicles attached to a planar SLB via XYWZ-DNA. Though the schematics are approximately proportional to real values, they are not drawn to scale. Detailed step-by-step schematics describing the experimental procedure for each system are given in Figure S2.
The values of all of the model parameters obtained from the analysis of the NR curves are summarized in Figure 2d, revealing that the size of the SiO$_2$ NPs is $2 \times r_{NP} = 142 \pm 2$ nm, which is in excellent agreement with the size estimation obtained by AFM ($143$ nm)$^{40}$ and DLS ($146 \pm 2$ nm in Figure S1) on the same NP batch. The slightly larger value obtained by DLS is expected as the technique probes the hydrodynamic size of particles, which is often larger than their actual size.

It is also worth mentioning that the model used for the data fitting allowed for direct determination of $r_{NP}$ and the water fraction in the SiO$_2$ NP layer, $f_{w,NP}$. This was achieved by directly including $f_{w,NP}$ and the VFP of a homogeneous sphere defined numerically as a function of $r_{NP}$ in the modeling (for details, see the Materials and Methods section). Adopting this approach avoided any ambiguity in defining the geometry of the NP layer, which would have otherwise arisen if a standard slab model were to be used. The fitted SLD value of the SiO$_2$ NPs, $\rho_{NP}$, is $3.47 \times 10^{-6}$ Å$^{-2}$.
\[ \rho_{\text{NP}} = 3.6 \pm 0.2 \times 10^{-6} \text{ Å}^{-2}, \] is in full agreement with the theoretical value of SiO\(_2\), \( \rho_{\text{SiO}_2} = 3.5 \times 10^{-6} \text{ Å}^{-2} \). Finally, based on \( f_{\text{NP}} \), we calculated the surface coverage of the SiO\(_2\) NPs to be \( \theta = 28.9\% \), which is in the expected range below the jamming limit of \( \sim 55\% \) for monodispersed spheres randomly adsorbing to a surface.\(^{27}\) A surface coverage lower than the jamming limit is likely due to adsorption of the NPs under stagnant, i.e., no flow, conditions. Under such conditions, one can expect local depletion of the NP concentration near the SLB interface, which may attenuate the final surface coverage.

NR also offers the unique opportunity to probe the NP layer while simultaneously monitoring changes in the underlying SLB. This is possible because the distance probed by NR is, to a good approximation, inversely proportional to \( Q \) and equals \( Q^2 / Q_{\text{min}}^2 \). Therefore, the NR signal of layers with thickness \( \gg 15 \text{ nm} \), such as the SiO\(_2\) NP layer, will primarily result in fringes in reflectivity.
in the low-Q range \((Q < 0.04 \, \text{Å}^{-1})\), and thinner layers, such as the SLB, will mainly contribute to the overall shape of the reflectivity curve, with features appearing, if any, in the mid and high-Q range \((Q > 0.05 \, \text{Å}^{-1})\). In fact, no detectable changes in the overall shape of the reflectivity curves were observed before and after the adsorption of the NPs (Figure S15), indicating that despite the formation of an NP layer in close proximity to the SLB, the NPs did not penetrate the SLB layer nor was there any indication of wrapping of lipids around the NPs.

To summarize, the NR data allowed us to construct a detailed structural description of the systems, including the separation distance, the size of the NPs, their material composition, their surface coverage, and the absence of detectable structural changes in the underlying planar SLB. By attentive selection of the modeling approach, all of these pieces of information were obtained using one experimental technique with unprecedented nanometric accuracy.

**ABCD-nanoSLBs.** Unlike the SiO\(_2\) NPs, which are single-component particles, nanoSLBs are two-component particles: a lipid shell and a silica core. To account for both, the VFP used to fit the NR data was built by superimposing the individual VFPs of a homogeneous sphere and a spherical shell. The resulting nanoSLB VFP was described by five fitting parameters: the radius and the SLD of the SiO\(_2\) core, \(r_{\text{core}}\) and \(\rho_{\text{core}}\), the thickness and the SLD of the lipid shell, \(t_{\text{shell}}\) and \(\rho_{\text{shell}}\), and the solution volume fraction, \(f_{\text{core,nanoSLB}}\). Using this approach, excellent fits to the experimental data were achieved (Figure 3a). One might at first thought assume that a thin lipid layer, <5 nm, coating a much larger NP, ~150 nm, would have an insignificant contribution to the NR signal; however, the reflectivity curves (Figure 3a) and the corresponding SLD profiles (Figure 3b) show otherwise. Unlike the SiO\(_2\) NPs, the reflectivity of the nanoSLB layer in SiO\(_2\)-MW exhibited fringes in the low-Q range (Figure 3a, inset), demonstrating that the presence of the lipid shell renders the nanoSLBs visible even if the SiO\(_2\) core is contrast-matched (Figure 3b). If one were to ignore the multicomponent structure of the nanoSLBs and try to model them as a one-component entity, no acceptable fitting of the data would be achievable (Figure S16). This ability to distinguish the different components of biological nanoparticles, e.g., the core and the shell, may be paramount to understanding biological processes occurring at the cell membrane interface, for example, the release of a viral genome from the capsid during membrane fusion.

