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Membrane-containing virus particles exhibit the mechanics of a composite material for genome protection†

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The protection of the viral genome during extracellular transport is an absolute requirement for virus survival and replication. In addition to the almost universal proteinaceous capsids, certain viruses add a membrane layer that encloses their double-stranded (ds) DNA genome within the protein shell. Using the membrane-containing enterobacterial virus PRD1 as a prototype, and a combination of nanoindentation assays by atomic force microscopy and finite element modelling, we show that PRD1 provides a greater stability against mechanical stress than that achieved by the majority of dsDNA icosahedral viruses that lack a membrane. We propose that the combination of a stiff and brittle proteinaceous shell coupled with a soft and compliant membrane vesicle yields a tough composite nanomaterial well-suited to protect the viral DNA during extracellular transport.

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1. Introduction

Virions (infectious virus particles) are nano-sized carriers of information whose objective is to infect a host cell and generate progeny that can, in turn, repeat the infection cycle. When traveling outside of the host cell, virions may encounter harsh environmental conditions. Knowledge of the deformability (stiffness, k), the energy required for mechanical failure (toughness, T), and the limits to fatigue for virions has been acquired through studies using atomic force microscopy (AFM), but mainly for viruses that do not have an internal

membrane. 1-4 The mechanical properties of these virions ulti-

Guided by the available genetic, biochemical, and structural information, we chose to utilize (i) wild type PRD1 (wt PRD1),

of a proteo-lipidic tube. 12,13

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mately recapitulate the molecular interactions across the viral icosahedral shell.^{5,6} There is no comparable information available for icosahedral virions in which an internal membrane vesicle surrounds the densely packaged DNA and, to our knowledge, such viruses with RNA genome are yet to be identified. Defining how a membrane and capsid provide protection to the genome within will provide new insights into the function and properties of biological materials, thereby inspiring the design of novel nanostructures. PRD1, the prototype for icosahedral dsDNA viruses with an internal membrane, is a large and complex enterobacterial virus (family Tectiviridae; \sim 65 nm mean diameter and triangulation number pseudo-T =25, Fig. 1a). Its crystal structure revealed the molecular interactions across the major capsid proteins (MCPs) P3 (395 residues) and with the membrane.^{7,8} P3 trimers (capsomers) are arranged in the group of nine (GON) layout in the virion facets (Fig. S1a†). The minor capsid protein P30 cements the edges of the icosahedron. Eleven of the twelve vertices are capped by peripentonal P3 trimers and by spike complexes⁷⁻¹¹ (Fig. 1a and S1a†). The internal membrane enclosing the densely packaged dsDNA is composed of a roughly equal mass of lipids and membrane proteins.8 The twelfth vertex is the unique, membrane-embedded portal used for DNA packaging into the procapsid and later for DNA ejection via the formation

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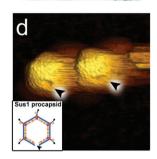
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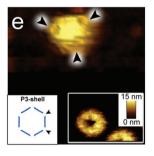
Membrane

Unique vertex

wt PRD1

Paper





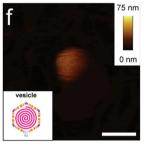


Fig. 1 (a) Schematic of PRD1 highlighting the main structural features of the virus particle. (b) Protein composition of wt PRD1 and subviral particles analysed by SDS-PAGE and Coomassie staining. Left: Molecular masses (M; kDa). Right: Positions of the most abundant proteins; the vertical black line refers collectively to membrane proteins (MPs, including P14, P16, P18, P20, P22, P31 and P32). Cementing protein P30 identified in the P3-shell by mass spectrometry is indicated by a black arrowhead. (c-f) Atomic force micrographs of wt PRD1 and subviral particles with scale bar (100 nm) and z range (75 nm) indicated in (f). (c) Wt PRD1 featuring different particle orientations. Inset: Crystal structure of wt PRD1.7 Yellow, green, cyan and blue denote the P3 pseudo-hexameric capsomers composing the icosahedral asymmetric unit; white lines delineate a virus facet; white pentagons, triangles, and ovals indicate icosahedral 5-, 3- and 2-fold symmetry axes, respectively; (d) Sus1 procapsids (no DNA within). Arrowheads indicate the visible depression due to the missing packaging portal in the unique vertex. Inset: Schematic of Sus1 procapsid; (e) P3-shell. Arrowheads highlight the depressions visible at all vertices due to the absence of the peripentonal capsomers and vertex complexes. Right inset: The star-shaped vertex depression at higher resolution (z range: 15 nm). Left inset: Schematic of P3-shell. Arrowheads indicate some of the de-capped vertices. (f) DNA-filled vesicle. Inset: Schematic of vesicle represented as an icosahedron for clarity and consistency with the wt PRD1 schematic representation - this shape, however, might not be retained in the purified membrane-vesicle.

(ii) a PRD1 mutant that forms procapsids devoid of DNA (Sus1 procapsid), (iii) the icosahedral P3-shell composed of MCP P3 and the minor capsid protein P30 (Fig. 1b and Table S1†), but lacking the pentons and peripentonal capsomers (P3-shell), and (iv) proteo-lipidic membrane vesicles enclosing a complete

genome (vesicle; Fig. 1b). We used AFM to examine the mechanical responses of these particles in an aqueous environment by assessing their stiffness and yield behaviour under an applied force and used finite element modelling, where possible, to aid the analysis.

