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The role of salt and shear on the storage and assembly of spider silk proteins

- [Lukas Eisoldt](#),
 - [John G. Hardy](#),
 - [Markus Heim](#),
 - [Thomas R. Scheibel](#)
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Abstract

Major ampullate silk fibers of orb web-weaving spiders have impressive mechanical properties due to the fact that the underlying proteins partially fold into helical/amorphous structures, yielding relatively elastic matrices that are toughened by anisotropic nanoparticulate inclusions (formed from stacks of β -sheets of the same proteins). *In vivo* the transition from soluble protein to solid fibers involves a combination of chemical and mechanical stimuli (such as ion exchange, extraction of water and shear forces). Here we elucidate the effects of such stimuli on the *in vitro* aggregation of engineered and recombinantly produced major ampullate silk-like proteins (focusing on structure–function relationships with respect to their primary structures), and discuss their relevance to the storage and assembly of spider silk proteins *in vivo*.

Keywords

- Spider silk;
 - Protein assembly;
 - Salt;
 - Shear
-

1. Introduction

Orb web-weaving spiders produce a number of task-specific silk protein-based fibers. One of these fibers is assembled from proteins produced by the major ampullate (MA) silk gland. MA silk fibers are used both in the construction of the web as the frame upon which to attach the capture spiral fibers (flagelliform silk fibers coated with aggregate silk) and as a lifeline to escape from predators ([Aprhisiart and Vollrath, 1994](#) and [Lin et al., 1995](#)). MA silk fibers of orb web-weaving spiders have impressive mechanical properties due to the fact that the underlying proteins partially fold into helical/amorphous structures yielding relatively elastic matrices that are toughened by anisotropic nanoparticulate inclusions (formed from stacks of β -sheets of the same proteins) ([Gosline et al., 1999](#)). The primary structures of MA proteins are reminiscent of amphiphilic multiblock copolymers ([Exler et al., 2007](#), [Geisler et al., 2008](#), [Zbilut et al., 2005](#) and [Zbilut et al., 2006](#)), in which the repetitive sequence elements (typically composed of multiple repeats of A_n , $(GA)_n$, $(GGX)_n$, and $(GPGXX)_n$) are flanked by highly conserved non-repetitive (NR) amino- and carboxy-terminal domains ([Bini et al., 2004](#), [Foo et al., 2006](#), [Hedhammar et al., 2008](#), [Lewis, 2006](#) and [Rising et al., 2006](#)).

In common with other silk proteins, the MA proteins of *Araneus diadematus* spiders (ADF-3 and ADF-4) are stored at very high concentrations (>30 wt.%) without the onset of undesirable aggregation within the lumen of the spider. However, our *in vitro* studies of the repetitive sequence elements of recombinantly produced proteins based upon the consensus sequences of the MA proteins of *Araneus diadematus* spiders ([Guerette et al., 1996](#)) without the NR terminal domains clearly demonstrate that these proteins are not soluble at such high concentrations as found *in vivo* ([Huemmerich et al., 2004a](#) and [Huemmerich et al., 2004b](#)). Our findings suggest that the NR terminal domains control both the storage of the proteins (by increasing their solubility and/or deterring undesirable aggregation events) and fiber production (allowing/facilitating controlled protein assembly into fibers upon demand) ([Hagn et al., in press](#)).

In vivo Raman spectromicroscopy ([Lefevre et al., 2008](#) and [Lefevre et al., 2007b](#)) and *in vitro*¹³C NMR ([Hijirida et al., 1996](#)) studies of the spinning dope indicate that the repetitive sequence elements adopt highly hydrated random coil and 3₁-helix (polyproline II-like) secondary structures which can be easily transformed into β -sheets. We ([Huemmerich et al., 2004b](#)) and subsequently others ([Huang et al., 2006](#), [Ittah et al., 2007](#), [Ittah et al., 2006](#), [Lin et al., 2009](#) and [Stark et al., 2007](#)) have shown the C-terminal NR domains to adopt a predominantly α -helical conformation *in vitro*. These domains have been postulated to form the outer layer of droplet-like structures observed in the spinning dope *in vivo* ([Jin and Kaplan, 2003](#), [Lin et al., 2009](#) and [Vollrath and Knight, 1999](#)). We have also shown that the C-terminal NR domain is a prerequisite of certain supramolecular self-assembly processes (manifested in the form of fully reversible lower critical solubilisation temperature behavior) of our recombinantly produced proteins ([Exler et al., 2007](#)).