The VFPs of the nanoSLB layer and the underlying layers are shown in Figure 3c, revealing \(t_{\text{gap}} = 7 \pm 1\) nm. The value is, as expected, larger than \(t_{\text{gap}}\) of the SiO\(_2\) NPs (1 ± 1 nm; Figure 2c, inset) and smaller than the theoretical length of completely extended 45 DNA base pairs of ~15 nm. The separation distance also suggests that the DNA linkers are either at a nonperpendicular angle to the surface or have a secondary winded structure. As indicated by the small error of 1 nm, the quality of the fit is very sensitive to any variations in \(t_{\text{gap}}\); trying to fit the data while forcing \(t_{\text{gap}}\) outside the optimal range results in fits with substantially lower quality (Figure S17). Figure 3d summarizes the results of the fitted parameters, showing the anticipated size and composition of both the core and the shell: \(2 \times r_{\text{core}} = 140 \pm 4\) nm (within the error ranges indistinguishable from the SiO\(_2\) NP size determined using AFM, DLS, and NR in Figure 2d), \(\rho_{\text{core}} = 3.3 \pm 0.2 \times 10^{-6} \, \text{Å}^{-2}\) (in perfect agreement with \(\rho_{\text{SiO}_2} = 3.5 \times 10^{-6} \, \text{Å}^{-2}\)), \(t_{\text{shell}} = 4.1 \pm 0.5\) nm (overlapping the range of values reported for planar SLBs, i.e., between 4.3 ± 0.1 nm and 4.7 ± 0.1 nm, Table S1), and \(\rho_{\text{shell}} = 0.33 \pm 0.05 \times 10^{-6} \, \text{Å}^{-2}\) (identical to the SLD of the POPC lipids composing the shell, \(\rho_{\text{POPC}} = 0.33 \times 10^{-6} \, \text{Å}^{-2}\)). The nanoSLBs had a surface coverage of \(\theta = 41.6\%\), also within the expected range. Similar to the SiO\(_2\) NPs, the results show no sign of change in the structure of the underlying planar SLB (Figure S18); similarly, the results for XYWZ-nanoSLBs (Figure S19), ABCD-vesicles (Figure S20), and XYWZ-vesicles (Figure S21) exhibit no signs of structural changes of the underlying planar SLB.

**XYWZ-nanoSLBs.** Figure 4a shows the reflectivity curves for the XYWZ-nanoSLBs. Similar to the ABCD-nanoSLBs, the curves exhibit high-frequency fringes in the low-Q range (Figure 4a, inset), indicative of the formation of a thick layer, much thicker than the planar SLB. However, the amplitude of the fringes in all aqueous solutions is damped compared to the ABCD-nanoSLBs, indicating substantially lower surface coverage. Under the same experimental conditions, it is expected that XYWZ-nanoSLBs will have lower surface coverage than ABCD-nanoSLBs due to differences in the configuration between the two DNA linkers. While ABCD-DNA is anchored with two connected cholesterol molecules per DNA linker at both the planar SLB and the nanoSLB ends, the XYWZ-DNA is anchored with only one cholesterol molecule per DNA linker at both ends, due to the separation into two linkers upon hybridization of the XY- with WZ-DNA, forming an XW linker and a YZ linker (Figure 1). It has been previously shown that DNA molecules attached with single cholesterol anchors bind reversibly to lipid bilayers, while those attached with double anchors bind essentially irreversibly. This difference is likely due to the continuous switching of cholesterol molecules between the aqueous and lipid phases. Such configuration allows only XYWZ-nanoSLBs with a sufficiently large number of linkers to remain attached to the planar SLB as illustrated in Figure S22.

At such low surface coverage, it is more challenging to resolve details such as the thickness of the lipid shell, as the layer contrast is directly proportional to the adsorbed amount. For data fitting, we have thus used the same approach as for the ABCD-nanoSLBs, but with fixing the shell thickness to the same value obtained from fitting the ABCD-nanoSLBs, \(t_{\text{shell}} = 4.1\) nm. The resulting SLD profiles are very similar to that of the ABCD-nanoSLBs, but with attenuated curvature at the \(z\) range where the nanoSLBs are located (Figure 4b), confirming the lower surface coverage of the XYWZ-nanoSLBs. Still, the corresponding VFPs reveal the key difference between the two nanoSLB systems, namely, \(t_{\text{gap}} = 3.5 \pm 0.5\) nm (Figure 4c), which is half of that of the ABCD-nanoSLBs. Indeed, the determined separation distance lies precisely in the expected range: larger than that of the SiO\(_2\) NPs (1 ± 1 nm) and smaller than ABCD-nanoSLBs (7 ± 1 nm). All other fitting parameters, i.e., \(r_{\text{core}}\), \(\rho_{\text{core}}\), and \(\rho_{\text{shell}}\) (Figure 4d), were in perfect agreement with both the expected values and the ones determined for the ABCD-nanoSLBs. As indicated from the reflectivity curves, SLD profiles, and VFPs, the surface coverage of XYWZ-nanoSLBs was substantially low at \(\theta = 15.9\%\).

