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Structure, dynamics and phase separation of measles virus RNA replication machinery

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Abstract

The measles virus replication complex represents a potentially important, but as yet relatively unexplored target for viral inhibition. Little is known about the molecular mechanisms that underpin replication and transcription in paramyxoviruses. In recent years it has become clear that conformational dynamics play an important role in paramyxoviral replication, and that a complete understanding of the viral cycle requires a description of the structural plasticity of the different components. Here we review recent progress in this direction, covering the dynamics of the nucleocapsid assembly process, high resolution structure and dynamics of protein:RNA interactions, and the investigation of the role of intrinsic conformational disorder in pre-assembly nucleoprotein/phosphoprotein complexes. Finally, we discuss the role of viral factories in the form of phase-separated membraneless organelles formed by measles virus phospho and nucleoproteins that promote the assembly of nucleocapsid structures.

Paramyxoviruses are non-segmented negative sense RNA viruses, including Measles (MeV), the most infectious human pathogen that still has a global mortality of 140,000 per year, as well as numerous emerging human pathogens with extremely high mortality rates. There are currently no effective treatments to combat MeV infection, whose impact is increasing throughout the world due to incomplete vaccine coverage [1]. The replication complex represents a promising, but relatively unexplored therapeutic target. Paramyxoviral replication machinery comprises the nucleoprotein (N), whose major role involves encapsidation of the viral genome, the RNA-dependent RNA polymerase (L) and its cofactor the phosphoprotein (P) [2–4].

Each MeV N protein binds to six nucleotides of the viral genome or antigenome during encapsidation, assembling into helical nucleocapsids that are thought to protect the viral RNA from recognition and degradation by the innate immune system. N is chaperoned in the unassembled form by P prior to encapsidation, in the so-called N⁰P complex, and is thought to co-localize N and L during the replication process [5,6].

N and P both present a significant amount of conformational plasticity, comprising structured domains as well as highly disordered domains. Approximately 30 and 75% of the primary sequences of N and P lack tertiary structure respectively (figure 1), and fall into the category of intrinsically disordered proteins. N consists of a folded domain (N_{CORE}), that binds the viral RNA in the nucleocapsid structure, preceded by disordered 34 and 17 amino acid (aa) N- and C-terminal arms (N_{ARM} and C_{ARM}), and followed by a 134 aa flexible domain (N_{TAIL}) [7] that remains dynamic in the context of assembled nucleocapsids [8]. The presence of N_{TAIL} appears to introduce a level of global flexibility in the nucleocapsid, resulting in an increased average distance between successive helical turns when the disordered domain is present [9,10]. Flanking regions of N_{CORE}, N_{ARM}, C_{ARM} and the N-terminal end of N_{TAIL}, stabilize the helical superstructure of nucleocapsids by binding to neighbouring N molecules upon interaction with RNA (figure 1). Deletion of the flanking regions of N_{CORE} also prevents assembly into nucleocapsid like particles (NCLPs).

The polymerase cofactor P is tetrameric, consisting of a central coiled-coil domain [3], whose structural properties have recently been linked to MeV gene expression [11], flanked by a long (304 aa) intrinsically disordered N-terminal domain (P_{TAIL}) [12] and a shorter (80 aa) disordered domain (P_{LOOP}) that is terminated by a small C-terminal three-helix bundle (XD) [13] which interacts with a short helical linear motif in N_{TAIL} [7,13,14]. This level of disorder is replicated, in morbilliviruses and other paramyxoviruses (P_{TAIL} in Nipah virus is 470 aa long). The N-terminal 30 amino acids of MeV P_{TAIL} are known to interact tightly with N, via a helix-kink-helix conformation [15], forming a heterodimeric chaperone (N^0P) that inhibits assembly of N into nucleocapsids (figure 1). The sites on N_{CORE} that interact with the neighbouring flanking arms in nucleocapsid structures coincide with the binding sites of the two helices of P_{TAIL} in the N^0P complex, suggesting an inhibitory mechanism that maintains N in the unassembled state.

The functional role of the remainder of this extensive disorder has remained unclear. The conserved presence of such long disordered domains in viruses is all the more remarkable considering the parsimonious nature of their genetic material. Intrinsic disorder in paramyxoviral proteins has been rendered yet more intriguing by the recent observation that disordered domains play important roles in the formation of membraneless organelles that appear to act as viral replication factories [16–18].

