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REPORT

Unraveling amyloid formation paths of Parkinson's disease protein α -synuclein triggered by anionic vesicles

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Abstract. Amyloid formation of the synaptic brain protein α -synuclein (α S) is related to degeneration of dopaminergic neurons in Parkinson's disease patients. α S is thought to function in vesicle transport and fusion and it binds strongly to negatively charged vesicles *in vitro*. Here we combined circular dichroism, fluorescence and imaging methods *in vitro* to characterize the interaction of α S with negatively charged vesicles of DOPS (1,2-dioleoyl-*sn*-glycero-3-phospho-L-serine, sodium salt) and DOPG (1,2-dioleoyl-*sn*-glycero-3-phospho-(1'-*rac*-glycerol), sodium salt) and the consequences of such interactions on α S amyloid formation. We found that lipid head-group chemistry modulates α S interactions and also affects amyloid fiber formation. During the course of the experiments, we made the unexpected discovery that pre-formed α S oligomers, typically present in a small amount in the α S starting material, acted as templates for linear growth of anomalous amyloid fibers in the presence of vesicles. At the same time, the remaining α S monomers were restricted from vesicle-mediated nucleation of amyloid fibers. Although not a dominant process in bulk experiments, this hidden α S aggregation pathway may be of importance *in vivo*.

1. Introduction

The protein alpha-synuclein (α S) is a major component of amyloid aggregates found in Lewy body inclusions, which are the pathological hallmark of Parkinson's disease (PD). PD is the second most common neurological disorder and the most common movement disorder. The assembly process of the 140-residue protein α S into amyloid fibrils, via oligomeric intermediates, has been linked to the molecular basis of PD (Uversky, 2007). Duplications, triplications and point-mutations in the α S gene are related to familial PD cases (Polymeropoulos *et al.* 1997; Spillantini *et al.* 1997).

The exact function of α S is unknown, but it is suggested that the protein is involved in synaptic vesicle release and trafficking, regulation of enzymes and transporters, and control

of the neuronal apoptotic response (Dev *et al.* 2003; Lassen *et al.* 2016). α S is present at presynaptic nerve terminals (Eliezer *et al.* 2001; Iwai *et al.* 1995; Maroteaux *et al.* 1988) and it is reported that the protein exists *in vivo* in both free cytosolic and membrane-bound states (Lee *et al.* 2002). Membrane-bound α S can generate nuclei/oligomers, which can seed the aggregation of cytosolic α S (Lee *et al.* 2002). The monomeric α S form in solution has a disordered structure, whereas the membrane-bound state is rich in alpha-helix structure (Bodner *et al.* 2009; Davidson *et al.* 1998; Eliezer *et al.* 2001; Fusco *et al.* 2014). PD pathology may be associated with change or disruption of α S–lipid interactions, and *in vitro* studies have indicated that early-onset PD mutations can affect the binding of α S to phospholipids (Choi *et al.* 2004; Jo *et al.* 2002; Perrin *et al.* 2000). It was suggested that the mechanism of toxicity in PD involves direct, disruptive interactions between α S and lipid bilayers involving an interplay between growth of α S aggregates and lipid extraction (Reynolds *et al.*

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2011). α S has also been shown to bind specifically to mitochondria, which can lead to disruption of their function (Nakamura *et al.* 2008).

Because of the potential relevance to both functional and pathological roles of α S, further characterization of interactions between α S and lipid membranes is of great importance. There exists a significant body of work aimed at understanding the nature of interactions between α S and lipid bilayers but it has proven difficult to characterize the membrane-bound state and the role of membranes on α S aggregation. This challenge is directly related to the difficulty in isolating the membrane-bound protein from brain samples. *In vitro* studies have also not yet provided a clear mechanism for α S aggregation in the presence of lipids in part due to variability of protein-starting sample and variety of conditions used in kinetic studies of the conversion of monomeric α S into amyloid fibrils (Cole *et al.* 2002; Comellas *et al.* 2012; Fink, 2006; Galvagnion *et al.* 2015, 2016; Grey *et al.* 2011; Jo *et al.* 2000; Lee *et al.* 2002; Martinez *et al.* 2007; Zhu & Fink, 2003). Here we present a biophysical analysis of α S–lipid interactions that unravels a role for the lipid head-group chemistry in both α S binding and amyloid formation. Our experiments led us to the unexpected discovery of a hidden pathway involving oligomeric α S species that, when bound to membranes, acted as templates for linear growth of thin α S amyloid fibrils.