**ABCD-Vesicles.** The results from the SiO\(_2\) NPs, ABCD-nanoSLBs, and XYWZ-nanoSLBs demonstrated the successful use of NR to resolve the separation distances between planar SLBs and silica-based nanoparticles with sub-nanometer accuracy while revealing insightful details about their structure and composition. Many biological NPs are, however, more polydisperse and less dense than the so far investigated systems. Encouraged by the results of the SiO\(_2\) NPs and nanoSLBs experiments presented above, we decided to investigate the use of...
of NR to resolve the separation distances of polydisperse, hollow lipids vesicles. In fact, with their hollow structure and relatively broad size distribution (Figure S1), the lipid vesicles used here are likely to be more challenging to obtain a sufficient signal than from most native biological NPs samples. This is so because most biological NPs, such as purified virus samples have a relatively tight size distribution, and their cores are at least partially filled with the viral genome, which makes them likely to be more visible to NR and easier to analyze than polydisperse and hollow lipid vesicles. This system thus represents a challenging experimental scenario, serving as a benchmark to define the limits of the approach.

Figure 4. Characterization of a nanoSLB layer attached to a planar SLB using XYWZ-DNA. (a) Neutron reflectometry data and fits in aqueous buffers with four different H_2O and D_2O mixing ratios: (i) D_2O: nominal SLD 6.35 x 10^{-6} Å^{-2}; (ii) SiO_2 MW: nominal SLD 3.47 x 10^{-6} Å^{-2}; (iii) SiMW: nominal SLD 2.07 x 10^{-6} Å^{-2}; (iv) H_2O: nominal SLD =0.56 x 10^{-6} Å^{-2}. The likelihood parameter \( \chi^2 \) is also indicated. The inset shows a zoomed-in section of the plot at the low-Q range. (b) SLD profiles versus the perpendicular distance from the surface of the silicon block, \( z \), in the four different aqueous solutions. (c) Volume fraction profiles (VFPs) of all components of the system: the silicon block (Si), the SiO_2 layer on top, the SLB, the XYWZ-DNA layer, and the nanoSLB layer. The separation distance, \( t_{gap} \), is also noted, which represents the distance between the inflection point of the VFP of the outer headgroups of the planar SLB and the first nonzero value for the VFP of the nanoSLB layer. The inset shows a zoomed-in section, depicting the VFPs of the SiO_2 cores, the lipid shells, and the sum of both, which represents the VFP of the nanoSLBs. *The XYWZ-DNA molecules had a negligible volume fraction. (d) Artistic schematics showing lateral and in-plane views of the system, with all of the fitted parameters noted on the schematic: thickness of the planar SLB, \( t_{SLB} \), radius and SLD of the core, \( r_{core} \) and \( \rho_{core} \), thickness and SLD of the shell, \( t_{shell} \) and \( \rho_{shell} \), separation distance, \( t_{gap} \), and surface coverage, \( \theta \). Though the schematics are approximately proportional to real values, they are not drawn to scale. All model parameters are listed in Table S4.
SiO$_2$ NP and the nanoSLB layers, indicative of the formation of a thinner layer. This result is expected given the smaller size of the vesicles (Figure S1). These fringes were almost invisible in the reflectivity curve collected in H$_2$O, which was expected due to the reduced contrast (difference in SLD) between H$_2$O and the POPC lipids composing the vesicles. In theory, one can use the same VFP-based fitting approach as was used for the SiO$_2$ NPs and nanoSLBs to fit the NR data collected upon the formation of a vesicle layer. However, this would require accounting for the size distribution when calculating $r_{core}$. Unfortunately, such an approach comes at a high computational cost, hindering successful data fitting within reasonable time frames. Therefore, the NR data were analyzed using a modified version of the standard slab model in which the vesicle layer was described by a slab with asymmetric roughness on both sides. The resulting SLD profiles (Figure Sb) reveal a small roughness at the interface close to the planar SLB and a much larger roughness at the interface between the vesicles and the bulk water solution.

![Figure 5](https://i.imgur.com/ABC5.png)