The combination of structured and highly flexible domains in functional complexes requires the application of multi-disciplinary approaches that are capable of capturing the atomic detail of both the structure and dynamics of these highly plastic assemblies [19]. Here we summarize recent structural, dynamic and functional descriptions of these highly dynamic assemblies using a combination of solution and solid-state structural biology.

In vitro reconstitution of the MeV nucleocapsid assembly process

One major reason for the relative lack of understanding of the RNA genome (and anti-genome) encapsidation process stemmed from the difficulty in isolating N, prior to assembly, due mainly to the high propensity of N to bind cellular RNA (for example when expressed in *E. coli* or insect cells) and spontaneously assemble into NCLPs. Experimentally this has allowed the

purification of MeV NCLPs and the determination of increasingly high-resolution structures of the N_{CORE}-RNA complex over the last 20 years [9,10,20,21], providing a great deal of insight into the mechanisms of stabilization described above, but the kinetic process of assembly remained poorly characterized. Inspired by the mechanism exploited by the virus itself to stabilize N⁰P, N and P were co-expressed as a fusion construct, separated by a TEV cleavage site, which allowed the purification of N⁰P, where P can be of any length as long as the N-terminal binding site is included in the construct. This tool provides access to conformational characterisation of N⁰P complexes comprising different levels of disorder (figure 1), direct investigation of NCLP assembly [22] and control of the RNA present in the complex.

Indeed, addition of short RNA molecules was shown to trigger formation of NCLPs (figure 1D). The process can be observed in real time using NMR – by observing the appearance of ¹⁵N-¹H cross-peaks from amino acids in the P-peptide that detach from N_{CORE} upon NCLP formation, and the disappearance of flexible peaks from N_{ARM} that become structured upon binding neighbouring N_{CORE} domains. The process can also be observed using negative staining electron microscopy (EM) and more importantly fluorescence anisotropy, with fluorescein-labelled RNA. This latter revealed the presence of two steps in the assembly kinetics, one significantly faster than the other. The physical origin of the biphasic assembly is not yet known, for example whether the two steps correspond to binding of RNA and then inter-protomer assembly, or vice versa. Importantly a clear dependence on RNA sequence was observed, with polyA and the 5' end of the genome assembling with high efficiency, while some sequences, for example polyU did not assemble. This specificity suggests a role for nucleation of assembly via specific sequences (5' sequences of genome and anti-genomes are highly conserved in *Paramyxoviridae*).

Cryo-EM structure of RNA-specific MeV nucleocapsids reveals new mechanistic insight

While interactions between N and RNA in MeV NCLPs have been investigated for using EM and cryo-EM [9,20,10,21] and in single turns crystallised as closed rings in a number of related viruses [23–26], in all cases the particles were purified in the assembled form after

overexpression in cellular systems. The particles therefore contained random cellular RNA at each position, so that only sequence-averaged structural information is obtained, precluding the determination of protein-base interactions. Nevertheless, the binding mode of the RNA to MeV N was shown to comprise three bases pointing into, and three bases pointing away from the RNA binding groove [21,26]. In order to gain further insight, NCLPs were polymerized on polyA and viral genomic 5' (ACCAGA) sequences using the approach described above, and their three-dimensional structures determined using cryo-EM to 3.4 Å resolution [27].

The 5' and 3' binding sites were unambiguously identified (figure 2), and thereby the register of binding of viral genomic RNA within the assembled NCLPs. This implies that the final three nucleotides at the 3' end of the genome are almost fully exposed in assembled nucleocapsids (figure 2), allowing access to the genome for the RNA-dependent RNA polymerase complex [28]. Putative protein-nucleobase interactions were identified in the RNA-binding groove (figure 2). In particular the sidechain of R195 was found to stabilize nucleic acid backbone and base via a tight bipartite interaction (figure 2). This interaction was then shown to be essential for assembly using mutagenesis, negative staining EM and real-time NMR [27].

N⁰P – A complex story of disorder and allostery

The role of P_{TAIL} domain was recently investigated by co-expressing integral N with P_{TAIL}(1-304), thereby stabilising the 90 kDa N⁰P_{TAIL} sub-complex, comprising the disordered regions P_{TAIL}, N_{ARM} and N_{TAIL} (in total over 450 disordered amino acids) as well as N_{CORE}. NMR spectroscopy allows the atomic resolution description of each disordered region in such highly dynamic complex (figures 1, 3) [19,29,30]. This study [12] revealed that, in addition to the known binding site at its N-terminus, P_{TAIL} contains an additional helical linear motif, ¹⁹¹HELL¹⁹⁴ (α 4), that binds weakly (with an intrinsic K_D~600 μ M) to the surface of the N-terminal lobe of N_{CORE}. This secondary binding site is the most evolutionarily variable part of N_{CORE} [31] and has been shown to be implicated in binding of cofactors of VSV [32] and RSV [33] and was exploited via chimeric constructs to be implicated in N_{TAIL}:XD interactions localised on the surface of N_{CORE} [34].