Experimental

2.1 PRD1 specimen production

Wt PRD1 and mutant PRD1 and mutant *sus1* [amber mutation in gene *IX*] were propagated on *Salmonella enterica* serovar Typhimurium strain DS88 or on suppressor strains PSA(pLM2) or DB7156(pLM2). Cells were grown in Luria–Bertani (LB) medium at 37 °C.

For the production of wt and mutant phage particles, DS88 cells were infected using a multiplicity of infection of 8–10. For mutant particle production, infected cells were collected 15 min after infection (Sorvall rotor F12, 5000 rpm, 10 min, 22 °C) and transferred to a fresh pre-warmed medium. Virus particles were purified by polyethylene glycol–NaCl precipitation and rate zonal ultracentrifugation in sucrose (Sorvall rotor AH629), as previously described. For AFM, wt PRD1 and Sus1 procapsids were further purified by equilibrium ultracentrifugation in sucrose (Sorvall rotor AH629). The particles were concentrated by differential centrifugation (Sorvall rotor T647.5, 32 000 rpm, 2 h, 5 °C). A buffer containing 20 mM potassium phosphate pH 7.2 and 1 mM MgCl₂ was used for purification and resuspension.

For P3-shell preparation, the rate zonal purified Sus1 mutant particles (2 mg ml $^{-1}$ in 20 mM Tris-HCl, pH 7.2, 1 mM MgCl $_2$) were treated with 1% (w/v) sodium dodecyl sulfate (SDS) for 15 min at 25 $^{\circ}$ C. 20 P3 shells were isolated by rate zonal centrifugation in a linear 5–20% (w/v) sucrose gradient using the Tris buffer (Sorvall rotor AH629, 24 000 rpm, 1 h 45 min, 20 $^{\circ}$ C). The particles were concentrated by centrifugation (Sorvall rotor T865, 34 000 rpm, 4 h, 5 $^{\circ}$ C) and resuspended in Tris buffer.

For membrane vesicle preparation, the rate zonal purified Sus607 particles devoid of the major membrane protein P11 (1 mg ml⁻¹ in 20 mM Tris-HCl, pH 7.2) were treated with 2.5 M GuHCl²¹ and membrane vesicles were purified by equilibrium centrifugation in a linear 20–70% (w/v) sucrose gradient (Sorvall rotor TH641, 22 000 rpm, 16–18 h, 20 °C). Protein concentrations were determined by Bradford assay.²² Particles were analyzed using sodium dodecyl sulfate polyacrylamide gel electrophoresis²³ (SDS-PAGE; 16% acrylamide; Fig. 1b).

Virus particles were stored at 4 °C for no more than 4 weeks. During this time, more than 95% of the wt PRD1 particles remained intact and without the loss of their genome as visualized by cryo-electron microscopy (cryo-EM) 2D imaging [a JEM-2200FS (JEOL) transmission electron microscope equipped with an UltraScan 4000 SP $4k \times 4k$ camera (GATAN)] (Fig. S2†). Sus1 procapsid, P3-shell particles and vesicles were similarly stored and also used within this time frame (Fig. S2†).

2.2 Protein identification by mass spectrometry

Silver stained²⁴ protein bands were cut out of the polyacrylamide gel and "in-gel" digested. Cysteine bonds were reduced with 0.045 M dithiothreitol (Sigma-Aldrich, USA) for 20 min at 37 °C and alkylated with 0.1 M iodoacetamide (Fluka, Sigma-Aldrich, USA) at room temperature. Samples were digested by adding 0.75 µg trypsin (Sequencing Grade Modified Trypsin, V5111, Promega) overnight at 37 °C. After digestion, peptides were purified with C18 microspin columns (Harvard Apparatus) according to the manufacturer's protocol. The dried peptides were reconstituted in 30 µl of buffer A [0.1% trifluoroacetic acid (TFA) in 1% acetonitrile (ACN)].

Liquid chromatography coupled to tandem mass spectrometry (LC-MS/MS) analysis was carried out on an EASYnLC1000 (Thermo Fisher Scientific, Germany) connected to a Velos Pro-Orbitrap Elite hybrid mass spectrometer (Thermo Fisher Scientific) with a nano-electrospray ion source (Thermo Fisher Scientific). The LC-MS/MS samples were separated using a two-column setup consisting of a 2-cm C18-Pepmap trap column (Thermo Fisher Scientific), followed by a 15-cm C18-Pepmap analytical column (Thermo Fisher Scientific). The linear separation gradient consisted of 5% buffer B in 5 min, 35% buffer B in 60 min, 80% buffer B in 5 min and 100% buffer B in 10 min at a flow rate of 0.3 µl min⁻¹ (buffer B: 0.1% TFA acid in 98% acetonitrile). Six microliters of sample were injected per LC-MS/MS run and analyzed. Full MS scan was acquired with a resolution of 60 000 in the normal mass range in an orbitrap analyzer and followed with CID-MS2 top 20 most intense precursor ions within the ion trap (energy 35). Data were acquired using LTQ Tune software.

The acquired MS2 scans were searched against the enterobacteria phage PRD1 (NCBI) protein database using the Sequest search algorithms in Thermo Proteome Discoverer. The allowed mass error for the precursor ions was 15 ppm and for the fragment 0.8 Da. A static residue modification parameter was set for carbamidomethyl +57 021 Da (C) of the cysteine residue. Methionine oxidation was set as dynamic modification +15 995 Da (M). Only full-tryptic peptides were allowed for the maximum of one missed cleavage.

