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Hamley, I. W. ORCID: <https://orcid.org/0000-0002-4549-0926>, Hutchinson, J., Kirkham, S., Castelletto, V., Kaur, A., Reza, M. and Ruokolainen, J. (2016) Nanosheet formation by an anionic surfactant-like peptide and modulation of self-assembly through ionic complexation. *Langmuir*, 32 (40). pp. 10387-10393. ISSN 0743-7463 doi: 10.1021/acs.langmuir.6b02180 Available at <https://centaur.reading.ac.uk/68326/>

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Published version at: <http://dx.doi.org/10.1021/acs.langmuir.6b02180>

To link to this article DOI: <http://dx.doi.org/10.1021/acs.langmuir.6b02180>

Publisher: American Chemical Society

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Nanosheet Formation by an Anionic Surfactant-like Peptide and Modulation of Self-Assembly through Ionic Complexation

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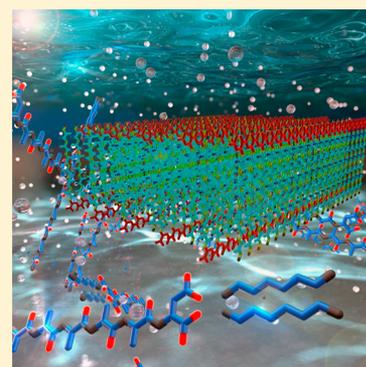
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S Supporting Information

ABSTRACT: The surfactant-like peptide (Ala)₆-(Asp) (A₆D) is shown to self-assemble into ultrathin (3 nm thick) nanosheets in aqueous solution above a critical aggregation concentration. A combination of circular dichroism and FTIR spectroscopy and X-ray diffraction shows that the nanosheets comprise interdigitated bilayers of the peptide with β -sheet conformation. The self-assembly can be modulated by addition of hexamethylenediamine which is expected to interact with the anionic C terminus (and C-terminal D residue) of the peptide. Multiple ordered nanostructures can be accessed depending on the amount of added diamine. Nanosheet and bicontinuous network structures were observed using cryogenic-TEM and small-angle X-ray scattering. Addition of hexamethylenediamine at a sufficiently large molar ratio leads to disruption of the ordered nanostructure and the formation of a solution of A₆D–diamine molecular complexes with highly charged end groups. The multiple acid-functionalized nanostructures that are accessible in this system are expected to have many applications in the fabrication of new nanomaterials.



INTRODUCTION

The formation of free-floating ultrathin nanosheet structures has been observed for several classes of systems including assemblies with graphene or graphene oxide,^{1,2} two-dimensional metal oxide sheets,^{3,4} various exfoliated inorganic structures,⁵ polymer⁶ and block copolymer crystals,⁷ and peptoids.^{8–10} Nanosheet formation by peptides is rarely reported,^{11–16} although such structures offer the potential to prepare functional nanostructures coated at high density with bioactive amino acids.

Surfactant-like peptides (SLPs) have a remarkable ability to self-assemble into different nanostructures, primarily due to their amphiphilic nature. For example, they can aggregate into high aspect ratio structures while displaying bioactive peptides. SLPs are a class of amphiphilic peptide comprising a headgroup which is a short sequence of charged residues attached to a tailgroup of neutral residues.^{17–19} Pioneering work on SLPs has been conducted by the Zhang group including a series of peptides with neutral amino acid tails (based on valine, alanine, or glycine) and charged residues such as D (aspartic acid) or K (lysine).^{20–23} We previously reported the unexpected observation that the SLP A₆R self-assembles into free-floating ultrathin (3 nm thick) nanosheets in dilute aqueous solution.¹² At higher concentration, the nanosheets wrap into helical ribbons and nanotubes. Here we report on the self-assembly of the related SLP A₆D with oppositely charged headgroup compared to A₆R, and we show that the self-assembly can be modulated by

addition of diamine, expected to bind to the peptide C terminus and the carboxyl group in the D residue.