Remarkably, mutation of this site to ¹⁹¹AAAA¹⁹⁴ in cell-based MeV mini-genome studies revealed that the HELL site is essential for viral transcription and replication. Equivalent motifs comprising two bulky hydrophobic residues at the C-terminus of a short helical motif are conserved across morbilliviruses and paramyxoviruses comprising P_{TAIL} domains of very different length, such as Nipah (470 amino acids comprising ³⁴³RELL³⁴⁶). These sequences show significantly higher sequence conservation over non-redundant MeV sequences than the majority of P_{TAIL}, suggesting that this motif may provide an important new target for treating paramyxoviruses, in particular in view of the relatively weak intrinsic affinity that renders this target accessible to peptide-based inhibitors.

This study therefore revealed a more complex picture of the N⁰P complex than previously envisaged – that comprises not one but (at least) two N:P binding sites, both of which are essential for viral function – and sheds new light on the conserved presence of long intrinsically disordered paramyxoviral P_{TAIL} domains [35]. When both sites are bound, the unfolded P_{TAIL} wraps around N_{CORE}, but because the affinity of the two binding sites differs by around 5 orders of magnitude, the HELL motif transiently binds and unbinds while the N-terminal site is bound (figure 3). The impact of the highly dynamic 150 aa separating the two sites, comprising a cluster of acidic amino acids, may enhance chaperone function by frustrating interaction of RNA with the surface of N_{CORE}, impacting self-assembly with other N monomers or possibly regulating interactions with host factors involved in the immune response [36]. It should be mentioned that the construct investigated here comprises the genome-edited viral protein V [37–40] which is equivalent to P_{TAIL} until residue 232 followed by a zinc finger domain. The observations made here are thus possibly equally valid for putative N⁰V complexes. Finally the essential role of the HELL motif in MeV replication exemplifies the growing number of examples of ultra-weak interactions involving intrinsically disordered proteins in biology [41].

N and P proteins form membraneless organelles that promote nucleocapsid formation

As illustrated above, N and P proteins exhibit multiple essential interactions during the viral cycle, a multivalency that, in combination with their extensive intrinsic disorder and the

transient nature of some of the associated interactions, may play a role in the formation of viral factories in infected cells [42,43]. Liquid-liquid phase separation (LLPS), by which micro-environments remain immiscible with respect to their immediate surroundings thereby maintaining distinct reaction or storage conditions within the condensate, are now thought to be prevalent throughout cell biology [44–46]. Intrinsic disorder plays a key role in the thermodynamic stabilisation of LLPS, due to the prevalence of weak and transient interactions that contribute to maintaining the liquid nature of the environment, and allowing rapid spatial and temporal control of biochemical processes [47].

The formation of membraneless organelles, in addition to concentrating viral partners in a confined volume, may also provide protection from factors involved in the host-immune response. Numerous examples of colocalization of Mononegaviral N and P have been observed *in vivo*, including rabies [48], VSV [49], human parainfluenza types 5 [40] and 3 [50,51], RSV [17], Nipah [52], human metapneumovirus [53], MeV [18,54–56] and Ebola [57]. A detailed study by Gaudin et al demonstrated the liquid-like nature of Negri bodies formed in rabies [16] and colocalization of N and P in infection-induced cytoplasmic inclusion bodies in MeV [18] suggests that LLPS represents a widespread mechanism exploited by Mononegaviruses during infection. These studies also investigate the role of transport of membraneless organelles along the cytoskeleton in the infection cycle [16,18].

The formation of membraneless organelles comprising MeV N and P proteins was recreated *in vitro* using co-expressed soluble N⁰P complexes and different constructs of P under conditions where neither N⁰P nor P undergoes phase separation (figure 4). Expression of combinations of the different domains (figure 1) identified the tetramerization domain of P, the disordered domain P_{LOOP} and the three-helix bundle XD as essential for droplet formation (P_{TAIL} is dispensable for LLPS). Integral N, comprising the disordered N_{TAIL}, is essential. These results align with those measured in MeV [18] and rabies [16] *in vivo* in terms of the essential domains required for LLPS.