**Figure 5.** Characterization of a vesicle layer attached to a planar SLB using ABCD-DNA. (a) Neutron reflectometry data and fits in aqueous buffers with four different H$_2$O and D$_2$O mixing ratios: (i) D$_2$O: nominal SLD $6.35 \times 10^{-6}$ Å$^{-2}$; (ii) SiO$_2$MW: nominal SLD $3.47 \times 10^{-6}$ Å$^{-2}$; (iii) SiMW: nominal SLD $2.07 \times 10^{-6}$ Å$^{-2}$; (iv) H$_2$O: nominal SLD $-0.56 \times 10^{-6}$ Å$^{-2}$. The likelihood parameter $\chi^2$ is also indicated. The inset shows a zoomed-in section of the plot at the low-Q range. (b) SLD profiles versus the perpendicular distance from the surface of the silicon block, $z$, in the four different aqueous solutions. (c) Volume fraction profiles (VFPs) of all components of the system: the silicon block (Si), the SiO$_2$ layer on top, the SLB, the ABCD-DNA layer, and the vesicle layer. The separation distance, $t_{gap}$, is also noted, which represents the distance between the inflection points of the VFP of the outer headgroups of the planar SLB and that of vesicle layer minus $t_{shell}/2$ (eq S15). The inset shows a zoomed-in section of the empirically determined VFP of the vesicles and an overlayed VFP based on the fitted Schutz distribution. (d) Size distribution of the vesicles and an artistic schematic showing lateral and in-plane views of the system, with all of the fitted parameters noted on the schematic: thickness of the planar SLB, $t_{SLB}$, SLD of the shell, $\rho_{shell}$, separation distance, $t_{gap}$, and surface coverage, $\theta$. Though the schematics are approximately proportional to real values, they are not drawn to scale. All model parameters are listed in Table S5.
indicative of the arrangement and broad size distribution of the attached vesicles. The resulting VFPs (Figure 5c) reveal a separation distance, \( t_{\text{gap}} = 6.5 \pm 0.5 \) nm, which within the error ranges is indistinguishable from the separation distance of the ABCD-nanoSLB layers.

We then used the empirically determined VFP of the vesicle layer (extracted from the SLD profiles) to determine the size distribution of the adsorbed vesicles (Figure 5c, inset). This calculation was performed assuming spherical vesicles with a shell thickness of 4.1 nm (equivalent to the shell thickness of the nanoSLBs) and size distribution described by the Schultz function.\(^8^1\) Detailed calculations are provided in the Materials and Methods section. The size distribution of the adsorbed vesicles shifted toward smaller sizes (Figure 5d) with respect to that obtained by DLS for the same batch of vesicles in solution (Figure S1). This observation is in line with the well-documented bias for adsorption of smaller entities in a pool of polydisperse particles, a phenomenon generally attributed to the

**Figure 6.** Characterization of a vesicle layer attached to a planar SLB using XYWZ-DNA. (a) Neutron reflectometry data and fits in aqueous buffers with four different H\(_2\)O and D\(_2\)O mixing ratios: (i) D\(_2\)O: nominal SLD \(6.35 \times 10^{-6}\) Å\(^{-2}\); (ii) SiO\(_2\)MW: nominal SLD \(3.47 \times 10^{-6}\) Å\(^{-2}\); (iii) SiMW: nominal SLD \(2.07 \times 10^{-6}\) Å\(^{-2}\); (iv) H\(_2\)O: nominal SLD \(-0.56 \times 10^{-6}\) Å\(^{-2}\). The likelihood parameter \(\chi^2\) is also indicated. The inset shows a zoomed-in section of the plot at the low-Q range. (b) SLD profiles versus the perpendicular distance from the surface of the silicon block, \(z\), in the four different aqueous solutions. (c) Volume fraction profiles (VFPs) of all components of the system: the silicon block (Si), the SiO\(_2\) layer on top, the SLB, the XYWZ-DNA layer, and the vesicle layer. The separation distance, \(t_{\text{gap}}\), is also noted, which represents the distance between the inflection points of the VFP of the outer headgroups of the planar SLB and that of vesicle layer minus \(t_{\text{shell}}/2\) (eq S15). The inset shows a zoomed-in section of the empirically determined VFP of the vesicles and an overlayed VFP based on the fitted Schutz distribution. *The XYWZ-DNA molecules had a negligible volume fraction. (d) Size distribution of the vesicles and an artistic schematic showing lateral and in-plane views of the system, with all of the fitted parameters noted on the schematic: thickness of the planar SLB, \(t_{\text{SLB}}\), SLD of the shell, \(\rho_{\text{shell}}\), separation distance, \(t_{\text{gap}}\), and surface coverage, \(\theta\). Though the schematics are approximately proportional to real values, they are not drawn to scale. All model parameters are listed in Table S6.
toward the cellular membrane to initiate fusion with neuronal change of the SNARE complex that brings synaptic vesicles to, for example, investigate the calcium-facilitated structural tempting to speculate about the potential possibility of using NR volume fraction coverages as low as 5%. With this precision, it is sub-nanometric accuracy, even for polydisperse samples at distance between the nanoparticles and the membranes with zonation. Most notable is the ability to measure the separation distance between the vesicle and the planar mimetic ■ interactions of biological nanoparticles with planar mimetic nanoSLBs; the lower surface coverage is expected due to the reasons discussed for the XYWZ-vesicles and illustrated in Figure S22, allowing only vesicles with a sufficient number of DNA linkers to remain attached. The VFPs (Figure 6c) reveal that the volume fraction of the vesicles peaked at ~5%, a remarkably low value. The corresponding separation distance between the vesicle and the planar SLB was determined to be tpdp = 2.3 ± 0.9 nm, which within the error ranges is indistinguishable from the distance determined for XYWZ-nanoSLBs, tsdp = 3.5 ± 0.5 nm. The size distribution determined by fitting the VFP of the vesicle layer to the Schultz distribution function (Figure 6c, inset, and d) reveals a shift to larger vesicle sizes compared to that of the ABCD-vesicles. This shift is attributed to the higher number of DNA linkers needed for the XYWZ-vesicles to remain attached to the surface, which is preferentially accommodated by larger vesicles (Figure S22). Finally, the computed surface coverage (Figure S6) was found to be θ = 20.9%. While being even higher than the surface coverage of the XYWZ-nanosSLBs (θ = 15.9%), the volume fraction, which is the main parameter contributing to the NR signal, occupied by the vesicles is less than one-third that of the nanoSLBs, due to the hollow structure of the vesicles.