The self-assembly of Ac–A₆D–OH (N-terminal capped; charge of –2 at pH 7) has been studied in phosphate-buffered solution (PBS)²⁴ or in water.²¹ In PBS, light scattering revealed a critical aggregation concentration (CAC), and above the CAC, atomic force microscopy revealed extended structures interpreted as nanotubes, or nanotubes with bulges (which alternatively appear to be like bead aggregate structures).²⁴ The results were compared to those for peptide Ac–A₆K–CONH₂ which it is also claimed self-assembles into nanotubes.²⁴ The influence on the CAC and on the self-assembled nanostructure was investigated for mixtures of Ac–A₆D–OH and Ac–A₆K–NH₂.²⁵ “Nanorope” structures were observed for the Ac–A₆D–OH/Ac–A₆K–NH₂ = 2:1 molar ratio mixture.²⁵ A study using circular dichroism to determine secondary structure found that Ac–A₆D–OH adopts a mostly random coil structure in dilute aqueous solution, with some β -sheet content at higher concentration.²⁶ Peptides including Ac–A₆D–OH have been used as surfactants in commercial *Escherichia coli* (*E. coli*) cell-free systems to produce soluble G-protein coupled receptors.²⁷ Another potential application involves nanovesicles for drug delivery since Ac–A₆D–OH has been reported to form

Received: June 13, 2016

Revised: August 30, 2016

Published: September 16, 2016

nanovesicles and nanovesicle clusters under defined phosphate buffered saline conditions (in contrast to the above-mentioned works).²⁸ This peptide has also been used along with a zwitterionic surfactant to template the formation and deposition (at nanotube or nanobelt walls) of gold nanoparticles.²⁹ The influence of SLPs such as A₆D on the structure of ternary mixtures with the lipid mono-olein and water has been examined, and it was shown to be possible to tune the size of the bicontinuous cubic phase lattice or to induce the formation of an inverse hexagonal structure.³⁰

Here we report on the self-assembly, and secondary structure, of NH₂-A₆D-COOH. We note that this differs from the peptide studied by Zhang et al. as it is uncapped at both termini. We also investigate self-assembly in mixtures of NH₂-A₆D-COOH with hexamethylenediamine (HEMED, 1,6-hexanediamine) and the effect of this diamine on peptide conformation. The diamine is expected to form complexes with A₆D which will have a charge of +1 at pH 7 as it contains an ionizable -NH₂ group at the N terminus, an aspartic acid residue, and a carboxyl group at the C terminus. Diamines such as tetramethylenediamine (putrescine) or pentamethylenediamine (cadaverine) are important biogenic amines with a variety of functions. Here, we chose to study the complexation of A₆D with hexamethylenediamine from the same series of short flexible amines.

The complexation of various amines including ethylene-, tetramethylene-, and pentamethylene diamine and spermine and spermidine with the anionic surfactant SDS has been investigated, and it has been shown that enhanced adsorption at the air-water interface results.^{31,32} However, to the best of our knowledge, complexation of surfactant-like peptides with diamines has not previously been examined. We show that addition of hexamethylenediamine significantly modifies the self-assembled nanostructure and that it is possible to produce nanotape and platelet structures in this way. As far as we are aware, platelet peptide nanostructures have not previously been noted.

EXPERIMENTAL SECTION

Materials. Peptide NH₂-AAAAAD-COOH, referred to as A₆D, was custom synthesized by CS Bio Co. (Menlo Park, CA, USA) and was received as the TFA salt variant. Purity was 99.87% by HPLC in water/acetonitrile (0.1% TFA). Electrospray-ionization mass spectroscopy (ESI-MS) indicated a molar mass of 559.33 g mol⁻¹ (559.59 g mol⁻¹, expected).

Pyrene Fluorescence. Fluorescence spectra were recorded with a Varian Cary Eclipse fluorescence spectrometer with samples in 4 mm inner quartz cuvettes. The assays were performed using 1.3 × 10⁻³ to 0.13 wt % A₆D, in 2.3 × 10⁻⁵ wt % pyrene solution, and the same A₆D concentration range with 2:1, 1:1, and 1:2 molar ratios to hexamethylenediamine and 1.3 × 10⁻⁴ wt % pyrene solution. The samples were excited at λ_{ex} = 339 nm, and the fluorescence emission was measured for λ = 360–500 nm.