In vivo the transition from soluble protein to solid fibers involves a combination of chemical and mechanical stimuli (such as ion exchange, acidification, extraction of water and shear forces ([Bini et al., 2004](#), [Foo et al., 2006](#), [Heim et al., 2009](#) and [Vollrath and Knight, 2001](#)) that have also been demonstrated to promote protein assembly into fibers *in vitro* ([Hardy and Scheibel, 2009a](#), [Hardy et al., 2008](#) and [Rammensee et al., 2008](#)). We were consequently interested in determining the effect of such stimuli upon the stability of our MA silk-like proteins ([Hardy and Scheibel, 2009b](#)).

A protein's solubility in aqueous solution is governed by many factors, of which, one of the most important is its primary structure, which also determines its secondary, tertiary and quaternary structures ([Levy et al., 2006](#), [Rossmann and Argos, 1981](#) and [SenGupta and Scheibel, 2007](#)). The presence of other solutes (such as ions) also has an effect upon a protein's solubility in water, as they interfere with the highly ordered layer of water (known as the hydration layer) on the protein's surface ([Gerstein and Chothia, 1996](#), [Kim and Cremer, 2001](#), [Lesk et al., 1980](#) and [Zhang and Cremer, 2006](#)). Low concentrations of salt tend to improve the solubility of proteins (known as 'salting in'), due to the formation of ion-rich hydration layers in the vicinity of charged and polar amino acid residues (as described by the Debye-Hückel theory), whereas high concentrations of salt tend to have the opposite effect, causing the protein to precipitate (known as 'salting out'). The magnitude of this effect is dependent upon the particular ions and usually follows the Hofmeister series, in which 'chaotropic' ions favor salting-in of proteins and 'kosmotropic' ions favor salting-out of proteins, and anions are well-known to have a much greater effect than cations ([Baldwin, 1996](#), [Geisler et al., 2008](#), [Gurau et al., 2004](#), [Horinek et al., 2008](#), [Pegram and Record, 2008](#), [Pirzer et al., 2009](#), [Zhang and Cremer, 2006](#) and [Zhang et al., 2005](#)).

Knight and Vollrath have investigated the MA gland of *Nephila edulis* spiders, finding that the physiological concentration of sodium chloride in the spinning dope in the lumen to be of the order of 150 mM, and the total concentration of salts was observed to decrease during the spinning process. They also found that as the dope flowed through the spinning duct, sodium cations were replaced by potassium cations, yet other metal cations (e.g. calcium, copper or magnesium) were almost undetectable in the lumen, duct and the naturally spun fibers. Furthermore, they observed that weakly chaotropic chloride anions are replaced with strongly kosmotropic phosphate anions (significantly increased concentrations) and sulfate anions (small increase with a relatively high level of experimental error) during the passage of the spinning dope along the spinning duct ([Knight and Vollrath, 2001](#)). Here we investigate the effects of salts (sodium chloride and sodium phosphate) and shear on the *in vitro* aggregation of our recombinantly produced MA silk-like proteins, and discuss their relevance to the storage and assembly of spider silk proteins *in vivo*.

In this study we use recombinantly produced silk proteins based on the consensus motifs of the MA silk *Araneus diadematus* fibroin 3 (ADF3) from the garden cross spider. The repetitive core domains of our proteins are composed of two different sequence modules, denoted A and Q, where module A is a hydrophobic polyalanine-rich motif, and module Q is a more hydrophilic glutamine- and glycine-rich motif (see materials and methods). In this study we use proteins containing 12 or 24 repeats of (AQ). The C-terminal NR3 domain is derived from *adf3* (gi|1263286, obtained from J. Gosline, Vancouver, BC) by using PCR. This domain is mainly α -helical and dimerises via disulfide bond formation between the single cysteine residues contained in two individual NR3 domains ([Huemmerich et al., 2004b](#)) ([Fig. 1](#)); therefore proteins bearing the NR3 domain are dimeric under non-reducing conditions.



Fig. 1.