2.3 Surface preparation and immobilization of PRD1 particles

Freshly cleaved surfaces (~1 cm²) of mica (Ted-Pella Inc., CA, USA) were functionalized with 3-(2,2-aminoethylamino)ethylaminopropyltrimethoxysilane (APTES; Sigma-Aldrich) by adapting a previously described procedure.²⁵ Briefly, mica was left to incubate with 30 µL of APTES in a 4 l gas incubator for 2 h. A small amount of water (10 µl) was deliberately added to generate a root-mean-square surface roughness of 1.7 to 2 nm, which proved beneficial to particle attachment. Following the APTES functionalization, surfaces were covered with 100 µl Hepes buffer (10 mM HEPES, pH 7.4, 150 mM NaCl, 2 mM MgCl₂) containing ~10 μg virus particles (wt PRD1, Sus1 procapsid, or P3-shell). PRD1 vesicles were attached to freshly cleaved mica instead of APTES-coated mica. Virus particles were

allowed to adhere for 30 min, and excess particles were removed by washing with buffer.

2.4 AFM imaging and nanoindentation

All PRD1 particles were found to attach as a monolayer to the prepared mica surfaces that could be readily visualized by AFM. Imaging and nanoindentation measurements were carried out using a MultiMode 8 AFM with a Nanoscope V controller (Bruker, CA, USA) at room temperature in Hepes buffer. Wt PRD1, Sus1 procapsids, and P3-shells were analysed with sharpened triangular Si₃N₄ cantilevers with a nominal spring constant of 0.7 N m⁻¹ (ScanAsyst-Liquid; Bruker). For PRD1 vesicles, Si₃N₄ cantilevers with a nominal spring constant of 0.1 N m⁻¹ (AC40; Bruker) were used. The real spring constants of the cantilevers were measured using the thermal noise method26 as implemented in the NanoScope software and were found to be close to the nominal values. Imaging was performed in Peak Force Tapping mode, with a typical tapping amplitude of 40 nm and a driving frequency of 2 kHz. For wt PRD1, Sus1 procapsids, and vesicles, the peak force was usually 100 pN; for P3-shells, it was 80 pN. Under these conditions, the image quality was sufficient to reliably identify the virus particles (and in some cases, their orientation and surface sub-features), and in general, intact PRD1 particles did not degrade or break upon repeated imaging. On already degraded particles, gradual further deterioration was observed, and particles were also found to detach from the surface. Images were plane fitted when required, using Gwyddion software (http://gwyddion.net/) without the application of noise filtering or sharpening.

Nano-indentation measurements were performed at individually selected particles using the 'point-and-shoot' function within the NanoScope software. Briefly, the area of interest was first imaged to localize the virus particle; the 'point-and-shoot' function was then activated and force curves were taken at the particle centre, subsequently, the area was imaged once more to verify successful nano-indentation. The accuracy of localization of the particle centre was found to be limited by piezo drifts and estimated to be within 5 nm. Force vs. distance (F/z)curves were acquired at a constant approach velocity of 200 nm s⁻¹. The approach and retract distances were 100 nm, corresponding to a total time of 1 s per complete approach and retract cycle. The maximal load was 4 nN, except for P3-shells, where the maximal load was lowered to 2 nN. F/z curves were analysed using the NanoScope software.

2.5 Force curve analysis

2.5.1 Selection of force curves. At maximal loads of 2 to 4 nN, the AFM tip typically reached the underlying support following the indentation and/or fracture of all PRD1-derived particles. We used the z distance travelled between the onset of a repulsive force and the closest approach as an indicator for successful indentation at a central particle position. The closest approach typically corresponded to hard-wall contact, though in some cases it was short of hard-wall contact by a few nm as estimated from force curves with hard-wall contact

acquired shortly before/after on a nearby mica area and taking advantage of the AFM's closed loop z scanner. Force curves showing distances of 60 ± 10 nm were retained for further analysis of wt PRD1, Sus1 procapsids, and P3-shells; and distances of 28 \pm 5 nm were considered acceptable for DNA-filled vesicles. When imaging vesicles, we found a second class of objects with a height of 12 ± 7 nm and lateral dimensions comparable to the DNA-filled vesicles. Since negative stain EM showed also deformed vesicles in some cases (Fig. S2†), most likely this second class represents vesicles that lost their DNA and had flattened on the surface. These flattened objects were not further analyzed.

To avoid including particles that might have been displaced or changed orientation upon indentation, force curves where force levels dropped and remained below 500 pN over distances of 20 nm and more before hard-wall contact were also discarded from analysis. A representative force curve for each PRD1 particle type with AFM micrographs before and after indentation is shown in Fig. S3,† and additional force curves illustrating sample to sample variations are shown in Fig. S4.†