Titration pH Measurements. A vial was filled with 1 wt % (17.88 mM) A₆D to which aliquots of a 71.52 mM diamine solution were added. pH was measured using a Mettler Toledo Five Easy FE20 instrument with a Sigma-Aldrich micro pH combination electrode, at various points as more hexamethylenediamine solution was added, until a ratio of 1:2 peptide:diamine was reached.

Circular Dichroism Spectroscopy. Circular dichroism (CD) spectra were recorded using a Chirascan spectropolarimeter (Applied Photophysics, Leatherhead, U.K.). The samples, with A₆D always at 1%, were placed in a coverslip cuvette (0.2 mm thick), spectra presented being with absorbance A < 2 at any measured point with a 0.5 nm step, 1 nm bandwidth, and 1 s collection time per step. The

CD signal from the background was subtracted from the CD data of the sample solutions.

Fourier Transform Infrared Spectroscopy. Spectra were recorded using a Thermo Scientific Nicolet ISS or a Nexus-FTIR spectrometer, both equipped with a DTGS detector. In the former, a PEARL liquid cell was used (sample contained between fixed CaF₂ plates). In the Nexus spectrometer, 20 μL of each sample was sandwiched between two CaF₂ plate windows (spacer, 0.012 mm thick). Spectra were scanned 128 times over the range of 900–4000 cm⁻¹.

Cryogenic Transmission Electron Microscopy. Imaging was carried out using a field emission cryo-electron microscope (JEOL JEM-3200FSC), operating at 200 kV. Images were taken in bright field mode and using zero loss energy filtering (Ω type) with a slit width of 20 eV. Micrographs were recorded using a Gatan Ultrascan 4000 CCD camera. The specimen temperature was maintained at -187 °C during the imaging. Vitrified specimens were prepared using an automated FEI Vitrobot device using Quantifoil 3.5/1 holey carbon copper grids with a hole size of 3.5 μm. Just prior to use, grids were plasma cleaned using a Gatan Solarus 9500 plasma cleaner and then transferred into an environmental chamber of a FEI Vitrobot at room temperature and 100% humidity. Thereafter 3 mL of sample solution was applied onto the grid, and it was blotted twice for 5 s and then vitrified in a 1/1 mixture of liquid ethane and propane at a temperature of -180 °C. The grids with vitrified sample solution were maintained at liquid nitrogen temperature and then cryotransferred to the microscope.

X-ray Diffraction. Measurements were performed on stalks prepared by drying filaments of solutions A₆D. The stalk for A₆D was prepared from a 1 wt % solution, but for the solutions containing mixtures of A₆D with diamine, 5 wt % solutions were used. Solutions of the mixtures were suspended between the ends of wax-coated capillaries and dried. The stalks were mounted (vertically) onto the four axis goniometer of an Oxford Diffraction Gemini Ultra instrument. The sample-detector distance was 44 mm. The X-ray wavelength was λ = 1.54 Å. The wavenumber scale (q = 4π sin θ/λ where 2θ is the scattering angle) was geometrically calculated. The detector was a Sapphire CCD.

Small-Angle X-ray Scattering. Collection of solution small-angle X-ray scattering (SAXS) data was performed on the bioSAXS beamline BM29 at the ESRF, Grenoble, France. Solutions containing 1 wt % A₆D or 1 wt % A₆D mixed with hexamethylenediamine were loaded in PCR tubes in an automated sample changer. SAXS data were collected using a Pilatus 1 M detector. The sample-detector distance was 2.84 m. The X-ray wavelength was 0.99 Å. The wavenumber q = 4π sin θ/λ scale was calibrated using silver behenate.

Zeta Potential Measurements. The ζ potential was measured using a Zetasizer Nano 2s from Malvern Instruments. A 1 mL aliquot of sample was placed inside a disposable folded capillary cell. The sample was left to equilibrate for 120 s before measuring the ζ potential, using an applied voltage of 50.0 V. The results presented are the average over three measurements.