**CONCLUSIONS**

In this work, we have shown that NR can be used to interrogate the interactions of biological nanoparticles with planar mimetic cell membranes, offering an integrated structural characterization. Most notable is the ability to measure the separation distance between the nanoparticles and the membranes with sub-nanometric accuracy, even for polydisperse samples at volume fraction coverages as low as 5%. With this precision, it is tempting to speculate about the potential possibility of using NR to, for example, investigate the calcium-facilitated structural change of the SNARE complex that brings synaptic vesicles toward the cellular membrane to initiate fusion with neuronal cells. Similarly, viruses have evolved equally fine-tuned spatiotemporal mechanisms to control cellular uptake and subsequent association and fusion with endosomal membranes to release their genetic cargo into the cytosol, a process that could also be addressed using NR.

The richness of the NR data also allowed us to determine the size and material composition of each component of the nanoparticles, the size distribution of the polydisperse samples, surface coverage, and even probe the possible structural changes in the underlying mimetic cell membrane. The possibility to characterize interactions, material composition, and structural changes within nanometer distances to cellular membranes is crucial for optimizing supramolecular assembly strategies for many biomedical applications, such as the intermolecular interactions at cellular membranes that inspired the design of the pH-sensitive lipid nanoparticles (LNPs) used for drug delivery, which was recently applied for mRNA vaccines that helped mitigate the COVID-19 pandemic. Additionally, applications for biosensor design and optimization, as well as interactions of nanoparticles at environmental interfaces, are also of direct relevance. Thus, given the unprecedented precision with which we quantified these crucial parameters, NR could offer an in-depth understanding of the interaction between nanoparticles and interfaces in general, and biological nanoparticles and cellular membranes in particular.

**ASSOCIATED CONTENT**

The Supporting Information is available free of charge at https://pubs.acs.org/doi/10.1021/jacs.2c08456.

Detailed materials and methods with explanatory figures; tables summarizing the values of all of the modeling parameters; figures for results from DLS experiments to characterize the SiO$_2$ NPs, nanoSLBs, and vesicles; preliminary experiments conducted using QCM-D; NR results for the silicon blocks and planar SLBs; before and after NR data; additional fitting results showing the sensitivity of the results to minor variations in the fitted parameters; and schematics showing the difference in binding configuration between the ABCD- and XYWZ-DNA (PDF)

**AUTHOR INFORMATION**

Corresponding Authors
Antonius Armanious — Department of Physics, Chalmers University of Technology, 41296 Gothenburg, Sweden; Present Address: ETH Zurich, 8092 Zurich, Switzerland; orcid.org/0000-0002-8809-4659; Email: armanioa@ethz.ch

Yuri Gerelli — Institut Max von Laue-Paul Langevin (ILL), 38042 Grenoble, France; Department of Life and Environmental Sciences, Università Politecnica delle Marche, 60131 Ancona, Italy; orcid.org/0000-0001-5655-8298; Email: y.gerelli@staff.univpm.it

Fredrik Höök — Department of Physics, Chalmers University of Technology, 41296 Gothenburg, Sweden; orcid.org/0000-0003-1994-5015; Email: fredrik.hook@chalmers.se

Authors
Samantha Micciulla — Institut Max von Laue-Paul Langevin (ILL), 38042 Grenoble, France

Hudson P. Pace — Department of Physics, Chalmers University of Technology, 41296 Gothenburg Sweden; orcid.org/0000-0001-5116-2577

Rebecca J. L. Welbourn — ISIS Facility, STFC, Rutherford Appleton Laboratory, Chilton, Didcot, Oxon OX11 0QX, United Kingdom; orcid.org/0000-0002-4254-5354

Mattias Sjöberg — Department of Physics, Chalmers University of Technology, 41296 Gothenburg Sweden; orcid.org/0000-0003-3753-2564

https://doi.org/10.1021/jacs.2c08456

J. Am. Chem. Soc. 2022, 144, 20726–20738
Complete contact information is available at: https://pubs.acs.org/10.1021/jacs.2c08456

Notes
The authors declare no competing financial interest.

ACKNOWLEDGMENTS

The authors acknowledge the support of the Knut and Alice Wallenberg Foundation (#2019-0577), the Swedish Research Council (#2018-04900), and Röntgen Ångstrom Cluster (2015-06139). They thank Emanuel Schneck (TU Darmstadt) for support with preliminary X-ray reflectivity experiments and valuable discussions. Experiments at the ISIS Neutron & Muon Source were supported by a beamtime allocation RB1710273 from the Science and Technology Facilities Council. NR data is available under: https://doi.org/10.5286/ISIS.E.RB1710273.