2.5.2 Determination of yield point and stiffness. After the first contact between the AFM probe and a PRD1 particle at the contact point z_c , the initial monotonous force (compression) was followed by an extended and steep drop (fracture), with subsequent, typically minor, compression and fracture phases (Fig. S4†). The onset of the first drop, defined as the yield point z_v , is determined by the yield force $F_v = F(z_v)$ and the yield strain $\varepsilon_v = (z_c - z_v)/h$, where h is the mean particle height. Neglecting the lowest forces (F < 200 pN), which would correspond to a non-linear Hertzian regime of capsid compression,²⁷ the major part of the first compression phase for wt PRD1, the Sus1 procapsid, and the P3-shell could, in general, be approximated well by a straight line - with the exception of short stochastic slips that we interpret as microfractures (Fig. S5†). Whilst for hollow-shell particles (Sus1 procapsid and P3-shell), the linear regime can be attributed to elastic shell bending, ²⁷ we focused analysis on this regime also for the full particles (wt PRD1 and vesicle). The particle stiffness k was determined from the slope in the initial compression phase, i.e., the slope up to the force at which the first slip or the fracture occurred. In any case, linear fits were not extended beyond 1500 pN (or strains above 10%) to avoid bias by non-linearity at larger compressions. Likewise, the stiffness was not quantified if the first slip occurred below 500 pN to avoid the large uncertainties associated with fitting a line to a small dataset. Stiffness values for vesicles were extracted from linear fits to the compression curves that did not extend beyond 600 pN, but typically up to a maximum of 50% strain, as strains below 15-20% fell within the noise limit of the measurements.

- 2.5.3 Estimation of particle toughness. Toughness T was defined as the amount of energy E_v per volume V that the PRD1 particles can withstand before breaking. Approximating the virus as a sphere of radius R, and with $T = E_v/V$, $E_v \approx F_v(z_c - z_v) =$ $F_{\nu} \varepsilon_{\nu} h$ and $V = 4\pi/3 R^3 \approx \pi/6 h^3$, we obtain $T \approx 6/\pi F_{\nu} \varepsilon_{\nu} / h^2$.
- 2.5.4 Statistical analysis. Origin data analysis and graphing software was used for statistical analysis of results (OriginLab,

Northampton, MA). Gaussians were fitted to the histograms in Fig. 2 to extract the means and standard deviations (s.d.). Oneway ANOVA tests were performed to determine the statistical significance of the differences of yield force and stiffness across the PRD1-derived particle populations. The stiffness of wt PRD1 against Sus1 procapsid yielded a P-value ≤ 0.001 (**), while the stiffness, yield force and yield strain of both PRD1 and Sus1 procapsid against P3-shell showed a P-value ≤ 0.0001 (***).

2.6 Finite-element analyses

To test how the membrane vesicle affects the stiffness and stability in a composite system of a proteinaceous capsid shell with an underneath membrane vesicle, modelling of the mechanical response of PRD1 was performed using a continuum-mechanics finite-element analysis (FEA) with the software ABAQUS 6.14 (ABAQUS, Fremont, CA) (Fig. S6 and S7†). In this approach, the proteinaceous capsid and the proteolipidic vesicle were represented as mechanically isotropic spherical shells, i.e., neglecting the structural details of each shell (see section 3.3). This method can effectively deal with the architectural complexity introduced by the presence of the vesicle whilst the number of adjustable parameters is kept small.

3. Results and discussion

Imaging PRD1 assemblies at the single particle level

When immobilized, the wt particles oriented with a 3-fold, 2-fold or 5-fold symmetry axis normal to the supporting surface (Fig. 1c). The preferred orientation (>67% of the particles) was with a facet down (i.e., a 3-fold axis normal to the surface). The average height was 67.8 ± 2.5 nm (mean \pm s.d.) for wt PRD1, 66.9 \pm 2.7 nm for the Sus1 procapsid (n = 75 each; Fig. 2a and b, and 62.8 \pm 3.5 nm for the P3-shell (n = 103; Fig. 2c), in good agreement with electron-microscopy and X-ray data^{7,28,29} (Fig. S1b†). Strikingly, we observed on the Sus1 procapsid a mild circular depression, ~13 nm in diameter at ~8% of the visualized vertices (n = 103) (Fig. 1d and S8†). This percentage, combined with the estimated dimensions, defines at the single-molecule level that this feature is the unique vertex that lacks the external part of the packaging portal. 12 This, however, was not detected on wt PRD1. Indeed, the structural difference in the capsid context between the 'wild type' unique vertex and the remaining 11 vertices is much smaller than that between the unique vertex and the other 11 vertices in the procapsid, 12 and apparently, it is too small to be resolved with our AFM set-up. P3-shells showed holes, ~25 nm wide, at each vertex (Fig. 1e) consistent with the missing peripentonal P3 capsomers, spike complexes, and internal vesicles^{28,29} (Fig. S1b†). The forces needed to be lowered from ~100 to ~80 pN to enable imaging of P3-shells indicating that these particles are more sensitive to breakage compared to wt PRD1. By contrast, vesicles displayed a featureless surface and a height of 28.2 \pm 4.1 nm (Fig. 1f and 2d). This height is less than the diameter of DNA-containing vesicles (~35 nm),8 indicating that vesicles readily deform when immobilized.