NMR spectroscopy was used to characterise the essential interactions, identifying the $N_{\text{TAIL}}:P_{\text{XD}}$ interaction [14,58–60] whose inhibition via a single mutation [61], results in abrogation of LLPS. A recent study also showed that XD deletion abolishes LLPS *in vivo* [18]. P_{LOOP} and N are also shown to interact via weak interactions (200–400 μM) that affect droplet fluidity, suggesting that the role of P_{LOOP} is not uniquely as a “spacer” [62] that ensures the physical separation of interacting components, but is also involved in regulating the dynamics of the droplets.

Investigation of the thermodynamics of droplet formation using NMR spectroscopy and the diffusion properties of the components using fluorescence imaging, revealed a probable mechanism of droplet stabilisation by which the tetrameric P protein interacts with N to form droplets, to which additional N proteins can be recruited, allowing for variable stoichiometry within the droplets [63]. P would then form a dynamic, multivalent scaffold comprising four branched dynamic XD interaction sites. Both N and P are highly dynamic within the droplets, but P diffuses slightly slower, in agreement with this model. As N is the most expressed MeV protein [64] and is required for assembly of nascent genomic material into nucleocapsids, such plasticity could be essential to allow for large variations of N concentration and fluid recruitment of N to droplets as nucleocapsids are produced.

Finally, and possibly most remarkably, membraneless organelles were shown to exhibit functional properties *in vitro*. Addition of RNA to a suspension of N:P droplets revealed preferential localisation to the droplets, and triggering of nucleocapsid assembly. The rate of assembly, as measured by real-time NMR, was significantly increased compared to the dilute phase, and EM revealed NCLP-like structures within one hour of addition of RNA. These results demonstrate that the condensed environment does not restrict the assembly of large superstructures and that droplet formation indeed enhances one of the essential processes of viral replication. The mechanisms observed here may be of general interest in characterising and inhibiting mononegaviral infection. The role of L, which is known to colocalize with N and P in mononegaviral puncta [49] remains to be characterised within viral coacervates. We note

that the presence of messenger RNA has been detected in puncta associated with rabies [48], RSV [17] and PIV3 [51], implying that both transcription and replication may be associated with LLPS in negative sense RNA viruses.

In conclusion we have described recent work centred on the biophysical characterisation of MeV N:P interactions involved in essential processes of mononegaviral infection. These aspects range from the observation of nucleocapsid assembly, to the identification of key N:RNA interactions and template architecture, to the development of a new understanding of the role of intrinsic disorder in paramyxoviruses and finally the formation of viral factories by this machinery, providing a functional microenvironment within which the preceding biophysical features should also be understood. In combination with the rapid increase in structural information concerning paramyxoviral and related RNA-dependent RNA polymerase structures and their interactions with phosphoproteins [65–69], our physical understanding of the molecular basis of viral replication is currently making major advances that will undoubtedly lead to the development of rational inhibitory strategies. We look forward to future structural studies of integral replication machinery comprising the N/NCLPs, P and the polymerase.

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Figure Legends

Figure 1. Architecture of MeV N and P proteins

- A. N and P both contain long, disordered strands (orange) that flank and are interspersed between structured domains (N_{CORE} , PMD, the multimerization domain of P and XD). The N-terminal peptide of P_{TAIL} binds N_{CORE} in the N^0P complex.
- B. Cryo-EM structure of nucleocapsid reveals that N_{ARM} stabilizes nucleocapsid assembly by binding neighbouring protomers (similarly the C-terminal flanking C_{ARM} 374-391). The N-terminal peptide of P_{TAIL} adopts a pose that inhibits these interactions thereby maintaining N in an unassembled state as determined in reference [15] and [31] in the case of Nipah virus.
- C. Co-expression of P_{TAIL} (either 1-50 or 1-304) with N_{CORE} or N_{FULL} chaperones N in a monomeric form (N^0P) and allows for studies of the pre-assembly complex. The nucleocapsid assembly process is triggered by addition of RNA .
- D. NMR and fluorescence spectroscopy and EM can be used to follow nucleocapsid assembly in real time as shown in references [22] and [27]. Following changes in peak intensity of NMR spectra of the complex it is possible to follow how the P peptide detaches from N_{CORE} and how N_{ARM} and C_{ARM} bind (left figure). The assembly process can be investigated by observing fluorescently labelled RNA, which, when bound to N exhibits higher anisotropy (central figure). Nucleocapsids can be imaged using electron microscopy (right figure).