Employed recombinantly produced proteins based on the consensus sequences of the major ampullate protein (ADF-3) of *Araneus diadematus* spiders. (A) A schematic representation of the repetitive sequence elements (A, hydrophobic polyalanine-rich motif and Q, hydrophilic

glutamine- and glycine-rich motif) and the C-terminal non-repetitive domain (NR3). (B) A schematic representation of the engineered and recombinantly produced major ampullate silk-like proteins composed of the elements shown above. For each protein the molecular weight in kDa is given in brackets. Note that proteins bearing the NR domain are dimeric due to interdomain disulfide bond formation. (C) The primary sequence of the C-terminal non-repetitive domain (NR3) of the major ampullate protein (ADF-3) of *Araneus diadematus* spiders, highlighting the residues involved in salt bridges and the cysteine residue responsible for interprotein dimerization. In the (AQ)₁₂NR3 (D93A) mutant the single aspartic acid residue is replaced by an alanine residue, inhibiting the formation of one of the two possible salt bridges.

[Figure options](#)

2. Materials and methods

2.1. Cloning, protein expression and purification

The proteins (AQ)₁₂NR3 (dimer of 116 kDa), (AQ)₂₄NR3 (dimer of 212 kDa) and (AQ)₂₄ (monomer of 95 kDa), containing the repetitive elements (A, hydrophobic polyalanine-rich motif: GPYGPASAAAAAAGGYGPGSGQQ; Q, hydrophilic glutamine- and glycine-rich motif: GPGQQGPGQQGPGQQGPGQQ) were cloned, expressed and purified as previously reported. A D93A mutant (numbering according to the residues of the NR domain; see [Fig. 1C](#)) of (AQ)₁₂NR3 was obtained according to the QuikChange protocol from Stratagene (USA).

2.2. Salt-induced aggregation assay

Experiments were carried out in accordance with our previously described protocol ([Huemmerich et al., 2004b](#)). Prior to the experiment the proteins were dissolved in guanidinium thiocyanate solution (6 M) and then dialyzed against 10 mM Tris/HCl, pH 8.0, 50 mM NaCl. The assay was started by the addition of buffered (10 mM Tris/HCl, pH 8.0) solutions of NaCl and NaH₂PO₄ (pH 8.0). Protein concentrations were, respectively, 8.8 μM, 5.3 μM and 4.7 μM for AQ₁₂NR3 (wt and D93A), AQ₂₄ and AQ₂₄NR3. After 1 h of incubation at 25 °C the samples were centrifuged at 17700 g to remove any visible aggregates, and the amount of soluble protein remaining in the supernatant was determined using a UV spectrometer (NanoDrop ND1000, ThermoFischer, USA).

2.3. Shear-induced aggregation assay

Prior to the experiment the proteins were dissolved in guanidinium thiocyanate solution (6 M) and then dialyzed against 10 mM Tris/HCl, pH 8.0. Then the proteins were diluted into buffer containing 10 mM Tris/HCl, pH 8.0 with 50 mM NaCl. Final protein concentrations were adjusted to 17.6 μM of AQ₁₂NR3 and 10.6 μM of AQ₂₄. The samples were incubated for 16 h at 25 °C without/with rotation at 25 rpm (Intellimixer RM-2, NeoLab, Germany). After rotation, the clearly visible aggregates were transferred onto glass slides for optical microscopy (DMI3000B, Leica, Switzerland). The samples were then centrifuged at 125000 g to remove non-visible aggregates, and the amount of remaining soluble protein in the supernatant was determined using a UV spectrometer (NanoDrop ND1000,

ThermoFischer, USA). The percentage of aggregated protein was calculated as described in section 2.2.

2.4. Fourier transform infrared (FTIR) spectroscopic studies

FTIR spectra of fibers were recorded on a liquid nitrogen cooled Bruker Tensor 27 FTIR spectrometer, fitted with a liquid nitrogen cooled Bruker Hyperion microscope with IR polarizers supplied by Optometrics Corporation (Ayer, MA, USA). Spectroscopy was performed on fibers formed during the previously described aggregation assay that were washed with water to remove salts, air dried and placed on highly polished optical grade calcium fluoride disks (Crystal GmbH, Berlin, Germany). Spectra of all aggregates were recorded in absorbance mode at 21 °C, with a 4 cm⁻¹ resolution and 60 scans (corrected for background and atmosphere using OPUS software), and polarized FTIR spectra of aggregates of μm scale (and larger) were recorded at three different orientations (0°, 45° and 90° relative to the long axis of the fibrous aggregates) to determine if any of the secondary structural elements were aligned with the fibers.