2. Materials and methods

2.1. Materials

DOPS (1,2-dioleoyl-*sn*-glycero-3-phospho-L-serine, sodium salt) and DOPG (1,2-dioleoyl-*sn*-glycero-3-phospho-(1'-*rac*-glycerol), sodium salt) chloroform solutions were purchased from Avanti Polar Lipids (Alabaster, USA). Sodium phosphate monobasic (NaH_2PO_4 ; $\geq 99.0\%$), sodium phosphate dibasic (Na_2HPO_4 ; $\geq 99.0\%$) and thioflavin T UltraPure Grade (ThT) ($\geq 95\%$) were purchased from Sigma-Aldrich (Stockholm, Sweden).

2.2. Protein and lipid vesicle preparation

α S was expressed and purified as described previously (Chorell *et al.* 2015). The protein was stored in a lyophilized form at -80°C . Monomeric α S was purified from the lyophilized powder dissolved in 8 M urea and 20 mM ethylene diamine tetra-acetic acid by size exclusion chromatography (SEC) on a Superdex 75 column (GE Healthcare, Uppsala, Sweden) into 20 mM phosphate buffer, pH 6.5. Aggregation kinetic studies were also performed with samples of lyophilized protein dissolved directly in 20 mM phosphate buffer, pH 6.5, followed by filtration through 0.2 μm syringe filters. These samples (found to contain both monomers and oligomers)

consisted of only full-length α S (no fragments) as assessed by SDS–PAGE. The protein concentration was determined spectrophotometrically using a molar extinction coefficient of $5960 \text{ M}^{-1} \text{ cm}^{-1}$ at 280 nm. Lipid vesicles were prepared by the lipid film hydration method. Appropriate volumes of chloroform solution of the lipid (DOPS or DOPG) were transferred to round bottom flask and the organic solvent was removed by rotary evaporation. The resultant film was further dried under vacuum for at least 3 h and then hydrated with 20 mM phosphate buffer, pH 6.5. The liposome suspensions were extruded through polycarbonate filters with pore diameters of 100 nm using Avestin LiposoFast-Basic extruder. The size of the liposomes was measured by dynamic light scattering using a Zetasizer Nano ZS instrument (Malvern Instruments Ltd., Worcestershire, UK).

2.3. Circular dichroism (CD) spectroscopy

Far-UV CD spectra of 5 μM α S in the presence of increasing concentrations of liposomes in 20 mM phosphate buffer, pH 6.5, were recorded at 37°C on a Chirascan (Applied Photophysics, Leatherhead, UK) equipped with a Peltier temperature control unit. CD spectra between 250 and 200 nm were obtained using quartz cuvettes with path lengths of 1 mm and the following parameters: a bandwidth of 1 nm, a step size of 1 nm, a time per point of 1 s and averaging three individual spectra. The CD signal of the buffer and liposomes was recorded and subtracted from the CD signal of the protein.

2.4. CD data analysis

CD titration curves were fitted using one-step binding model (Galvagnion *et al.* 2015): $F + \text{Lipid}_L \xrightleftharpoons{K_D} B(\text{Lipid})_L$, where F and B represent α S-free in solution and bound to the vesicles, respectively, and L is the number of lipid molecules interacting with one molecule of α S. Then:

$$K_D = \frac{[F][\text{Lipid}_L]}{[B(\text{Lipid})_L]} \quad (1)$$

$$[\alpha\text{S}] = [F] + [B] \quad (2)$$

$$[\text{Lipid}] = L([\text{Lipid}_L] + [B(\text{Lipid})_L]) \quad (3)$$

Detected CD signal can be expressed as a sum of contributions from α S bound to the vesicles ($x_B = [B]/[\alpha\text{S}]$) and α S free in solution ($x_F = [F]/[\alpha\text{S}]$):

$$x_B + x_F = 1 \quad (4)$$

$$\text{CD} = x_B \text{CD}_B + x_F \text{CD}_F \quad (5)$$

where CD_B and CD_F are the CD signals of the α S bound and free states, respectively.