REFERENCES

(1) Carlson, L. A.; Briggs, J. A. G.; Glass, B.; Riches, J. D.; Simon, M. N.; Johnson, M. C.; Müller, B.; Grünewald, K.; Kräusslich, H. G. Three-Dimensional Analysis of Budding Sites and Released Virus Suggests a Revised Model for HIV-1 Morphogenesis. Cell Host Microbe Microbe 2008, 4, 592–599.

(2) Maurer, U. E.; Sodeik, B.; Grünewald, K. Native 3D Intermediate of Membrane Fusion in Herpes Simplex Virus 1 Entry. Proc. Natl. Acad. Sci. U.S.A. 2008, 105, 10559–10564.

(3) le Bihan, O.; Bonnafous, P.; Marak, L.; Bickel, T.; Trépout, S.; Mornet, S.; de Haas, F.; Talbot, H.; Taveau, J. C.; Lambert, O. Cryo-Electron Tomography of Nanoparticle Transmigration into Liposome. J. Struct. Biol. 2009, 168, 419–425.

(4) Xia, Z.; Zhou, Q.; Lin, J.; Liu, Y. Stable SNARE Complex Prior to Evoked Synaptic Vesicle Fusion Revealed by Fluorescence Resonance Energy Transfer. J. Biol. Chem. 2001, 276, 1766–1771.

(5) Hogue, I. B.; Hoppe, A.; Ono, A. Quantitative Fluorescence Resonance Energy Transfer Microscopy Analysis of the Human Immunodeficiency Virus Type 1 Gag-Gag Interaction: Relative Contributions of the CA and NC Domains and Membrane Binding. J. Virol. 2009, 83, 7322–7336.

(6) Chen, H.; Kim, S.; Wang, S.; Park, K.; Cheng, J. X. Release of Hydrophobic Molecules from Polymer Micelles into Cell Membranes Revealed by Förster Resonance Energy Transfer Imaging. Proc. Natl. Acad. Sci. U.S.A. 2008, 105, 6596–6601.

(7) Betzig, E.; Patterson, G. H.; Sougrat, R.; Lindwasser, O. W.; Olenych, S.; Bonifacino, J. S.; Davidson, M. W.; Lippincott-Schwartz, J.; Hess, H. F. Imaging Intracellular Fluorescent Proteins at Nanometer Resolution. Science 2006, 313, 1642–1645.

(8) Li, D.; Shao, L.; Chen, B. C.; Zhang, X.; Zhang, M.; Moses, B.; Milkie, D. E.; Beach, J. R.; Hamer, J. A.; Pasham, M.; Kirchhausen, T.; Baird, M. A.; Davidson, M. W.; Xu, P.; Betzig, E. Extended-Resolution Structured Illumination Imaging of Endocytic and Cytoskeletal Dynamics. Science 2015, 349, No. aab3500.

(9) Lelek, M.; di Nuzzo, F.; Henriques, R.; Charneau, P.; Arhel, N.; Zimmer, C. Superresolution Imaging of HIV in Infected Cells with PALM. Proc. Natl. Acad. Sci. U.S.A. 2012, 109, 8564–8569.

(10) Taylor, R. W.; Mahmoodadjadi, R. G.; Rauschenberger, V.; Giessl, A.; Schambony, A.; Sandoghdar, V. Interferometric Scattering Microscopy Reveals Microsecond Nanoscopic Protein Motion on a Live Cell Membrane. Nat. Photonics 2019, 13, 480–487.

(11) Richter, R. P.; Hock, K. K.; Burkhartsmeyer, J.; Boehm, H.; Bingen, P.; Wang, G.; Steinmetz, N. F.; Evans, D. J.; Spatz, J. P. Membrane-Graded Hyaluronan Films: A Well-Defined Model System of Glycoconjugate Cell Coats. J. Am. Chem. Soc. 2007, 129, 5306–5307.

(12) Pace, H. P.; Hannestad, J. K.; Armanious, A.; Adams, M.; Agranarsson, B.; Gunnarsson, A.; Macciulla, S.; Sjövall, P.; Gerelli, Y.; Höök, F. Structure and Composition of Native Membrane Derived Polymer-Supported Lipid Bilayers. Anal. Chem. 2018, 90, 13065–13072.

(13) Holt, S. A.; le Brun, A. P.; Majkrzak, C. F.; McGillivray, D. J.; Heinrich, F.; Lösch, M.; Lakey, J. H. An Ion-Channel-Containing Model Membrane: Structural Determination by Magnetic Contrast Neutron Reflectometry. Soft Matter 2009, 5, 2576–2586.

(14) Liu, H. Y.; Pappa, A. M.; Pavia, A.; Pitsalidis, C.; Thiburse, Q.; Salleo, A.; Owens, R. M.; Daniel, S. Self-Assembly of Mammalian-Cell Membranes on Bioelectronic Devices with Functional Transmembrane Proteins. Langmuir 2020, 36, 7523–7531.