Figure 2. High resolution structure of RNA-specific MeV nucleocapsid-like particles (as determined in the recent publication [27]).

- A. Helical reconstruction of nucleocapsid-like particles assembled in vitro on known RNA sequences determines the binding register of the RNA in the binding cleft, revealing that the last three 3' nucleotides in the genome are exposed. Three copies of N are shown, the central is shown in yellow cartoon format, with neighbouring copies shown

in blue. The RNA (orange) is shown in the binding cleft, with 5 phosphate groups associated with the hexameric RNA molecule shown as spheres.

- B. Detail of the protein-RNA interactions identified in the binding cleft due to the higher resolution structure and the unambiguous assignment of the electron density to the RNA sequence. The key interacting amino acids are shown in red. (reproduced from Desfosses et al *PNAS* 2019, **116**:4256–4264).
- C. R195 in particular is shown to adopt a bipartite interaction with the adenine in position 5. When this site is mutated no assembly is observed by NMR, fluorescence or EM. (reproduced from Desfosses et al *PNAS* 2019, **116**:4256–4264).

Figure 3. The role of intrinsic disorder in the MeV N⁰P complex (as determined from the recent NMR study of this 90 kDa assembly [12])

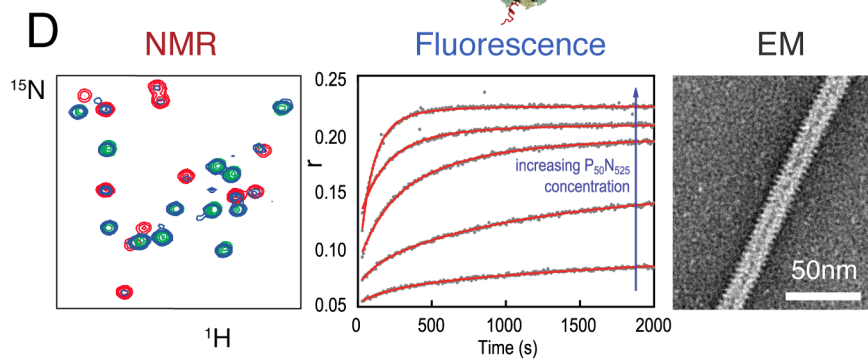
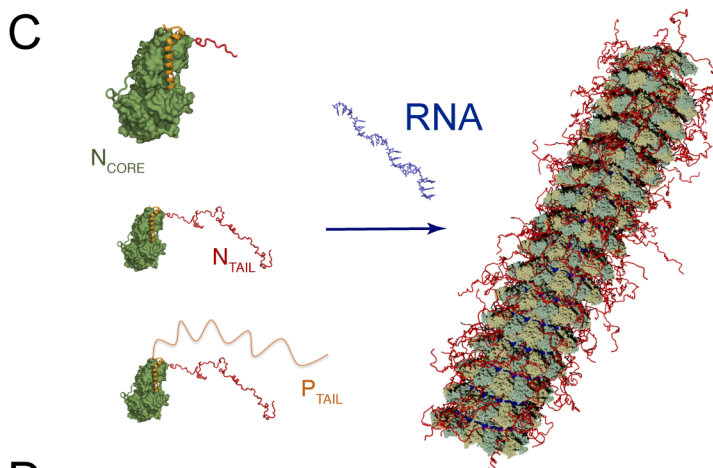
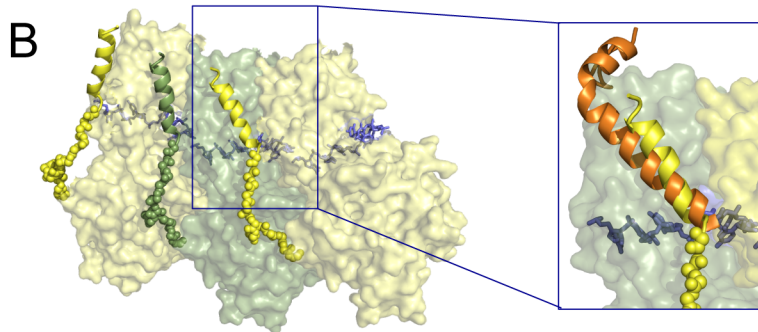
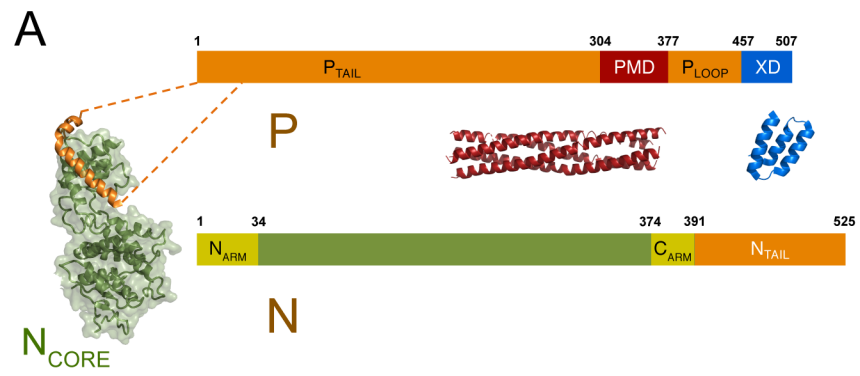
- A. Structural characterisation of MeV P_{TAIL} at amino acid specific resolution. Four helical elements are found along the sequence (α 1 and α 2 that correspond to the N-terminal peptide that binds N_{CORE}, α 3 and α 4). Reproduced with permission from Milles et al *Science Advances* 2018, **4**:eaat7778.
- B. Heteronuclear single quantum coherence (HSQC) spectrum of the N₁₋₅₂₅⁰P₁₋₃₀₄ complex comprising the entire P_{TAIL} and integral N. Reproduced with permission from Milles et al *Science Advances* 2018, **4**:eaat7778.
- C. HSQC peaks intensities provide insight into nature of the interactions between P_{TAIL} and N. Top - In addition to the N-terminal binding peptide, P_{TAIL} binds N via a second interaction site that corresponds to helix α 4 (comprising the sequence HELL). Bottom – mutation of HELL to AAAA abrogates the second interaction site, and renders the virus incompetent for replication. Reproduced with permission from Milles et al *Science Advances* 2018, **4**:eaat7778.
- D. The intrinsic affinity of the HELL interaction site for N is approximately 5 orders of magnitude weaker than the N-terminal site, such that the binding site exchanges