From Eqs. (4) and (5):

$$x_B = \frac{\text{CD} - \text{CD}_F}{\text{CD}_B - \text{CD}_F} \quad (6)$$

And from Eqs. (1)–(4):



$$x_B = \frac{\left([\alpha S] + \frac{[\text{Lipid}]}{L} + K_D \right) - \sqrt{\left([\alpha S] + \frac{[\text{Lipid}]}{L} + K_D \right)^2 - \frac{4[\alpha S][\text{Lipid}]}{L}}}{2[\alpha S]} \quad (7)$$

Matlab software package was used to fit CD titration curves using Eqs. (6) and (7).

2.5. Aggregation kinetics

αS purified by SEC just before incubation or αS dissolved directly in buffer was incubated at 37 °C in 20 mM phosphate buffer, pH 6.5, in the presence of 20 μM thioflavin T (ThT) and different concentrations of DOPG and DOPS liposomes. The samples (60 μl per well) were incubated in non-binding polystyrene 96-well half-area plates with clear bottom (Corning Life Sciences, Tewksbury, USA) ThT fluorescence signal was monitored using a microplate reader (FLUOstar Optima, BMG Labtech, Ortenberg, Germany). The fluorescence (excitation wavelength 440 nm, emission wavelength 480 nm) was measured at every 20 min during the incubation and the plates were agitated for 5 min before each measurement to assure proper mixing of the samples. In the absence of liposomes, aggregation of αS does not occur within the timeframe of our experiments (100 h).

2.6. Atomic force microscopy (AFM)

For AFM imaging, samples were diluted with Milli-Q water (10–20 times) and deposited on freshly cleaved mica. After 10 min, the mica was rinsed with Milli-Q water and dried under a gentle nitrogen stream. Images were recorded in intermittent contact mode in air using an NTEGRA Prima setup (NT-MDT Spectrum Instruments, Moscow, Russia), a gold-coated single-crystal silicon cantilever (NT-MDT, NSG01, spring constant of $\sim 5.1 \text{ N m}^{-1}$) and a resonance frequency of $\sim 180 \text{ kHz}$. 512 pixel images were acquired with 0.5 Hz scan rate. Images were analyzed using the WSxM 5.0 software (Horcas *et al.* 2007).

3. Results

3.1. Binding of αS to anionic lipid vesicles is modulated by lipid head-group chemistry

We investigated the binding of monomeric αS to lipid vesicles composed of DOPS or DOPG lipids by far-UV CD spectroscopy. Titration of large unilamellar liposomes (LUVs) to αS induced a change from unordered to alpha-helix structure in the protein, as indicated by characteristic negative bands at 208 and 222 nm in the CD spectrum (Fig. 1a). The CD signal at 222 nm of αS scales almost linearly with increasing lipid concentration until saturation is reached (Fig. 1b) and the data can be well described by a one-step binding model. Fitting the binding curves with this model, we estimated the αS –lipid binding stoichiometry

L (how many lipid molecules per one αS molecule) and the apparent lipid–protein dissociation constant K_d . For both types of lipid vesicles, the αS dissociation constant was $\sim 0.1 \mu M$ ($0.10 \pm 0.08 \mu M$ for DOPG and $0.14 \pm 0.12 \mu M$ for DOPS), while the binding stoichiometry L was approximately four times higher for DOPS compared with DOPG vesicles (83 *versus* 21, respectively, lipids/ αS). This result demonstrates that the binding of αS to negatively charged lipid vesicles is not solely based on electrostatic forces but must in part be defined also by interactions between the hydrophobic face of the amphipathic helix and the membrane interior, which are affected by properties such as surface chemistry and/or local structure of the lipid bilayer (Perrin *et al.* 2000).