2.5. Transmission electron microscopy

Prior to the experiment the proteins were dissolved in guanidinium thiocyanate solution (6 M) and then dialyzed against 10 mM Tris/HCl, pH 8.0. A droplet (3 μL) was placed on pioloform-coated copper grids and the excess liquid was removed by wicking using a Kimwipe (Kimberly-Clark Corporation, USA). The grid was then stained with uranyl acetate (1% in water, pH 4.5) and the excess liquid was removed by wicking using a Kimwipe, and allowed to dry in a dust free dessicator. The grids were viewed with a JEOL JEM-2100 transmission electron microscope (Jeol, Eching, Germany).

3. Results and discussion

3.1. The effect of salt on protein aggregation

As noted in the introduction, MA spidroins are stored in the lumen of the gland in the presence of sodium and chloride ions, and during fiber spinning these ions are exchanged for potassium and phosphate ions ([Knight and Vollrath, 2001](#)). Here we investigate the effects of chloride and phosphate anions on the *in vitro* aggregation of our recombinantly produced MA silk-like proteins ([Fig. 1](#)).

Our solubility/aggregation assays in the absence of shear forces (similar to the situation of the silk proteins in the lumen of the gland) show that the extent of protein aggregation is determined by the concentration of salt. Our assays with sodium chloride (found in the lumen) indicate that the level of aggregation is low (less than 20%) for all of the tested engineered proteins, even at extremely high salt concentrations (up to 500 mM); whereas in the presence of equivalent concentrations of sodium phosphate (found in the spinning duct) the level of aggregation for each protein is notably higher (up to 100%) due to the more kosmotropic nature of the phosphate anion ([Fig. 2](#)). Therefore, weakly chaotropic chloride anions aid the long term storage of MA spidroins in the lumen (salting-in) and are substituted for strongly kosmotropic phosphate anions (salting out) during the natural fiber spinning process.

Fig. 2.

Salt-induced aggregation of engineered and recombinantly produced major ampullate silk-like proteins after incubation for 1 h at RT: induced by the presence of sodium chloride (grey circles) and by the presence of sodium phosphate (black circles), the concentration regime required to aggregate the proteins is indicated by striped boxes. (A) (AQ)₁₂NR3 (wild type). (B) (AQ)₁₂NR3 (D93A). (C) (AQ)₂₄NR3 (wild type). (D) (AQ)₂₄. (E) A comparison of the degree of aggregation of our spider silk proteins with and without shear; induced in the presence of 50 mM sodium chloride (grey bars), or 500 mM sodium chloride (black bars).

[Figure options](#)

3.2. The effect of shear on protein aggregation

During the passage of the spinning dope from the lumen through the spinning duct to the spigot (the point at which the fiber exits the animal), the dope is exposed to shear forces which are known to encourage aggregation in a variety of globular and fibrillar proteins ([Bekard and Dunstan, 2009](#), [Cromwell et al., 2006](#), [Hamilton-Brown et al., 2008](#), [Hill et al., 2006](#), [Vezy et al., 2009](#), [Yamaura et al., 1982](#) and [Yamaura et al., 1985](#)); we were therefore keen to find out if shear forces have an effect upon the aggregation of our recombinantly produced MA silk-like proteins *in vitro*.

Our aggregation assays in the presence of shear forces (as the silk proteins are exposed to in the duct during fiber spinning) show that the extent of protein aggregation is dramatically increased in relation to corresponding assays in the absence of shear, in the presence of both low and high concentrations of salt; moreover, the aggregates formed after longer periods of exposure to shear were fibrous with dimensions of μm to mm scale ([Fig. 3](#)).