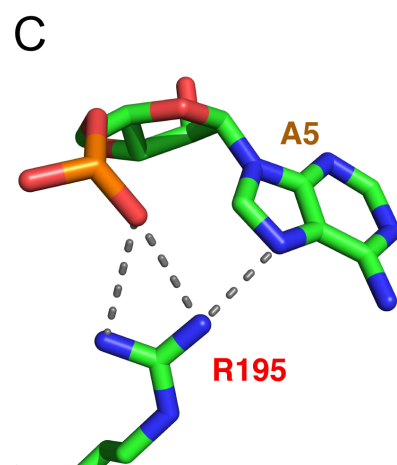
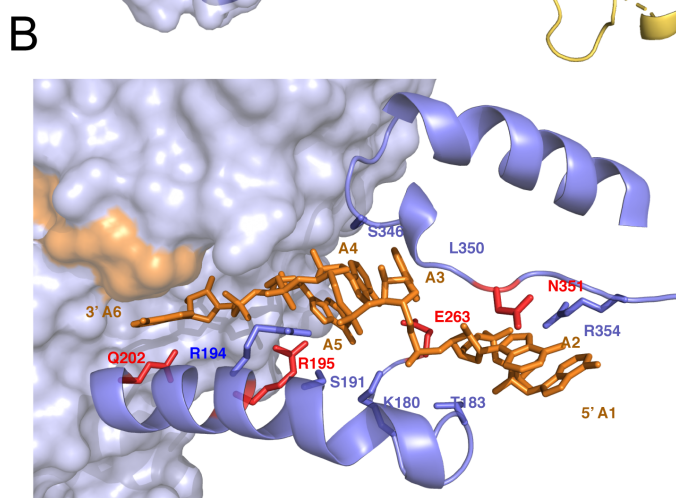
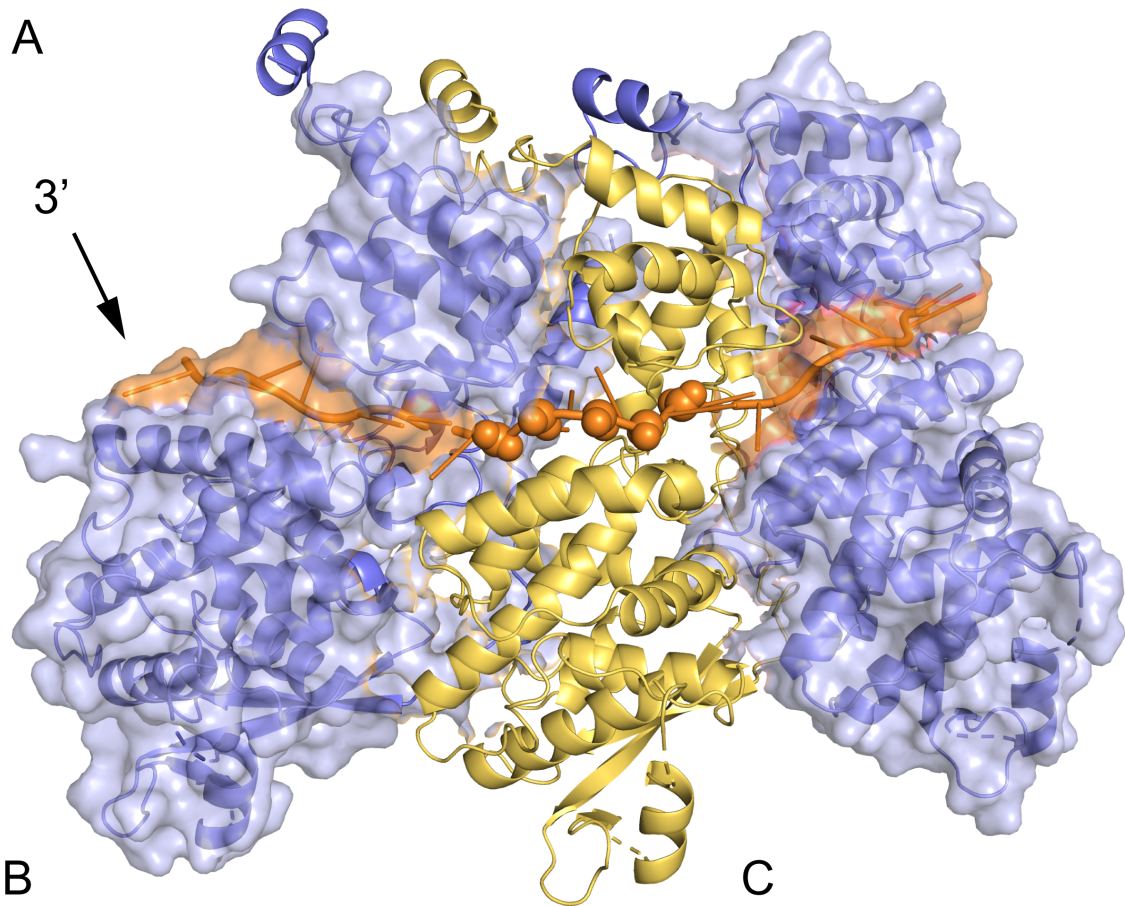
rapidly between bound and unbound conformations. Reproduced with permission from Milles et al *Science Advances* 2018, **4**:eaat7778.