RESULTS

We first determined whether the addition of hexamethylenediamine has any influence on the potential aggregation behavior of A₆D via critical aggregation concentration (CAC) assays. Pyrene is a dye whose fluorescence is sensitive to local hydrophobic environment, widely used for such assays as in our previous work for example on A₆RGD.³³ Figure 1 shows results from pyrene fluorescence CAC assays. A discontinuous increase in intensity occurs at the CAC. A₆D itself shows a CAC = 0.030 ± 0.005 wt % with a similar value for 2:1 A₆D:diamine (in this and the following where concentration ratios are given, they refer to molar concentrations). The value for 1:1 A₆D:diamine mixtures is lower, CAC = 0.0063 ± 0.003 wt %. This presumably reflects the enhanced hydrophobicity resulting from the screening of the charge on the carboxyl group on the D residue and at the C terminus under these conditions. Most

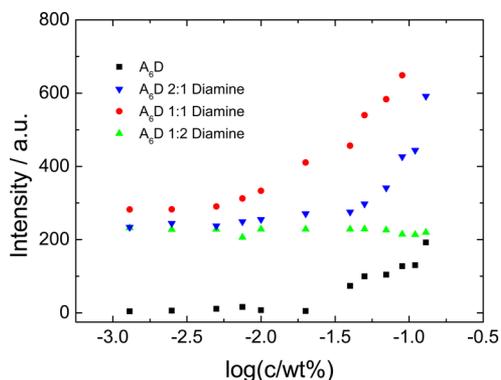


Figure 1. Critical aggregation concentration (CAC) assay using concentration-dependent pyrene (first vibronic band) fluorescence, I_1 .

remarkably, however, the CAC for the 1:2 A_6D :diamine system appears to be entirely suppressed.

This influence of molecular conformation on this unexpected phenomenon was examined using circular dichroism (CD) spectroscopy. The measured spectra are shown in Figure 2a.

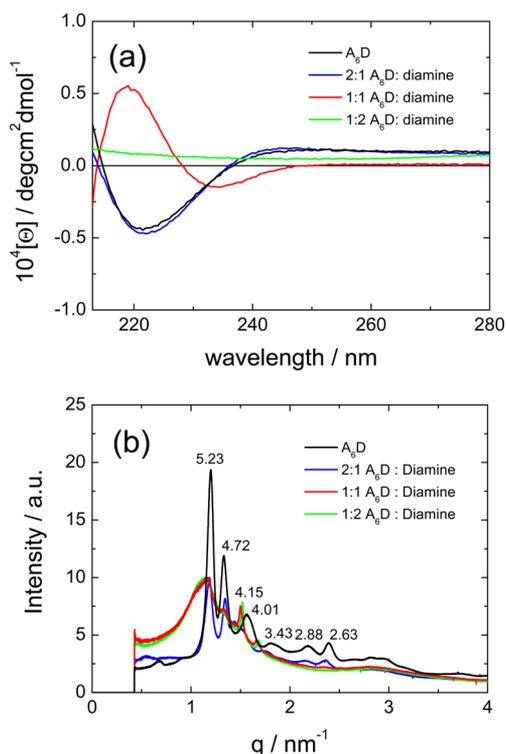


Figure 2. (a) CD spectra for mixtures of A_6D with hexamethylenediamine for the solutions indicated (constant A_6D concentration, 1 wt %; ratios of A_6D :diamine are molar ratios). (b) X-ray diffraction intensity profiles for A_6D with hexamethylenediamine under the conditions indicated. For the A_6D profile, d -spacings (\AA) are indicated as well as the 4.15 \AA spacing for two of the other samples. Stalks were prepared from solutions with a constant A_6D concentration, 1 wt %.

The spectra for A_6D and 2:1 A_6D :diamine exhibit minima near 220 nm and can be assigned to peptide β -sheet conformation.^{34,35} In contrast, the spectrum for a 1:1 A_6D :diamine mixture is distinct with a maximum at 219 nm and a minimum at 234 nm and is ascribed to disordered conformation and/or a contribution from a turn structure.³⁶ Remarkably, CD reveals the absence of secondary structure for the 1:2 A_6D :diamine