3.2. Amyloid formation of αS is differentially triggered by the two kinds of anionic lipid vesicles

Our CD results showed different stoichiometry L of αS for DOPG and DOPS liposomes. To assess the importance of this for αS amyloid formation in the presence of negatively charged vesicles, we exploited the typical ThT assay that utilizes the increase in fluorescence quantum efficiency of ThT when bound to amyloid fibrils (Groenning, 2010; LeVine, 1993). We chose a fixed concentration of αS (70 μM) and selected the concentrations of lipids according to the CD titration curves in order to create conditions at which αS populates both the free in solution and bound to the vesicle states. In effective terms, this means similar alpha-helix content of αS but different lipid concentrations of DOPG and DOPS. We also tested saturating conditions at which essentially all αS molecules are bound to the vesicles.

αS (70 μM) incubated at 37 °C for 100 h did not show an increase in ThT fluorescence, indicating the absence of amyloid fibrils detectable by ThT (Fig. 2). AFM confirmed the lack of amyloid fibrils at this condition. In contrast, the presence of negatively charged DOPG vesicles at lipid/protein molar ratios of 2:1 (10% αS bound to vesicles), 4:2 (20% αS bound) and 11:2 (55% αS bound) triggered the formation of amyloid fibrils by αS (Figs 2a and 2c). At higher DOPG/ αS molar ratios ($\geq 22:4$), where all αS were bound to the vesicles, no increase in ThT fluorescence was observed. DOPS vesicles also promoted aggregation of αS but with different features. At a DOPS/ αS ratio of 10:5 (13% αS bound), protein aggregation was detected, but at higher ratios of DOPS/ αS (i.e., 21 and 56) no ThT fluorescence was observed albeit, according to the CD titration, both free and bound αS states should be present (Figs 2b and 2d). AFM imaging of end point samples (after 100 h) indicated that in the plateau region of ThT fluorescence, the fibrils

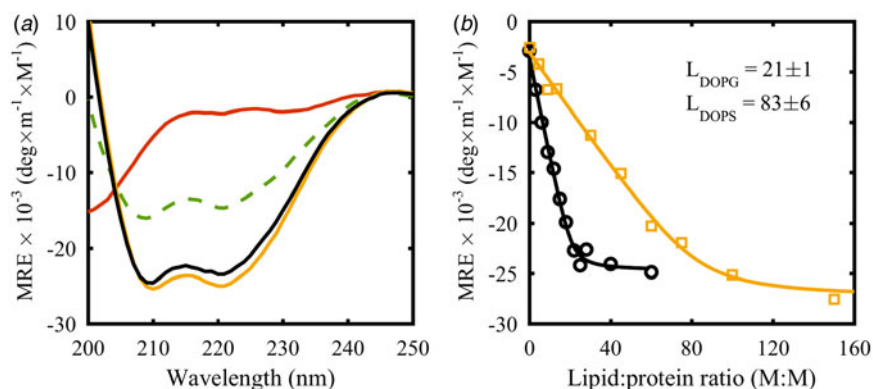


Fig. 1. Secondary structure of α S in the presence of lipid vesicles. (a) CD spectra of 5 μ M α S alone (red) and 5 μ M α S in the presence of 200 μ M of DOPG (black) and 200 μ M (dashed green) and 530 μ M (yellow) of DOPS. (b) Mean residue ellipticity at 222 nm upon titration of DOPG (black, circles) and DOPS (yellow, squares) vesicles into 5 μ M α S solution; the experimental data are shown as symbols and a one-step binding model fit is shown as solid lines.