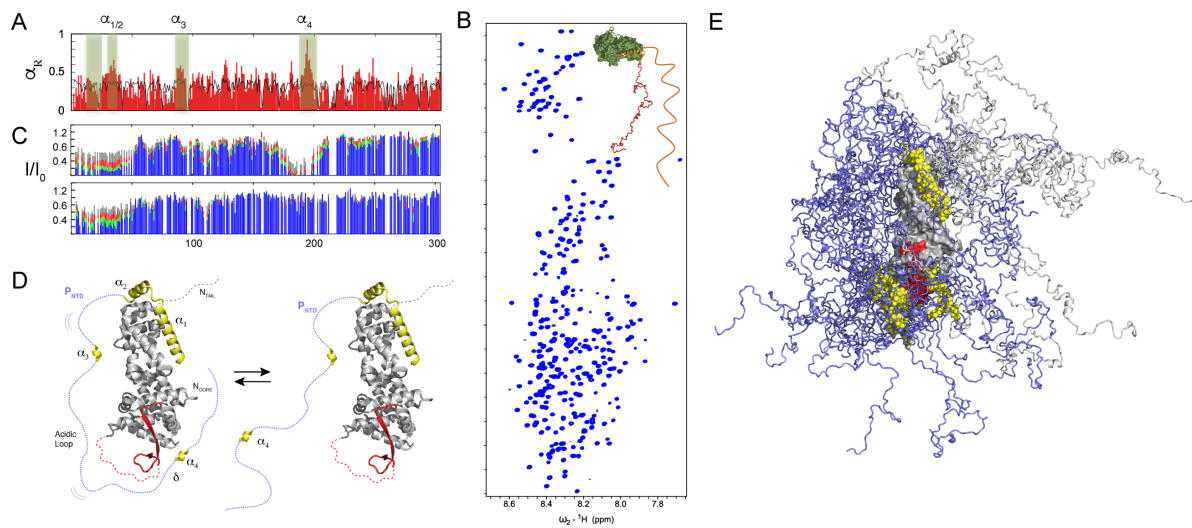
- E. An ensemble of conformations sampled by the P_{TAIL} - N_{CORE} complex. This highly dynamic process may contribute to protection against non-specific binding of host RNA or factors involving the host immune system. Reproduced with permission from Milles et al *Science Advances* 2018, **4**:eaat7778.