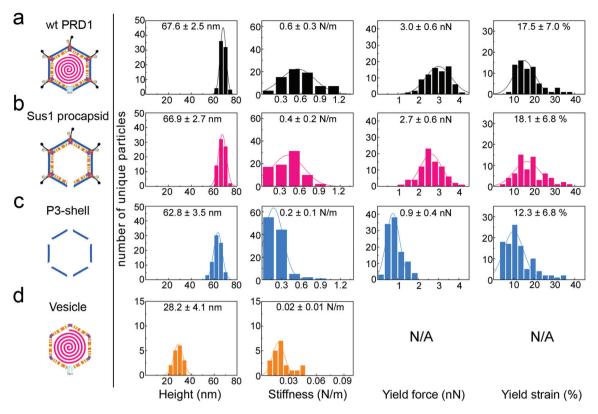


Fig. 2 Histograms of various properties of the four viral particle types, reporting mean values and standard deviations for each. y-Axis: number of unique particles (75 for wt PRD1 and Sus1 procapsid, 103 for P3-shell and 21 for vesicles). x-Axis: property denoted below each column. Specimen heights closely match the particle dimensions previously obtained by averaging techniques, except for the vesicle, which is slightly flattened upon immobilization. Yield force and strain are not shown for vesicles, as these do not have a defined yield point.

Stiffness of PRD1 particles

The AFM force vs. distance (F/z) curves (Fig. S3 and S4†) generated by the nanoindentation testing of individual particles quantified the resistance to small elastic deformation (stiffness, k), as well as the force and strain at the point of mechanical failure (yield force, F_v ; yield strain, ε_v) (Fig. 2 and 3a-c).

The F/z curves typically exhibited a relatively small nonlinear regime at the smallest indentation forces (F < 200 pN), which is likely to represent the Hertzian deformation of the capsid shell.²⁷ This was followed by an extended linear regime, justifying the quantification of elastic properties in terms of the stiffness k. For quantitative stiffness analysis, we considered this linear regime for strains up to 10%, which was well below the yield point. For quasi-spherical shells such as the Sus1 procapsid and the P3-shell, this linear regime can be associated with shell bending.²⁷

Wt PRD1 possessed the greatest stiffness (0.57 ± 0.03) N m⁻¹, mean \pm s.e.m.), followed by the Sus1 procapsid (0.39 \pm 0.02 N m^{-1}) and the P3-shell ($0.22 \pm 0.01 \text{ N m}^{-1}$; Fig. 3a). Most likely, the enhanced resistance of wt PRD1 to elastic deformation arises from the pressure exerted by the DNA.^{8,30} PRD1 genome packaging produces a radial expansion of the internal membrane (e.g., the radius of the outer leaflet increases by ~6%), which presses the vesicle closer to the capsid. 12,28,30

The DNA-filled vesicle displayed the least stiffness (0.022 \pm 0.002 N m⁻¹; Fig. 3a), indicating that its direct effect on virion stiffness is marginal but that it contributes to virion stiffness by transmitting pressure from the DNA to the capsid. The volume inside the rigid capsid is reduced upon indentation accentuating the effect of DNA pressure on the stiffness of wt PRD1. In contrast, the soft membrane can stretch upon indentation, thus reducing any effect of DNA pressure on the vesicle stiffness.

3.3 Modelling of PRD1's shell elasticities

To confirm the above findings on PRD1's elastic properties, we confronted the experimental data with continuum-mechanics finite-element computational modelling (see Experimental and Fig. S6†). In this simulation, the particle was placed between a rigid plane and a rigid indenter (Fig. 4a). The displacement of the rigid plane was constrained, and the indenter apex was modelled as a sphere of 10-nm radius positioned coaxially with the virus-derived particle in the direction normal to the plane. For indentation, the indenter applied a force in the direction normal to the plane. The contact between all bodies was assumed to be hard in the direction normal to the contact (*i.e.*, interpenetration was not allowed)

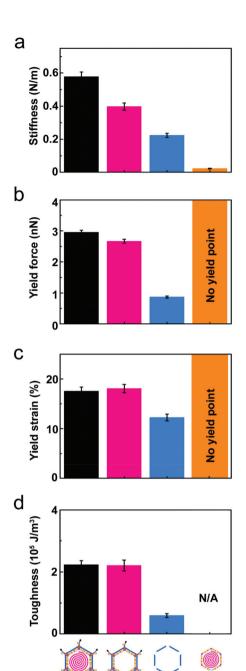
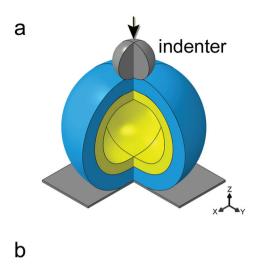


Fig. 3 Comparison of the average mechanical properties across the different PRD1-derived particles. In all panels: black bar, PRD1 wt; pink bar, Sus1 procapsid; blue bar, P3-shell; orange bar, vesicle. (a) Stiffness (k); (b) yield force (F_y); (c) yield strain (ε_y); (d) toughness (T). All panels report the mean values and standard errors of the mean that were derived from the data shown in Fig. 2 (for the vesicle, calculation of toughness is not possible because there is no defined yield point).

and frictionless in the tangential direction (*i.e.*, relative displacement and rotation were allowed with no constraint).