(15) Daniel, S.; Diaz, A. J.; Martinez, K. M.; Bench, B. J.; Albertorio, F.; Cremer, P. S. Separation of Membrane-Bound Compounds by Solid-Supported Bilayer Electrophoresis. J. Am. Chem. Soc. 2007, 129, 8072–8073.

(16) Shi, J.; Chen, J.; Cremer, P. S. Sub-100 Nm Patternning of Supported Bilayers by Nanoshaving Lithography. J. Am. Chem. Soc. 2008, 130, 2718–2719.

(17) Liu, H. Y.; Grant, H.; Hsu, H. L.; Sorkin, R.; Bolković, F.; Wuite, G.; Daniel, S. Supported Planar Mammalian Membranes as Models of in Vivo Cell Surface Architectures. ACS Appl. Mater. Interfaces 2017, 9, 35526–35538.

(18) Cremer, P. S.; Groves, J. T.; Kung, L. A.; Boxer, S. G. Writing and erasing barriers to lateral mobility into fluid phospholipid bilayers. Langmuir 1999, 15, 3893–3896.

(19) Groves, J. T.; Ulman, N.; Cremer, P. S.; Boxer, S. G. Substrate-Membrane Interactions: Mechanisms for Imposing Patterns on a Fluid Bilayer Membrane. Langmuir 1998, 14, 3347–3350.

(20) Kam, L.; Boxer, S. G. Formation of Supported Lipid Bilayer Composition Arrays by Controlled Mixing and Surface Capture. J. Am. Chem. Soc. 2000, 122, 12901–12902.

(21) Ferhan, A. R.; Yoon, B. K.; Park, S.; Sut, T. N.; Chin, H.; Park, J. H.; Jackman, J. A.; Cho, N. J. Solvent-Assisted Preparation of Supported Lipid Bilayers. Nat. Protoc. 2019, 14, 2091–2118.

(22) Chun, M.; Lowe, R. D.; Chan, Y. H. M.; Ganesan, P.; Boxer, S. G. DNA-Tethered Membranes Formed by Giant Vesicle Rupture. J. Struct. Biol. 2009, 168, 190–199.

(23) Shen, W. F.; Boxer, S. G.; Knoll, W.; Frank, C. W. Polymer-Supported Lipid Bilayers on Benzophenone-Modified Substrates. Biomacromolecules 2001, 2, 70–79.

(24) Chan, Y. H. M.; Boxer, S. G. Model Membrane Systems and Their Applications. Curr. Opin. Chem. Biol. 2007, 11, 581–587.

(25) Groves, J. T.; Ulman, N.; Boxer, S. G. Micropatterning Fluid Lipid Bilayers on Solid Supports. Science 1997, 275, 651–653.

(26) Cremer, P. S.; Boxer, S. G. Formation and Spreading of Lipid Bilayers on Planar Glass Supports. J. Phys. Chem. B 1999, 103, 2554–2559.

(27) Richards, M. J.; Hsia, C. Y.; Singh, R. R.; Haider, H.; Kumpf, J.; Kawate, T.; Daniel, S. Membrane Protein Mobility and Orientation Preserved in Supported Bilayers Created Directly from Cell Plasma Membrane Blebs. Langmuir 2016, 32, 2963–2974.

(28) Hughes, A. V.; Holt, S. A.; Daulton, E.; Soliakov, A.; Charlton, T. R.; Roser, S. J.; Lakey, J. H. High Coverage Fluid-Phase Floating Lipid Bilayers Supported by α-Thiolid Self-Assembled Monolayers. J. R. Soc. Interface 2014, 11, No. 20140447.

(29) Peerboom, N.; Schmidt, E.; Trybala, E.; Block, S.; Bergröten, T.; Pace, H. P.; Bally, M. Cell Membrane Derived Platform to Study Virus Binding Kinetics and Diffusion with Single Particle Sensitivity. ACS Infect. Dis. 2018, 4, 944–953.

(30) Pace, H.; Simonsson Nyström, L.; Gunnarsson, A.; Eck, E.; Monson, C.; Geschwindner, S.; Snijder, A.; Höök, F. Preserved Transmembrane Protein Mobility in Polymer-Supported Lipid Bilayers Derived from Cell Membranes. Anal. Chem. 2015, 87, 9194–9203.

(31) Melby, E. S.; Mensch, A. C.; Lohse, S. E.; Hu, D.; Orr, G.; Murphy, C. J.; Hamers, R. J.; Pedersen, J. A. Formation of Supported Lipid Bilayers Containing Phase-Segregated Domains and Their
Interaction with Gold Nanoparticles. Environ. Sci.: Nano 2016, 3, 45–55.

(32) Kam, L.; Boxer, S. G. Spatially Selective Manipulation of Supported Lipid Bilayers by Laminar Flow: Steps toward Biomembrane Microfluidics. Langmuir 2003, 19, 1624–1631.