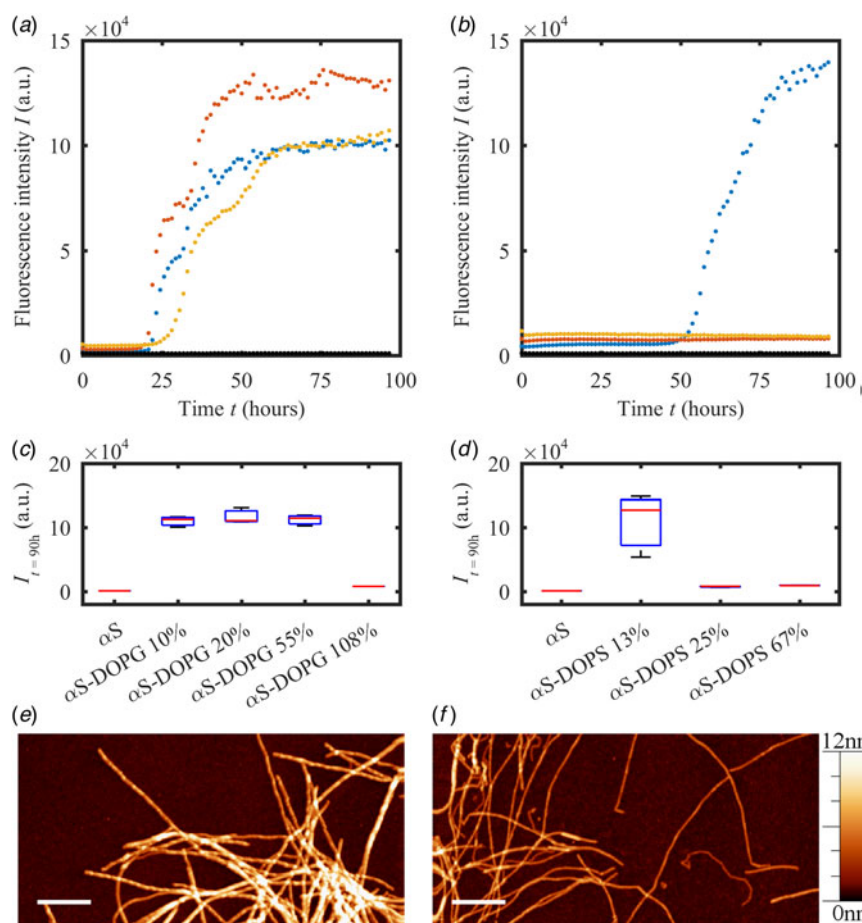


Fig. 2. Aggregation of α S in the presence of negatively charged vesicles. ThT fluorescence of α S alone (black) (a, b) and of α S incubated in the presence of 150 μ M (10% α S bound, blue), 300 μ M (20%, red) and 790 μ M (55%, yellow) of DOPG (a) or in the presence of 735 μ M (13%, blue), 1470 μ M (25%, red) and 3920 μ M (67%, yellow) of DOPS (b). Boxplots comparing ThT intensity at plateau ($t = 90$ h) of α S incubated in the presence of DOPG vesicles (c) and DOPS vesicles (d). AFM images of α S amyloid fibrils at the end point of the aggregation experiment after incubation of monomeric α S in the presence of 300 μ M DOPG (e) and 735 μ M DOPS (f) vesicles. Scale bars 500 nm; height color-scale is shown on the right.



formed in the presence of both types of negatively charged vesicles were heterogeneous, with some fibers of dimensions of 7–9 nm in height as well as some with dimensions <5 nm in height (Figs 2e and 2f). For comparison, these amyloid fiber properties are in agreement with those of α S amyloids formed in the absence of vesicles but in the presence of a small glass bead and agitation to speed up the reaction (Fig. S1).

3.3. α S oligomer ‘contamination’ dramatically alters α S aggregation in the presence of negatively charged vesicles

In the above experiments, stock lyophilized α S was dissolved and purified by SEC prior to use as a standard step to always assure monomeric starting material. Indeed, SEC analysis of lyophilized α S samples showed the presence of protein species with a larger apparent size, indicative of an assembled and oligomeric fraction (Fig. S2). Moreover, AFM analysis demonstrated the presence of spherical structures with heights in the range of 1–3 nm in the lyophilized α S sample (Fig. S2). Based on the SEC analysis, the oligomer fraction corresponds to approximately 5% of total protein and the oligomer size appears to be about 80 kDa (Fig. S2). The oligomers appeared stable as they could not be dissociated by 8 M urea (Fig. S2). As expected for a sample containing mostly monomers in mixture with a small fraction of oligomers, CD titrations curves with negatively charged vesicles using this α S source matched the data collected for monomer-only α S (Fig. S3, cf. Fig. 1).