To extract material properties from the experimental data, we first considered the proteinaceous capsid individually. For the capsid, we estimated an outer radius R = 33.2 nm and a thickness d = 8 nm from the rotationally averaged electron density maps of the PRD1 virion. ^{28,30} By treating the capsid as



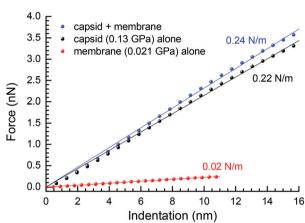


Fig. 4 (a) Schematic of the spherical two-shell model (outer protein shell in blue, inner vesicle shell in yellow) for the finite-element modelling; three-quarters of the spheres are shown with the indenter apex represented as a grey sphere with the direction of the applied force as indicated by the black arrow; in the bottom right corner, the Cartesian coordinate system: (b) predicted curves of force vs. indentation for a protein shell (with E = 0.13 GPa, representing the P3-shell; black dots), a vesicle shell (with E = 0.021 GPa, representing the proteo-lipidic vesicle; red dots) and a composite of these protein and vesicle shells (blue dots). In matching colours: lines are linear fits through the origin, and texts the stiffness values corresponding to the slopes. The stiffness values of protein and vesicle shells match the experimental data for P3-shell and vesicle (Fig. 3a), respectively, confirming the correct choice of Young's modulus values; the stiffness of the composite is well approximated by the sum of the stiffness values of the constituent shells and thus only marginally larger than the stiffness of the protein shell alone.

isotropic and linearly elastic, its properties can be described by two independent parameters: Young's modulus E and the Poisson ratio ν . By neglecting a non-linear regime at very small strains (ε < 3%), the predicted relationship between force and particle indentation δ was approximately linear and scaled with the Young's modulus E for indentations up to 2d (ε up to 24%; Fig. S6a†). Our results were rather insensitive to the Poisson ratio (Fig. S7a-c†), and we thus fixed ν = 0.4.

These features are consistent with previous computations by others and with the predictions of thin-shell theory, 4,31,32

and thus validate the numerical model. The linear relationship between force and indentation is also consistent with our experimental data (Fig. S3b, e, h, and S4a-d†), where we note that the strain regime of ε < 10% used for the analysis of experimental data lies well within the range over which theory predicts a linear response. This implies that stiffness analysis is far away from any buckling transition.³³ It justifies the use of linear elasticity in our simple theoretical model, and also the use of stiffness $k = F/\delta$ to characterize the shell's elastic properties. With $F/E \sim \delta$ and $k = F/\delta$, it is also clear that $k \sim E$, and a linear fit to the data in Fig. S6a† gives $k \approx 1.65$ nm $\times E$. With this equation, we estimate E = 0.13 GPa for the P3-shell from the experimentally determined mean stiffness value for this particle ($k = 0.22 \text{ N m}^{-1}$; Fig. 3a).

A recent computational modelling study on a smaller nonmembrane-containing virus has shown that capsid proteins can dynamically re-structure upon capsid indentation with appreciable effects on capsid elasticity compared to an idealized homogeneous shell.³⁴ Our experimental data do not allow deconvoluting these effects. However, the extended linear regime in the F/z curves observed experimentally for the P3-shell and the Sus1 procapsid (Fig. S4c and d†) indicates that the bending elasticity of the PRD1 capsid shell remains appropriately described by a Young's modulus (where this would effectively include the possible re-structuration effects).

The elasticity of the proteolipidic membrane was estimated analogous to that of the proteinaceous capsid, assuming an outer radius identical to the inner radius of the capsid (R =25.2 nm) for direct contact of the two shells, and a membrane thickness d = 5.5 nm from a rotationally averaged electron density map.³⁰ Fig. S6b† shows the dependence of F/E on vesicle deformation for these geometrical parameters, from which $k \approx 1.07$ nm \times E can be derived. Assuming to a first approximation that the stiffness of the membrane shell is similar to the experimentally accessible value for the genomecontaining vesicle ($k = 0.022 \text{ N m}^{-1}$; Fig. 3a), we can estimate E = 0.021 GPa.

To predict the elastic behaviour of the composite capsidmembrane system, we modelled a system of two concentric shells with the inner shell adopting the geometry and Young's modulus of the membrane and the outer shell adopting the geometry and Young's modulus of the P3-shell (Fig. 4a). The stiffness of this composite system was $k = 0.24 \text{ N m}^{-1}$, that is the presence of the vesicle enhanced the stiffness only marginally, by about 10%, compared to the P3-shell alone. More generally, the stiffness values shown in Fig. 4b exemplify that the stiffness of the composite (0.24 N m⁻¹) is well approximated by the sum of the stiffness values of the constituent shells (0.22 N m⁻¹ + 0.02 N m⁻¹). The small enhancement in stiffness was virtually independent of the Poisson ratios of both the capsid and membrane (Fig. S7d†).

The above modelling exercise provides reasonable predictions about the trends that can be expected based on the experimental AFM-derived shell elastic mechanical properties. Here, we have operated with the P3-shell as a reference system because experimental data for this single-shell system were

readily available. Using the above-identified stiffness relationship, we can now also estimate the Young's modulus of the complete capsid shell from the experimentally determined stiffness values of the Sus1 procapsid and the membrane. The closure of 11 of 12 vertices by additional proteins enhances the elasticity of the Sus1 procapsid shell compared to the P3-shell, whilst a further enhancement of the capsid elasticity by the unique vertex - missing in the Sus1 procapsid - is likely to be marginal. $k_{\text{capsid}} \approx k_{\text{capsid+membrane}} - k_{\text{membrane}} \approx 0.39 \text{ N m}^{-1}$ 0.02 N m⁻¹ = 0.37 N m⁻¹ (Fig. 3a) and $E \approx k/1.65$ nm (Fig. S5a†) give $E \approx 0.22$ GPa, a value comparable to that for some non-enveloped and enveloped viral capsids.35

Indeed, the two-shell modelling confirmed that the stiffness is only marginally affected, by less than 10%, by an internal soft membrane contacting the capsid (Fig. 4b). The reduced stiffness of the P3-shell (E = 0.13 GPa) is likely due to the absence of the stabilizing pentons and peripentonal capsomers.