(33) Hovis, J. S.; Boxer, S. G. Patterning and Composition Arrays of Supported Lipid Bilayers by Microcontact Printing. Langmuir 2001, 17, 3400–3405.

(34) Hidari, K. I. P. J.; Shimada, S.; Suzuki, Y.; Suzuki, T. Binding Kinetics of Influenza Viruses to Sialic Acid-Containing Carbohydrates. Glycoconjugate J. 2007, 24, 583–590.

(35) Altgäude, N.; Eriksson, C.; Peerboom, N.; Phan-Xuan, T.; Moeller, S.; Schnabelrauch, M.; Svedhem, S.; Trybala, E.; Bergström, T.; Bally, M. Mucin-like Region of Herpes Simplex Virus Type 1 Attachment Protein Glycoprotein C (GC) Modulates the Virus-Glycosaminoglycan Interaction. J. Biol. Chem. 2015, 290, 21473–21485.

(36) Rydell, G. E.; Dahlin, A. B.; Höök, F.; Larson, G. QCM-D Studies of Human Norovirus VLPs Binding to Glycosphingolipids in Supported Lipid Bilayers Reveal Variant-Specific Characteristics. Glycobiology 2009, 19, 1176–1184.

(37) Szklarczyk, O. M.; González-Segredo, N.; Kukura, P.; Oppenheim, A.; Choquet, D.; Sandoghdar, V.; Helenius, A.; Sbalzarini, I. F.; Ewers, H. Receptor Concentration and Diffusivity Control Multivalent Binding of SV40 to Membrane Bilayers. PLoS Comput. Biol. 2013, 9, e1003310.

(38) Bally, M.; Block, S.; Höök, F.; Larson, G.; Parveen, N.; Rydell, G. E. Physicochemical Tools for Studying Virus Interactions with Targeted Cell Membranes in a Molecular and Spatiotemporally Resolved Context. Anal. Bioanal. Chem. 2021, 413, 7157–7178.

(39) Parveen, N.; Rydell, G. E.; Larson, G.; Hytönen, V. P.; Zhadanov, V. P.; Höök, F.; Block, S. Competition for Membrane Receptors: Norovirus Detachment via Lectin Attachment. J. Am. Chem. Soc. 2019, 141, 16303–16311.

(40) Armanios, A.; Agrnarsson, B.; Lundgren, A.; Zhadanov, V. P.; Höök, F. Determination of Nano-Sized Adsorbate Mass in Solution Using Mechanical Resonators: Elimination of the so Far Inseparable Liquid Contribution. J. Phys. Chem. C 2021, 125, 22733–22746.

(41) Lochbaum, C. A.; Chew, A. K.; Zhang, X.; Rotello, V.; van Lehnn, R. C.; Pedersen, J. A. Lipophility of Cationic Lipids Promotes Irreversible Adsorption of Nanoparticles to Lipid Bilayers. ACS Nano 2021, 15, 6562–6572.

(42) Melby, E. S.; Solse, S. E.; Park, J. E.; Vartanian, A. M.; Putans, R. A.; Abbott, H. B.; Hamers, R. J.; Murphy, C. J.; Pedersen, J. A. Cascading Effects of Nanoparticle Coatings: Surface Functionalization Dictates the Assemble of Complexed Proteins and Subsequent Interaction with Model Cell Membranes. ACS Nano 2017, 11, 5489–5499.

(43) Sieben, C.; Kappel, C.; Zhu, R.; Wozniak, A.; Rankl, C.; Hinterdorfer, P.; Grumbümler, H.; Herrmann, A. Influenza Virus Binds Its Host Cell Using Multiple Dynamic Interactions. Proc. Natl. Acad. Sci. U.S.A. 2012, 109, 13626–13631.

(44) Merkel, R.; Nassoy, P.; Leung, A.; Ritchie, K.; Evans, E. Energy Landscapes of Receptor-Ligand Bonds Explored with Dynamic Force Spectroscopy. Nature 1999, 397, 50–53.

(45) Gunnarsson, A.; Jönsson, P.; Marie, R.; Tegenfeldt, J. O.; Höök, F. Single-Molecule Detection and Mismatch Discrimination of Unlabeled DNA Targets. Nano Lett. 2008, 8, 183–188.

(46) Lee, D. W.; Hsu, H. L.; Bacon, K. B.; Daniel, S. Image Restoration and Analysis of Influenza Virions Binding to Membrane Receptors Reveal Adhesion-Strengthening Kinetics. PLoS One 2016, 11, No. e0163437.

(47) Block, S.; Zhadanov, V. P.; Höök, F. Quantification of Multivalent Interactions by Tracking Single Biological Nanoparticle Mobility on a Lipid Membrane. Nano Lett. 2016, 16, 4382–4390.

(48) Müller, M.; Lauster, D.; Wildenauer, H. H. K.; Herrmann, A.; Block, S. Mobility-Based Quantification of Multivalent Virus-Receptor Interactions: New Insights into Influenza A Virus Binding Mode. Nano Lett. 2019, 19, 1875–1882.