mixture. This correlates with the results from the pyrene fluorescence CAC assay for this sample which shows no break point and thus no evidence for aggregation. These findings are supported by FTIR (Supporting Information (SI) Figure S1) which show a clear amide I' peak assignable to β -sheet structure^{37–39} at 1625 cm^{-1} for A_6D and the 2:1 mixture. However, this peak is replaced by a broad peak at 1644 cm^{-1} for the 1:1 and 1:2 mixtures, which is due to unordered conformation. Also evident is a small peak at 1700 cm^{-1} for the A_6D and 2:1 mixture, which is absent for the other two mixtures. This peak is assigned to C=O stretch vibrations,^{37,40} and its absence for the 1:1 and 1:2 mixtures is likely due to binding of the diamine. All spectra also have a peak at 1672 cm^{-1} due to TFA counterions bound to the N terminus of the peptide.⁴¹ There are also shifts in the intensity of the amide II' peak at 1550 cm^{-1} due to N–H/C–N deformations upon variation of diamine concentration.³⁷ The FTIR spectra in the 3000–3800 cm^{-1} range also show a dramatic transition comparing the spectra for A_6D and the 2:1 mixture with those for the 1:1 and 1:2 mixtures—the spectra in each pair showing features similar to each other. In particular, the sharp peak near 3290 cm^{-1} for A_6D and the high A_6D content mixture can be ascribed to N–H stretching vibrations within peptide chains (non-H-bonded), whereas this peak is absent from the spectra for the 1:1 and 1:2 mixtures which contain a broad peak near 3400 cm^{-1} which is due to H-bonded N–H or O–H stretch modes.^{37,40}

Confirmation of β -sheet structure is provided by fiber X-ray diffraction (XRD). The isotropic scattering patterns were reduced to one-dimensional intensity profiles as shown in Figure 2b. A_6D exhibits a series of reflections from an alanine-rich β -sheet structure which have unusually small β -sheet spacings^{12,42–45} (here 5.23 \AA) along with the 4.72 \AA β -strand spacing, and other peaks from the well ordered packing of the alanine residues. The 2:1 mixture sample shows the same features. In complete contrast, for the 1:1 and 1:2 A_6D :diamine samples these β -sheet features are absent and instead there is a broad halo in the XRD intensity profile corresponding to a d -spacing around 5.5 \AA along with a peak corresponding to 4.15 \AA . These features can be assigned to unordered peptide.

The distinct aggregation behavior in the mixtures with different diamine content was examined by cryogenic transmission electron microscopy (cryo-TEM; Figure 3). Cryo-TEM images for 1 wt % A_6D reveal the presence of thin nanosheet structures (Figure 3a and SI Figure S2) coexisting with irregular continuous network-like structures. We previously reported the formation of peptide nanosheets (under similar concentration conditions) for the related peptide with cationic headgroup A_6R .¹² Nanosheet-like structures were also observed for solutions of the 2:1 mixture although here with a more defined width; i.e., nanotape-like structures are observed (Figure 3b and SI Figure S3). For the 1:1 A_6D :diamine solutions there was evidence for a high density of plate-like structures (Figure 3c and SI Figure 4) as well as a region of irregular mesh-like structure. To our knowledge, such platelets have not previously been reported for self-assembling peptides. In contrast, no self-assembled structures were imaged for the 1:2 A_6D :diamine mixtures, consistent with the lack of a CAC and the absence of observed secondary structure characteristics in the CD spectrum.

Small-angle X-ray scattering (SAXS) was used to further probe the nature of the self-assembled nanostructures in solution. The corresponding measured intensity profiles with