3.4 Mechanical stability of PRD1 particles

The presence of the genome and the ensuing particle stiffening had no appreciable effect on the mechanical stability of the virion: the yield force and yield strain of wt PRD1 (F_v = 3.0 ± 0.1 nN, $\varepsilon_{\rm v} = 17.5 \pm 0.8\%$; mean \pm s.e.m.) and the Sus1 procapsid ($F_v = 2.7 \pm 0.1$ nN, $\varepsilon_v = 18.1 \pm 1.4\%$) were similar (Fig. 3b and c). This is in contrast to other phages, such as phage λ , where the DNA augments both stiffness and mechanical stability.³⁶ As for the P3-shell, it was much more sensitive to breakage than wt PRD1 or the Sus1 procapsid ($F_v = 0.9 \pm 0.1$ nN, $\varepsilon_v =$ $12.3 \pm 0.7\%$; Fig. 3b and c).

The mechanical failure of the PRD1 particles upon indentation was frequently accompanied by the loss of capsomers from the capsid shell (Fig. S3a-c,† insets). In addition, F/z curves of wt PRD1, the Sus1 procapsid, and the P3-shell revealed slip events coincident with the occurrence of microfractures during force loading likely reflecting the local displacements of MCPs (Fig. S5†). While all three particle populations presented a similar density of micro-fractures (average 1.3 per nN of applied compressive force), wt PRD1 and the procapsid withstood more of these fractures before yielding.

Previous studies on binary component viral systems genome encapsulated by a protein shell - have highlighted the role of DNA or RNA in contributing to the capsid stiffness where the mechanical reinforcement is achieved by the genome anchoring the protein shell from the interior.³⁵ In other cases, such as the herpes simplex virus type 1 (HSV-1) nucleocapsid, the stiffness and yield force remain practically the same whether the particle is fully packaged or devoid of the genome, and stabilizing viral proteins appear to be responsible for this assembly type. 37,38 In PRD1, the Sus1 procapsid and the mature particle display similar yield forces whereas the relative increase in the stiffness of the wt PRD1 can be attributed to the presence of DNA. The packaging of the genome leads to an expansion of the membrane-vesicle increasing the membrane's interactions with the capsid proteins. 12,28-30

3.5 Toughness analysis of PRD1 and other icosahedral dsDNA viruses

The resistance to material fracture is typically expressed as toughness T, here defined as the amount of energy per unit volume that can be absorbed before mechanical failure. The toughness of PRD1, 2.2×10^5 J m⁻³, is relatively high compared to that reported for other icosahedral dsDNA viruses (see Fig. 3d and 5, and Table S2†). Thus, PRD1 together with human adenovirus is one of the toughest among all icosahedral dsDNA viruses. 7,39-41 Adenovirus shares a common MCP fold and assembly mechanism with PRD1; however, whilst it lacks a membrane, it possesses a complex set of cementing proteins stabilizing the structure - about 300 in total (composed of sixty copies of each of the proteins IIIa, V, VI, VIII and IX) instead of the mere 60 copies of the single protein species P30 guiding the assembly in PRD1.7,42

Altogether, these comparisons highlight that the layered complex of the capsid and membrane vesicle relieves the genome from a stabilizing role and endows PRD1 with remarkably high mechanical stability.

To explore how the PRD1 vesicle and capsid together influence toughness, we compared the toughness of the PRD1derived particles: the Sus1 procapsid, the P3-shell, and the vesicle. The P3-shell ($T = 0.54 \times 10^5 \text{ J m}^{-3}$) was more susceptible to mechanical failure than the Sus1 procapsid ($T = 2.1 \times$ 10⁵ J m⁻³; Fig. 3d). The fact that the P3-shell's toughness was ~4-fold less but its stiffness was only 2-fold less revealed its rather stiff, brittle nature (Fig. 3a). The P3-shell lattice is held together by the (C-I type and C-II type) interactions established by the GON within each facet and along the facets via the MCPs C-termini and by the P30 proteins⁷ (Fig. S1a†). The relative ease of breakage of this lattice might facilitate morphological corrections as the capsomers assemble on the vesicle mould during the procapsid formation. Closing the icosahedral vertices with the peripentonal capsomers and penton proteins (Fig. S1a†) and plugging the unique vertex with the portal complex produce a stable procapsid that can withstand dsDNA translocation powered by the packaging ATPase P9.12

More generally, the brittle nature of the capsid is not only manifested in the small yield force of the P3-shell but also in the above-mentioned microfractures. The DNA-filled vesicles, on the other hand, did not show a clear yield point and typically recovered their original shape even after strains exceeding 60%, which indicated that they were ductile and very soft (Fig. 3a).

Thus, comparative analysis with other dsDNA viruses indicates that the toughness of PRD1 is superior to most other dsDNA viruses with comparable capsid organization but lacking the membrane, and is only rivalled by adenovirus - a non-membrane-containing virus - which is exceptionally rich in cementing proteins stabilizing the capsid (Fig. 5).