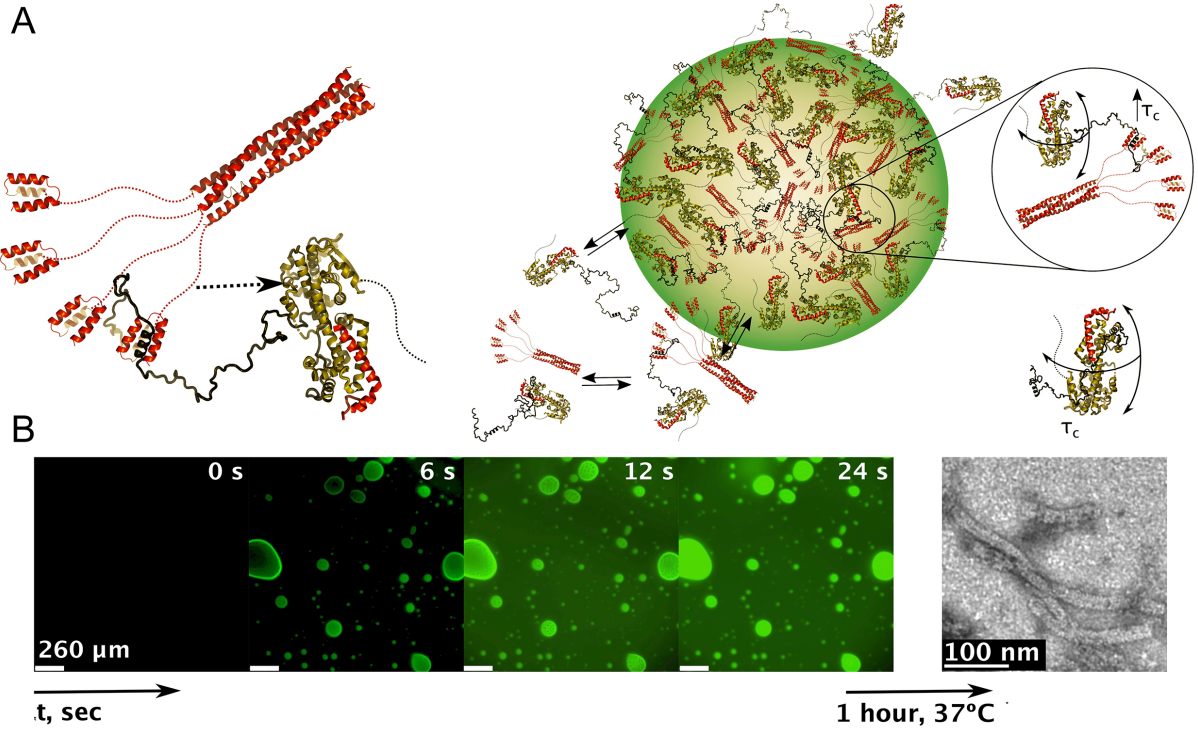
Figure 4. MeV N and P proteins form membraneless organelles that colocalize RNA and promote assembly of NCLPs.

- A. PMD, P_{LOOP} and XD and integral N constitute the minimal elements that can form droplets. Upon mixing with N, P appears to act as a dynamic scaffold, to which additional N proteins can be recruited.
- B. RNA colocalizes to N:P droplets (10 nucleotide polyA RNA is labelled with fluorescein in the example shown) and forms nucleocapsid like particles – a process that can be observed by NMR and EM. Reproduced with permission from Guseva et al *Science Advances* 2020, **6**:eaaz7095.









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