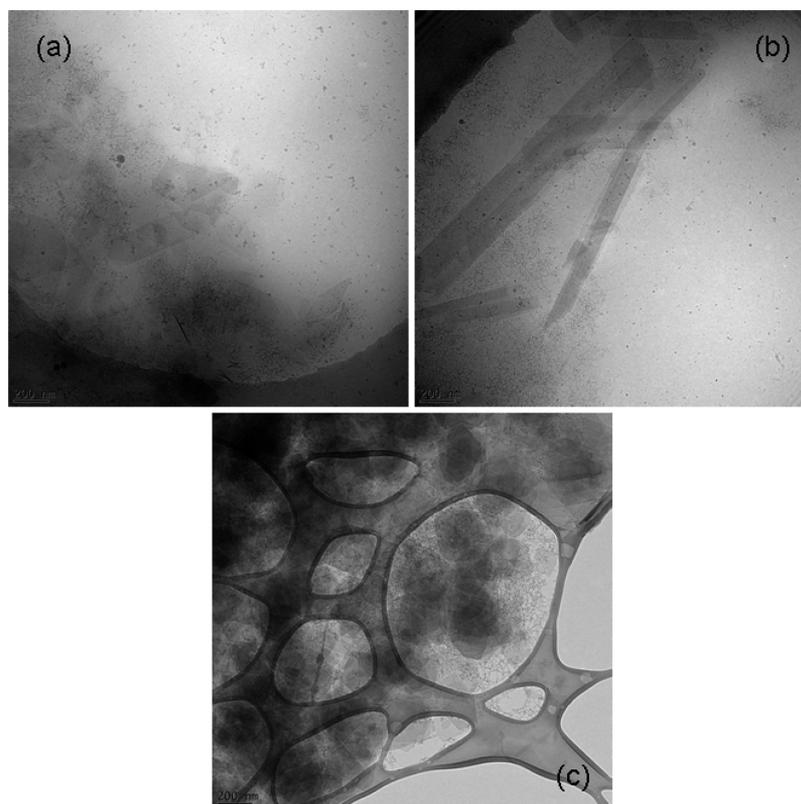


Figure 3. Cryo-TEM images: (a) 1 wt % A_6D ; (b) 2:1 A_6D :diamine; (c) 1:1 A_6D :diamine. (Constant A_6D concentration, 1 wt %; ratios of A_6D :diamine are molar ratios.)

model form factor fits are shown in Figure 4. The fit parameters are listed in Table 1. The profile for 1 wt % A_6D is very similar

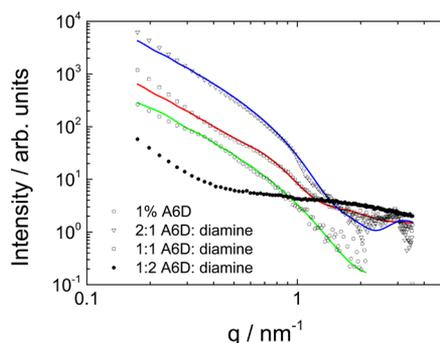


Figure 4. SAXS data (open symbols) with model form factor fits (solid lines). The fit parameters are listed in SI Table 2. For clarity, only every fifth measured data point is shown.

to that previously reported for 1 wt % A_6R .¹² The data can be fitted to a “Gaussian bilayer” form factor,⁴⁶ as described elsewhere,^{47–49} this form factor describes the electron density profile of a planar bilayer structure with three Gaussian functions representing the electron density of the core (here, hydrophobic A residues) and two for the surfaces (here, hydrophilic D residues). Here, similar to the case of A_6R ,¹² we propose that A_6D actually self-assembles into “monolayers” with a core of interdigitated A_6 chains and surfaces of D residues (Figure 5). This is consistent with the fact that the layer thickness is determined to be 2.9 nm, which is slightly larger than the estimated length of the chain in a parallel β -sheet ($7 \times 0.32 \text{ nm}^{50} = 2.2 \text{ nm}$) indicating interdigitation of the

opposed chains (Figure 5). The low q part of the SAXS data for the 2:1 A_6D :diamine mixture can also be fitted using the Gaussian bilayer model, consistent with the cryo-TEM image which shows flat and wrapped layered structures (Figure 3b). At high q , a broad Bragg peak at $q_0 = 2.9 \text{ nm}^{-1}$ is observed, corresponding to a structure with a periodicity $d = 2.2 \text{ nm}$. This is close to (but a little smaller than) the bilayer thickness obtained from the form factor fits, but identical to the estimated molecular length. Its presence suggests that some stacked layer structures (as well as isolated “monolayer” nanosheets) exist in the 2:1 mixture solution. The spacing of the stacked layer structures is smaller than that of the nanosheets, indicating complete interdigitation of the A_6D peptide chains in this case. This Bragg peak was not accounted for in the form factor fitting. The SAXS data for the 1:1 mixture show a scaling law-type decay of the intensity at low q similar to that for the A_6D and 2:1 mixtures (Figure 4). Because the cryo-TEM image for the 1:1 mixture shows flat platelets coexisting with irregular mesh structures (Figure 3c), the Gaussian bilayer form factor was supplemented with a contribution to allow for the mesh structure. For this purpose, we used the “broad peak” function (described in the footnotes in Table 1) to account for the presence of a broad peak in the measured SAXS data centered around $q_0 = 0.64 \text{ nm}^{-1}$ (Figure 4). The SAXS profile for the 1:2 A_6D :diamine solution showed a shape completely different from those for the other solutions, in particular a lack of any peak at higher q due to a bilayer form factor (Figure 4), and was not fitted to a Gaussian bilayer form factor, consistent with the absence of aggregated structures in the cryo-TEM images. The SAXS profile for diamine alone (not shown) was not distinguishable from background (as expected for a dilute molecular solution).