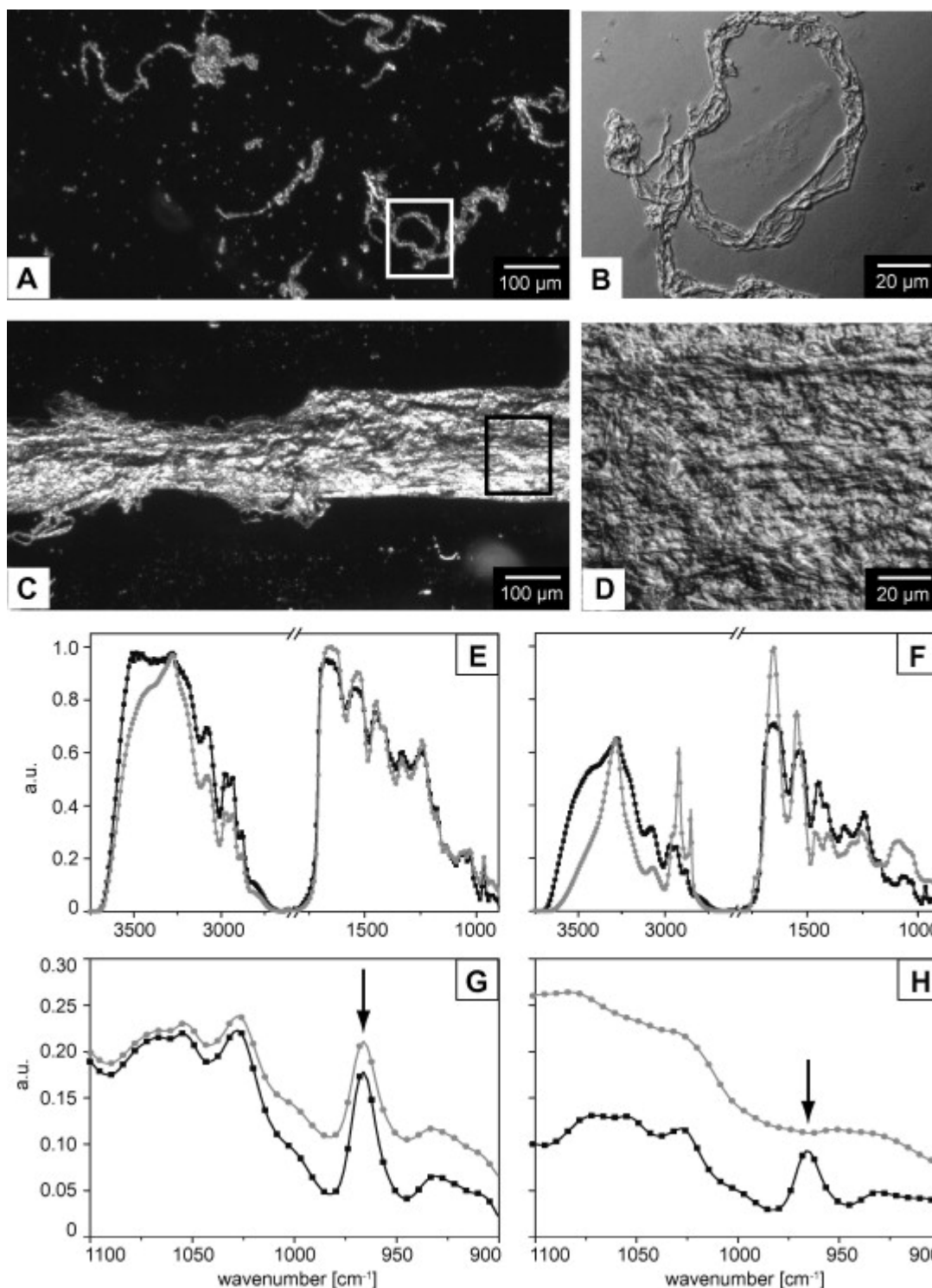


Fig. 3.

Shear-induced aggregation of our recombinantly produced major ampullate silk-like proteins. (A) Optical microscope image of aggregates produced by shear-induced aggregation of aqueous solutions of (AQ)₂₄. (B) Magnified image of the (AQ)₂₄ aggregates shown in A) (white frame). (C) Optical microscope image of well-defined fibrous aggregates produced by shear-induced aggregation of aqueous solutions of (AQ)₁₂NR3. (D) Magnified image of one part of the (AQ)₁₂NR3 fiber (black frame). (E) Polarized FTIR spectra from (AQ)₂₄ aggregates from A/B recorded at 0° (black circles) and 90° (grey squares) relative to the long

fiber axis. (F) Polarized FTIR spectra from AQ₁₂NR3 fibrils from C/D recorded at 0° (black circles) and 90° (grey squares) relative to the long fiber axis. (G) Magnification of the region between 1100 and 900 cm⁻¹ from the AQ₂₄ spectra (E). The arrow indicates the characteristic polyalanine β-sheet band at around 960 cm⁻¹ (Papadopoulos et al., 2009). (H) Magnification of the region between 1100 and 900 cm⁻¹ from the (AQ)₁₂NR3 spectra (F). The arrow indicates the characteristic polyalanine β-sheet band at around 960 cm⁻¹.