Lyophilization has been reported to induce the formation of α S oligomers and it has been proposed that these species may affect α S aggregation (Fredenburg *et al.* 2007; Lashuel *et al.* 2002; Volles & Lansbury, 2002). Therefore, we also investigated the aggregation kinetics of the heterogeneous α S preparation (i.e., lyophilized α S dissolved without further purification) in the presence of negatively charged vesicles. The ThT signal of lyophilized α S reconstituted in buffer without vesicles did not change within 100 h at aggregation conditions, indicating no amyloid formation and, importantly, is the same result as that for the monomer-only α S sample in the absence of vesicles (Figs 3a and 3b).

In contrast, for this α S sample, the ThT signal increased almost linearly with time without lag time in the presence of DOPG LUVs at lipid/protein ratios of 2:1, 4:2, 11:2 and DOPS/protein ratios of 10:5 (Figs 3a and 3b). The amplitudes of these ThT changes were however very low, but nonetheless clearly distinct from the flat ThT curves for α S without vesicles. AFM analysis revealed that thin amyloid fibrils, 3–4 nm in height, were present in these samples after the reactions (Figs 3c and 3d). Thus, if a small fraction of oligomers is present along with α S monomers, together with negatively charged vesicles, elongation of a small number of fibrils takes place, most likely with the pre-formed

‘contaminating’ oligomers as starting nuclei. At the same time, and most surprisingly, the typical cooperative aggregation seen for the monomer-only α S samples in the presence of vesicles (Fig. 2) is absent.

4. Discussion

The binding of α S to lipid membranes is described to depend on numerous factors such as solution conditions, chemical properties of the lipids, lipid bilayer phase state and membrane curvature (Galvagnion *et al.* 2016; Middleton & Rhoades, 2010; Ouberaï *et al.* 2013; Shvadchak *et al.* 2011a, b). Under the experimental conditions of this study, induction of the same amount of alpha-helical content in α S required four times higher concentration of DOPS than DOPG vesicles. The area per lipid molecule of DOPS (65.3 Å² (Petrache *et al.* 2004)) is almost 10% smaller than that of DOPG (70.8 Å² (Pan *et al.* 2012)), indicating that attractive interactions between PS head-group are stronger than between PG head-group. Stronger attractive interactions will result in higher molecular order in the polar head-group region of DOPS relative to DOPG lipid layers (Lindblom *et al.* 1991) and increased exposure of hydrophobic parts of DOPG relative to DOPS lipid bilayers. It has been previously described that α S binds with higher affinity to model membranes with defects, indicating that lipid packing strongly affects α S–membrane interactions (Nuscher *et al.* 2004; Ouberaï *et al.* 2013; Shvadchak *et al.* 2011a). Thus, DOPG may expose more hydrophobic areas and α S may insert deeper into the DOPG membrane than the DOPS membrane. This will increase the effective area of binding, since when the bilayer expands, the effective surface area per lipid will increase. This reasoning may explain the lower number of lipid molecules per α S molecule for DOPG compared with DOPS. Different reported affinities of α S for negatively charged head-group, going beyond the total charge, also emphasize specificity of α S among acidic phospholipids (Nakamura *et al.* 2008; Rhoades *et al.* 2006).

CD of α S binding to anionic lipid vesicles shows a transition from mostly disordered to largely alpha-helical conformation of α S. At an excess of lipids, most of α S monomers are bound to the vesicles. We observed a maximum value of mean residue ellipticity (MRE) at 222 nm of -25×10^3 and -27×10^3 deg m⁻¹ M⁻¹ for α S bound to the DOPG and DOPS vesicles, respectively. Using the MRE values and assuming that all α S molecules are bound to the vesicles, we estimate that 68–74% of the protein is in alpha-helical structure (Scholtz *et al.* 1991), which corresponds to ~100 out of the 140 residues of α S. This result agrees with structural studies that have shown that the first 100 N-terminal residues of α S interact with lipid vesicles (Bodner *et al.* 2009, 2010). The number of α S residues bound to the vesicles can be estimated by assuming that the dimensions of the alpha-helix are ~0.13 nm per residue