Table 1. SAXS Fitting Parameters (Fitted Using Gaussian Bilayer Form Factor⁴⁶)

sample	t^a /nm	σ_{core}^b σ_{out}^b /nm	ν_{core}^c	ν_{out}^d	Gaussian polydispersity params (t^e)	background ^f
A ₆ D	2.9	0.5, 0.5	0.030	0.010	$N = 0.20$, $\sigma = 0.7$	0.1
2:1 A ₆ D:diamine	2.9	0.3, 0.3	0.027	0.025	$N = 0.8$, $\sigma = 2.9$	0.1
1:1 A ₆ D:diamine ^g	2.9	0.3, 0.3	0.0062	0.012	$N = 0.8$, $\sigma = 0.7$	1.2

^aBilayer thickness. ^bWidth for core (lipid chain) electron density, σ_{core} , and outer (peptide) electron density, σ_{out} . Gaussian functions (fixed). ^cScattering contrast of hydrophobic core of bilayer (with respect to solvent) (arbitrary units). ^dScattering contrast of hydrophilic surface of bilayer (with respect to solvent) (arbitrary units). ^eGaussian peak height N and half-width σ . ^fAs well as the constant fixed background, the radius of the bilayer objects was fixed at 50 nm. Since this is much larger than t , it does not influence the shape of the scattering profile. ^gAn additional “broad peak” function⁴⁶ was included in the fit due to the observed mesh structure in the cryo-TEM images and the broad peak in the SAXS profile:

$$I(q) = \frac{I_0}{(1 + (lq - q_0|\xi|^m)^p)}$$

The parameters fitted were $I_0 = 13.1$, $\xi = 2.7$, $q_0 = 0.64$, $m = 1.82$, and $p = 1.65$.

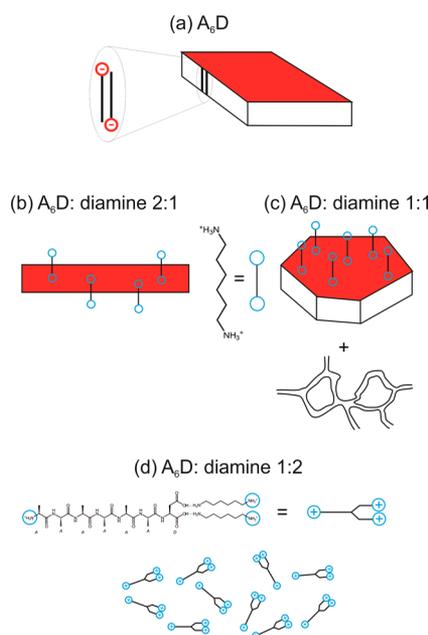


Figure 5. Schematic of self-assembly process: A₆D versus A₆D with hexamethylene diamine. (a) A₆D forms nanosheets 2.2 nm thick comprising interdigitated molecules. (b) In the 2:1 mixture, binding of some diamine molecules drives the formation of nanotape structures. (c) For the 1:1 mixture, a higher degree of binding leads to platelets coexisting with a disordered mesh structure. (d) For the 1:2 mixture, H-bonding interactions between the peptide C terminus and the -NH₂ groups at one end of the diamine molecules lead to complexes with charged end groups as shown in solution, and these are not able to undergo self-assembly due to electrostatic repulsion.