[Figure options](#)

Particularly noteworthy is that water is absorbed from the dope during its passage along the spinning duct (Kojic et al., 2004 and Tillinghast et al., 1984), which increases the concentration of proteins and consequently increases the shear forces between the proteins in the dope due to strengthened hydrophobic interactions. This leads us to suggest that shear forces are likely to play a greater role than salt effects during the fiber spinning process *in vivo* (Casem et al., 2002, Knight and Vollrath, 2002, Perez-Rigueiro et al., 2001a and Perez-Rigueiro et al., 2001b).

3.3. The role of the C-terminal NR domain in fiber storage and assembly

It is widely accepted that higher order supramolecular assemblies are required to achieve silk protein storage at high concentrations within the lumen, and as noted in the introduction, we have previously shown the highly conserved C-terminal NR domain to be a prerequisite of certain supramolecular self-assembly processes (manifested in the form of fully reversible lower critical solubilisation temperature behavior) of our recombinantly produced proteins (Exler et al., 2007). Interestingly, our transmission electron microscope studies of freshly dialyzed solutions of (AQ)₁₂NR3 show evidence of vesicle-like supramolecular assemblies (of nm to μm scale), which is not the case for (AQ)₂₄, a protein of similar molecular weight without the C-terminal NR domain (Fig. 4), which has also been observed by light microscopy in a previous study (Rammensee et al., 2008).

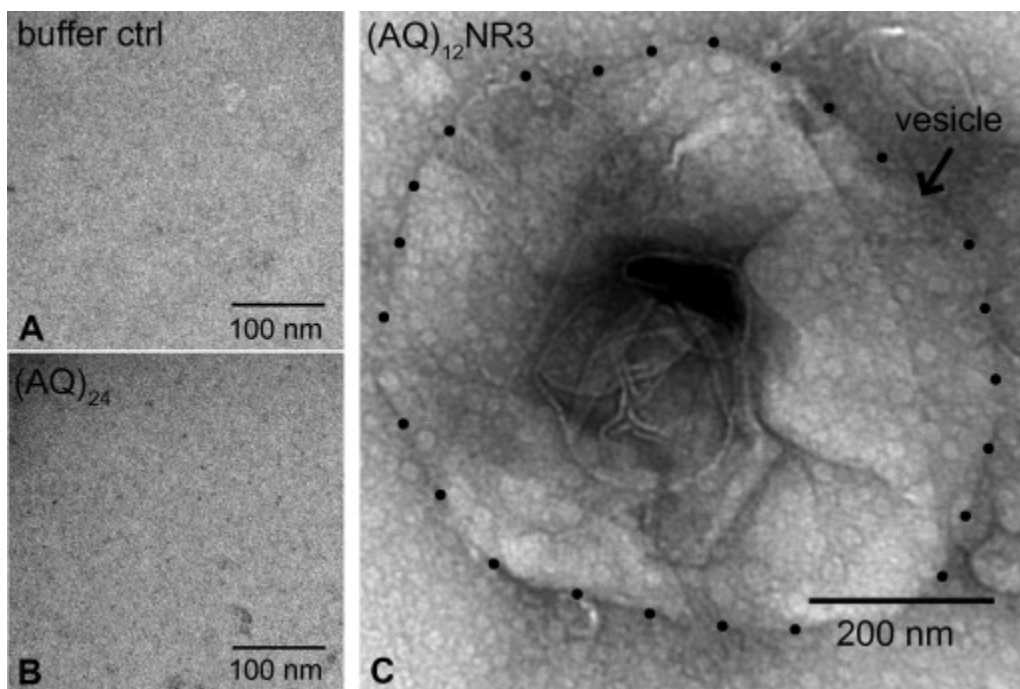


Fig. 4.

Images obtained via transmission electron microscopy. (A) Dialysis buffer. (B) Freshly dialyzed solutions of (AQ)₂₄. (C) Freshly dialyzed solutions of (AQ)₁₂NR3, showing evidence of higher order supramolecular assemblies.