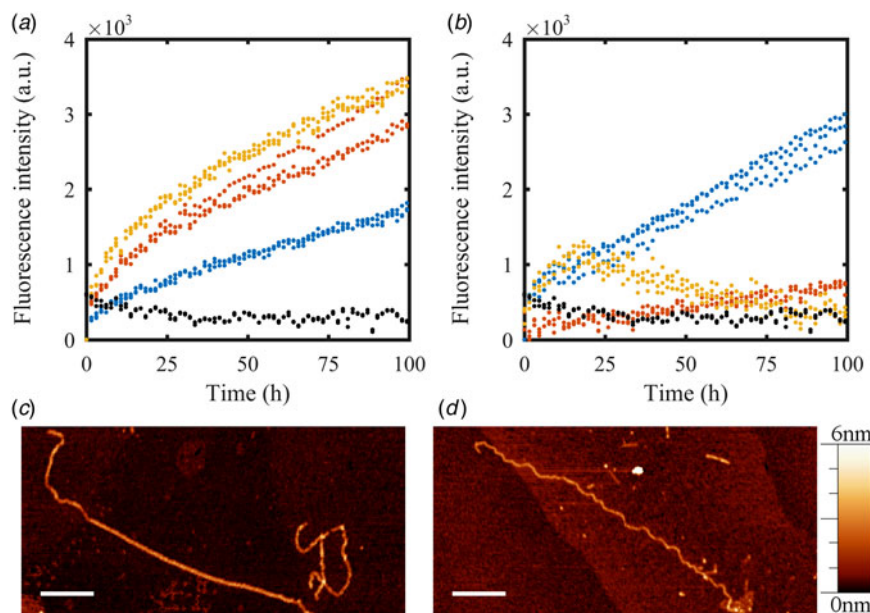


Fig. 3. Aggregation of αS , consisting of monomers and a small fraction of oligomers, in the presence of negatively charged vesicles. ThT fluorescence of αS alone (black) (a, b) and of αS incubated in the presence of 150 μM (blue), 300 μM (red), 790 μM (yellow) of DOPG vesicles (a) or in the presence of 735 μM (blue), 1470 μM (red) and 3920 μM (yellow) of DOPS vesicles (b). We note that the ThT intensity in these graphs are approximately 20-fold lower than in Figs 2a and 2b. AFM images of αS amyloid fibrils at the end point of the aggregation experiment after incubation in the presence of 300 μM DOPG (c) and 735 μM DOPS (d) vesicles. Scale bars 500 nm; height color-scale is shown on the right.

in length (Lee *et al.* 2004) and 1 nm in width in combination with the lipid-to-protein ratio at saturation determined here of 21 (lipid area of 7.5 nm²) for DOPG vesicles and 83 (lipid area of 27 nm²) for DOPS. We arrive at the result that only about 58 residues of each αS can be accommodated on DOPG vesicles at saturation, whereas on DOPS surfaces the available area per αS monomer corresponds to that of 200 residues in alpha-helical conformation.

This simple analysis may suggest that αS induces a rearrangement of the DOPG bilayer resulting in lateral expansion such that accommodation of 100 αS residues per molecule is possible. In accord, it has been proposed that αS binding to lipid bilayers can involve insertion of helical segments in the head-group region resulting in lateral expansion; importantly, this effect was highly dependent on the degree of order in the membrane (Ouberaï *et al.* 2013). Another possibility is that part of the αS helix is protruding from the membrane surface of the DOPG vesicles. It was previously reported that binding of αS to PS-containing vesicles requires all residues 1–102, whereas the binding to phosphatidic acid containing vesicles can be mediated by just one of the sequence segments 1–42, 43–56 or 56–102 (Perrin *et al.* 2000).