The information on nanostructure provided by cryo-TEM and SAXS (supported by the data on peptide conformation obtained from CD, FTIR, and XRD data) leads to the proposed schematic model shown in Figure 5. A₆D forms nanosheets 2.9 nm thick comprising interdigitated molecules. In the 2:1 mixture, binding of some diamine molecules (to the carboxyl groups on the D residue and C terminus) drives the formation of nanotape structures. For the 1:1 mixture, a higher degree of binding leads to platelets coexisting with a disordered mesh structure. At the molecular level, the peptide loses its β -sheet conformation. For the 1:2 mixture, FTIR shows the presence of H-bonding interactions which stabilize complexes of one A₆D molecule with two diamine molecules (Figure 5), leading to highly charged species which due to electrostatic repulsion of

the free diamine -NH₂ groups are stabilized in solution and do not undergo self-assembly. Data from measurements of pH titrating the hexamethylenediamine into A₆D are shown in SI Figure S5 and show that in the range up to around pH 10 the carboxyl and diamine amine groups are expected to be ionized since the pK_a values of hexamethylenediamine are $pK_{a,1} = 11.02$ and $pK_{a,2} = 10.24$,^{51,52} however, for the 2:1 mixture the diamine may not be fully charged; indeed the FTIR spectra suggest the presence of H-bonding interactions between uncharged NH₂ groups and COOH groups on the peptide. Note that the peptide N terminus is not necessarily expected to be fully charged for higher diamine content mixtures as its pK_a is typically considered to be approximately 8,^{50,53} although values up to pK_a 9.1 have been reported.⁵⁴ ζ potential measurements (SI Table S1) indicate the presence of negatively charged species in all A₆D:diamine mixtures and also show that the ζ potential in the 1:2 A₆D:diamine mixture is higher than in 1:1 or 2:1 mixtures suggesting that the peptide N terminus remains charged.

CONCLUSIONS

In summary, A₆D self-assembles into ultrathin (3 nm thick) nanosheets above a critical aggregation concentration. The nanosheets are bilayers comprising interdigitated antiparallel A₆D molecules in a β -sheet arrangement, leading to the covering of the nanosheet surfaces with aspartic acid residues. The presentation of such acidic residues at high density on a planar surface may be useful in future applications. It is presumed that the high charge density at the surfaces stabilizes a colloidal suspension of freely floating nanosheets.

Addition of a diamine is shown to be a valuable tool to modulate the self-assembly behavior and is a route to prepare acid-coated nanotapes and platelets upon addition of an appropriate amount of diamine. Addition of further diamine leads to the loss of β -sheet secondary structure and the formation of platelets coexisting with a bicontinuous structure. Upon addition of further amounts (two molar equivalents) of diamine, the self-assembled structure (and secondary structure) are lost completely and charged A₆D–diamine complexes are stable as isolated species in solution. The tight packing of the alanine residues (as evidenced by the very small β -sheet stacking distance revealed by the XRD studies discussed above) in A₆D (and A₆R studied previously¹²) is believed to favor the formation of densely packed nanosheets. The A₆D nanosheets have multiple possible uses as biocompatible materials or substrates due to their functional bioactive and/or highly acidic

nature. Acid-functionalized nanotubes, nanosheets, and platelets also have great potential as templates to create new nanomaterials.

■ ASSOCIATED CONTENT

Supporting Information

The Supporting Information is available free of charge on the ACS Publications website at DOI: 10.1021/acs.langmuir.6b02180.

ζ potential measurement results, FTIR spectra, additional cryo-TEM images, and pH titration results (PDF)

■ AUTHOR INFORMATION

Notes

The authors declare no competing financial interest.

■ ACKNOWLEDGMENTS

This work was supported by EPSRC Grant EP/L020599/1 and STFC Grant ST/L502480/1 to I.W.H. The studentship of J.H. was supported by Medimmune and the University of Reading. We thank Ashkan Dehsorkhi for preliminary measurements on A₆D and Sam Burholt for assistance with SAXS. We thank the ESRF for the award of bioSAXS beamtime on BM29 (ref MX-1769) and Gabriele Giachin for assistance with the measurements. We are grateful to Nick Spencer (University of Reading) for the XRD measurements.

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