[Figure options](#)

Our salt-induced aggregation studies revealed no significant differences in the behavior of (AQ)₁₂NR3 and (AQ)₂₄ in the presence of sodium chloride, whereas in the presence of sodium phosphate (>200 mM) (AQ)₂₄ is much more prone to aggregation ([Fig. 2A–D](#)) suggesting that the C-terminal NR domain dependent supramolecular assemblies are stabilized to a certain degree against aggregation. Shear stress had an even pronounced effect on protein aggregation ([Fig. 2E](#)). Furthermore, exposure of solutions of (AQ)₂₄ to shear stress resulted in the formation of ill-defined β -sheet rich aggregates ([Fig. 3A](#) and [B](#) and [Supplementary Table 1](#)), whereas exposure of solutions of (AQ)₁₂NR3 to shear stress yielded relatively well-defined fibrous aggregates which optical microscopy revealed to be composed of ordered (aligned) bundles of long fibrous aggregates ([Fig. 3C](#) and [D](#)). Interestingly, polarized Fourier Transform Infrared Spectra recorded at two different orientations (0° and 90° relative to the long axis of the fibrous aggregates) showed that the β -sheets within the fibrous aggregates of (AQ)₂₄ are randomly aligned, whereas the β -sheets within the fibrous aggregates of (AQ)₁₂NR3 were predominantly aligned with the long axis of the fibers (in analogy to naturally spun fibers ([Boulet-Audet et al., 2008](#), [Lefevre et al., 2007a](#), [Rousseau et al., 2009](#) and [Rousseau et al., 2004](#))) due to a controlled assembly process ([Fig. 3E](#) and [F](#)) ([Hagn et al., in press](#)). This finding was further confirmed by FTIR in the region between 1100 and 900 cm⁻¹ where specific signals for β -polyalanine crystals (964 cm⁻¹) can be observed, as reported recently ([Papadopoulos et al., 2009](#)). Interestingly, a clear difference at the two employed angles using the polarizer can be seen for (AQ)₁₂NR3 aggregates ([Fig. 3H](#)), while there is no spectral difference for the (AQ)₂₄ aggregates, pointing out that the polyalanine crystals are oriented in the fibers formed from the protein with the NR domain, but not in the fibers formed from the protein without the NR domain. These findings suggest that the C-terminal NR domain controls both the storage of the proteins (by increasing their solubility and/or deterring undesirable aggregation events) and fiber formation (by facilitating controlled protein assembly into fibers upon demand).

3.4. The role of the N-terminal NR domain in fiber storage and assembly

The N-terminal NR domains of silk proteins have been shown to be highly conserved between different spider species ([Hedhammar et al., 2008](#), [Lin et al., 2009](#), [Motriuk-Smith et al., 2005](#) and [Rising et al., 2006](#)). However, due to a lack of any published sequence data of the N-terminal NR domains of MA silk proteins of *Araneus diadematus*, this domain was not included in our engineered protein constructs. Despite this fact, our *in vitro* salt-induced aggregation assays with proteins with higher molecular weights and with longer repetitive sequence elements ((AQ)₂₄NR3 as opposed to (AQ)₁₂NR3) showed them to be more prone to aggregation ([Fig. 2](#)), which leads us to speculate that the N-terminal NR domain might also have an effect on higher order supramolecular assembly of silk proteins and therefore play an important role in protein storage and fiber assembly

in vivo, as the natural proteins are larger than our recombinantly produced analogues ([Guerette et al., 1996](#) and [Huemmerich et al., 2004b](#)).

4. Conclusions

Our salt-induced aggregation assays clearly demonstrate that chloride (which is present in the lumen during silk protein storage) does not induce significant levels of protein aggregation, whereas phosphate (present in the spinning duct) does induce significant levels of protein aggregation, due to the strongly kosmotropic nature of the phosphate anion. Our shear-induced aggregation assays clearly show that shear forces are likely to play a greater role than salt effects during the fiber spinning process *in vivo*, and moreover, a key role in the alignment of the secondary structural elements within the fibers (which controls their mechanical properties). Our transmission electron microscope studies show that the C-terminal NR domain is a prerequisite for higher order supramolecular assembly of the proteins, which is important for both protein storage and the controlled assembly of fibers. Finally, we speculate that the N-terminal NR domains also play a role in protein storage (and potentially fiber formation).

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