We observed that DOPG and DOPS vesicles modulate differently amyloid formation of monomeric αS . At lipid/protein ratios high enough to have all αS molecules bound to the vesicles, the protein did not aggregate within the time

frame of our experiments (100 h). However, for both vesicle types, when free monomers are present in solution, fibril formation took place and the process showed an initial lag phase, which is attributed to a nucleation-dependent mechanism (Fink, 2006; Wood *et al.* 1999). As the nucleation of αS occurred faster in the presence of DOPG vesicles than in their absence, we can assume that it is the lipid-bound αS fraction that forms the initial nuclei. Probably, the lipid environment increases the local concentration of αS , promoting self-assembly of membrane-bound αS molecules and consequently leading to the formation of nuclei on the bilayer surface. If we make the assumption that alpha-helix content corresponds to bound αS , then DOPG vesicles induce amyloid formation of monomeric αS as long as both bound and free protein states are present. In contrast, in the case of DOPS, aggregation of αS occurred only for 13% of αS bound to lipids and the lag phase with DOPS was longer than for αS with DOPG at the same percentage of vesicle-bound protein. Higher DOPS-bound fraction of αS than 13% resulted in the absence of any amyloid formation. Also when comparing the DOPS and DOPG results for similar absolute lipid concentrations, corresponding to lipid/protein ratio of 10.5 and 11.2, respectively, the lag times of αS amyloid formation are still much longer in the case of DOPS than DOPG. These data suggest that efficiency of αS nucleation on vesicles is dependent on vesicle surface chemistry, dynamics and structure, which together define the αS binding mode. It was previously suggested



that α S aggregation is promoted only by lipids with short saturated hydrocarbon chains (12 or 14 carbon chain) (Galvagnion *et al.* 2016). Our results here show that α S aggregation can also be promoted by long unsaturated hydrocarbon chains (i.e., DOPS, DOPG; 18 carbon chain with one double bond) at some conditions. Based on previous studies, it is possible that α S aggregation on vesicles involves extraction of lipids from the bilayer and partial protein–lipid co-aggregation (Hellstrand *et al.* 2013; Reynolds *et al.* 2011).

Unexpectedly, we found that α S amyloid formation in the presence of vesicles changes dramatically when a small fraction of oligomers is present. The AFM data suggested the formation of a small fraction of amyloid fibrils that were thin (Fink, 2006; Khurana *et al.* 2003). These amyloid fibrils grew without lag phase implying that the pre-formed ‘contaminating’ oligomers were the source of nuclei. Notably, this process blocked the aggregation of the remaining monomers, which contrasts the result for monomers-only α S samples in the presence of negatively charged vesicles. Since amyloid fibrils were absent from aggregation reactions of this α S sample without vesicles, the ‘contaminating’ oligomers must depend on the vesicles for nucleation of amyloid formation. α S oligomers are described to interact with negatively charged bilayers and may cause membrane disruption (Fredenburg *et al.* 2007; van Rooijen *et al.* 2009). The molecular details of oligomer–membrane interactions and the effect on surrounding monomeric species by those interactions are still unclear. If α S interactions with lipid vesicles are cooperative, then even a few (or one) oligomers may alter the way many monomers arrange on the surface of the liposome. Notably, a study on α S aggregation on supported lipid bilayers showed that monomeric α S has a higher affinity for protein aggregates on the surface than for the lipid bilayer itself (Reynolds *et al.* 2011). Moreover, similar to our results here, the study showed that after the initial formation of the first aggregates, these grew with time but the formation of new α S seeds occurs at much lower frequency.

Our findings point out the importance of careful assessment of membrane properties, and synergistic effects due to protein–lipid interactions, as well as both lipid vesicle and protein structural dynamics. Our study also highlights the importance of protein preparation procedures. The literature around α S aggregation includes a wide variation of results and at least some of these may be explained by differences in protein starting material.

4.1. Speculation box

Based on this work, we speculate that consequences of α S interactions with synaptic vesicles and membranes *in vivo* will depend on simultaneous interactions with other proteins that may modulate vesicle surface properties. It

appears that binding of pre-formed oligomers to vesicles alters surface properties such that monomers are restricted from independent formation of amyloid seeds. Our results also highlight the question of what *in vitro* conditions are most relevant for the *in vivo* scenario that ultimately results in PD. Finally, we note that to gain real biological bearing of these *in vitro* findings, vesicles composed of mixed lipids mimicking those in neuronal cells should be investigated.

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Conflict of interest

None.

Supplementary material

The supplementary material for this article can be found at <https://doi.org/10.1017/S0033583517000026>

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