Composite material model of PRD1

Our findings indicate that the capsid is stiff and brittle, whereas the membrane vesicle is ductile and soft. Might the hierarchical combination of these contrasting material properties be responsible for the high toughness of PRD1 particles? Quantifying accurately the contribution of the membrane vesicle to the overall toughness displayed by the PRD1 particle remains a challenge. To our knowledge, no PRD1 intact capsid particles can be biochemically or genetically produced that would allow the AFM probing, and yet the resulting information might be still limited for an exhaustive modelling of the yield behaviour of the full capsid as a brittle material. However, in nature, other macroscopic systems use similar arrangements to create tough materials. One such illustrative example is the coat protecting bird egg, in which a stiff, yet

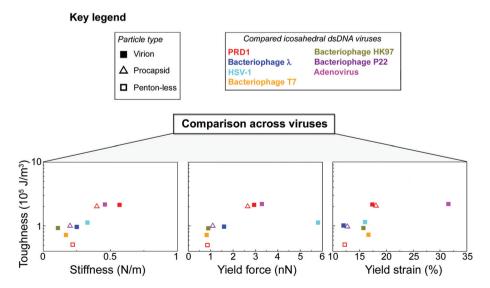


Fig. 5 PRD1's mechanical properties compared to other dsDNA viruses. Key for the viruses and particle types shown in the other panels (in the case of HSV-1 virion, we refer to its nucleocapsid). Comparison across viruses via Ashby plots of toughness (y-axis in log scale) vs. stiffness, yield force and yield strain (x-axis), respectively (see the Table S2† for references).

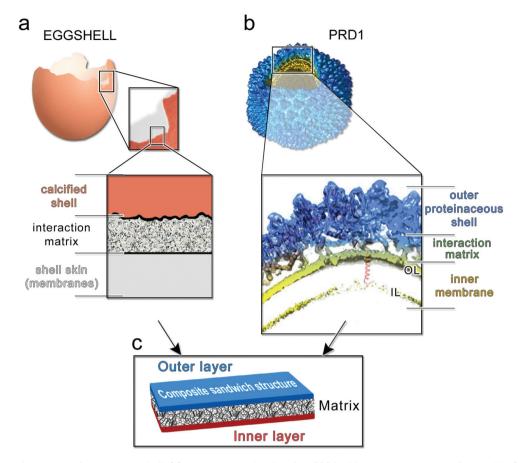


Fig. 6 (a) Multilayered structure of an avian eggshell; (b) cryo-electron density of wt PRD1 with an octant removed displayed in Chimera⁴⁵ to show (inset) the layered/composite structure of the virion with the proteinaceous shell in blue, the membrane-vesicle in yellow-lime (OL: outer leaflet; IL: inner leaflet) and with the horizontal black lines marking the distinct layers and the interacting matrix region between the capsid and the membrane. As cartoon representation in red fitted in density the P3 MCPs with N-terminal \alpha-helices interacting with the OL and the P16 transmembrane protein crossing the membrane vesicle. The dsDNA has been removed from within the membrane vesicle for clarity; (c) schematic of a composite sandwich material to which (a) and (b) recapitulate.

brittle, calcified shell is bonded to a soft and compliant proteinaceous inner skin⁴³ (Fig. 6a).

Altogether, these layers constitute a tough composite material that protects the progeny against mechanical stress while carrying out other essential functions. Analogously, the PRD1 capsid and vesicle are bonded to protect the integrity of the virion and its genome (Fig. 6b). Specifically, the connection between the capsid and membrane vesicle is made of polypeptide stretches: the capsid shell - composed of 240 copies of interacting trimers of the MCP P3 and glued at the icosahedral edges by the cementing protein P30 - anchors the membrane through several N-termini of MCPs P3; this connectivity is further augmented by the anchoring N-terminal transmembrane helix of protein P16 located at the base of the peripentonal MCPs P3 7 (inset in Fig. 6b).

Moreover, it is conceivable that during force loading onto the capsid, these connectors will act as nano-staples increasing the capability of the system of absorbing energy before mechanical failure (an additional energetic cost would be necessary to disrupt protein-membrane interactions⁴⁴).

Conclusions

Composites have evolved in nature driven by selection for the efficient use of materials, adaptability, and multi-functionality. Viruses can be seen as composite biological entities where nucleic acids, proteins and lipids assemble to produce functional particles for infection. We propose that, as in a composite sandwich material (Fig. 6c), an interfacial protein/polypeptide matrix in PRD1 generates a tight connection that mechanically couples the capsid and the membrane (Fig. 6b). The flexibility of this matrix and possibly also the fluidity of the membrane facilitate the displacement of the two shells relative to each other, and thus assist in maintaining the capsomers in place whilst allowing for the correction of small scale defects in the (re-)assembly process.

In summary, we here presented the nanomechanical characterization of a virus that features a membrane inside its capsid. The combination of a stiff, yet brittle, proteinaceous capsid with a soft proteolipidic vesicle in PRD1 facilitates multiple stages of the virus life cycle, including virus assembly,

mechanical protection for the extracellular virion, and the vesicle-to-tube transformation during DNA ejection. 13 From a broader perspective, it appears that the evolution of membrane-containing viruses has yielded, at the nanometer scale, composite properties comparable to those known for macroscopic natural materials, where the capsid and vesicle are bonded into a tough composite that protects the integrity of the virus and its genome.

Our findings provide both foundational quantitative information and inspiration that can encourage the engineering of tough nanoscale devices and particles capable of protecting fragile cargos.

Conflicts of interest

There are no conflicts to declare.

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