

1 **Raman study of lysozyme amyloid fibrils suspended on super-hydrophobic surfaces by shear flow**

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10 **Abstract**

11 The shear flow generated at the rim of a drop evaporating on a micro-fabricated super-hydrophobic
12 surface has been used to suspend and orient single/few lysozyme amyloid fibrils between two pillars for
13 substrate-free characterization. Micro Raman spectroscopy performed on extended fibers evidenced a
14 shift of the Amide I band main peak to the value attributed to β -sheet secondary structure, characteristic
15 of the amyloid fibers. In addition, given the orientation sensitivity of the anisotropic molecule, the Raman
16 signal of the main secondary structure was nicely enhanced for a fiber alignment parallel to the
17 polarization direction of the laser. The substrate-free sample generated by this suspending technique is
18 suitable for other structural analysis methods, where fiber crystals are investigated. It could be further
19 employed for generation of arrays and patterns in a controllable fashion, where bio-compatible material is
20 needed.

21 Keywords: super-hydrophobic micro-patterned surfaces; Raman spectroscopy; amyloid fibrils; β -sheet
22 secondary structure

23 **Introduction**

24 Micro-patterned super-hydrophobic surfaces (SHS) are an ideal platform for the manipulation and
25 characterization of single biological molecules and cells in background free environment [1-5]. The key
26 element for suspending long molecules between two adjacent pillars is the shear flow generated at the rim
27 of the evaporating drop [6]. This allows the single molecules to extend across the pillars top, ready for
28 analysis. For example, literature reports DNA and DNA-protein complexes stretching over the micro-
29 pillars substrates using this method, providing suitable samples to be analyzed by different techniques
30 such as scanning electron microscope (SEM), transmission electron microscope (TEM) and Raman
31 Spectroscopy [7-10]. In essence, when a drop of the solution containing the molecules of interest is
32 deposited over this kind of super-hydrophobic surface, it will shrink until a residue accumulates at its
33 center. Through the process, suitably long molecules might pin at the edge of the pillars, and due to the
34 evaporation, they are pulled until they bridge the adjacent pillars in the direction of the flow. This is
35 possible because the shape and design of the micro-fabricated surface maintain the drop quasi-spherical
36 while shrinking, favoring the hopping of the receding meniscus from one pillar's edge to the next.
37 Importantly, by structuring the surface it is possible to spatially organize the stretched molecules.

38 The three dimensional structure of a functional protein has a central role for its activity in the
39 cellular environment and is characterized by the co-presence of three main types of secondary structure,
40 namely α -helix, random coil, and β -sheet. Several pathologies are associated with the disarrangement of
41 the tertiary structure of the protein, for example the diseases related to amyloidosis [11]. Amyloid
42 deposits are characterized by the presence of highly oriented fibers made-up of β -strands organized into
43 β -sheets and further into β -helix or solenoids, generated from the aggregation of misfolded or unfolded

44 proteins which escaped the cellular quality-control [12]. The presence of β -sheet secondary structure can
45 be confirmed by micro Raman spectroscopy, which is a non-destructive, label free technique especially
46 suited for the localized biochemical analysis of biological samples. Appreciably, the results of Raman
47 analysis can complement structural data obtained with other techniques such as NMR and x-ray
48 spectroscopy, their most significant limitation being size (< 40 kDa) and the need for single crystals,
49 respectively [13].

50 In Raman spectroscopy, the Amide I band position, largely shifting around 1650 cm^{-1} , is routinely
51 used to assign the α -helix, random coil, or β -sheet composition of the proteins. The recent literature
52 agrees in assigning to the β -sheet secondary structure a Raman shift in the range $1665\text{-}1675\text{ cm}^{-1}$ [14, 15].
53 An upshift of the Amide I band is evidenced in the transition from functional protein to amyloid structure,
54 where α -synuclein and insulin were considered [16, 17]. In 2007, a first report on functional amyloids
55 derived from different types of silk fibers, highlighted the potential of polarized Raman spectroscopy,
56 while assigning to the β -sheet secondary structure the Raman shift value in the range of $1666\text{-}1670\text{ cm}^{-1}$
57 [18]. They observed that the spectra were dominated by the contribution of the β -sheet, and used the peak
58 height of the polarized Amide I band collected in different scattering geometries to demonstrate that the
59 main Raman tensor axis of the Amide I is highly oriented perpendicular to the fiber direction. A more
60 recent polarized Raman spectroscopy study takes into consideration insulin amyloid fibers aligned on a
61 planar substrate by the coffee ring stain effect [19, 20]. There, an increased intensity of the Raman peak
62 centered at $1672\text{-}4\text{ cm}^{-1}$ emerges in the Raman spectrum when the fiber is aligned parallel to the laser
63 polarization, an indication of the cross- β structure of the fiber core.

64 Here we verify by micro Raman spectroscopy the shift to a predominant β -sheet composition of
65 the hen egg white lysozyme (HEWL) after its conversion to amyloid fibrils. The samples were prepared
66 suspended across the gaps between neighboring pillars, and thus in a substrate and debris free
67 environment. In addition, and for each suspended singular fiber, we observe a clear dependence of the

68 intensity of the peak assigned to the β -sheet secondary structure to its orientation with respect to the
69 exciting laser polarization, with a three-fold counts increase when oriented parallel to the laser
70 polarization as compared to the perpendicular case. These results remark the ability of this technique to
71 align single molecules along one given direction. Fiber suspension by shear flow on super-hydrophobic
72 surfaces provides a fast and reproducible mean for the characterization of single molecules, purified from
73 the solution buffer and debris, which typically represent a serious problem in standard preparation for
74 laboratory analysis. The coupling of this preparation technique with Raman spectroscopy for the analysis
75 of the single suspended fiber is then a powerful combination for the study of any anisotropic fiber crystal,
76 where the structural characterization is not trivial by means of, e.g., x-ray crystallography or NMR.

77 **Materials and Methods**

78 **1. Super-hydrophobic surface fabrication**

79 Super-hydrophobicity emerges as a result of the patterning and functionalization of the surface,
80 whose combination provides very high contact angle displayed by these substrates ($>150^\circ$). In this case,
81 the pattern is a sequence of concentric circular arrays of pillars with radial pitch of $18\ \mu\text{m}$, each of the
82 pillars having a diameter of $6\ \mu\text{m}$ and height of 7 to $10\ \mu\text{m}$. To maintain a uniform surface coverage, the
83 number of pillars in each circular array was increased according to the recursive formula $N_i = N_{i-1} + 4$,
84 where N_i is the number of pillars of the i th circle. The patterning of the surface was obtained with a
85 combination of optical lithography and deep reactive ion etching (DRIE) technique. The original substrate
86 was a 4" standard Si $\langle 100 \rangle$ wafer. In the first step, the pillars pattern was defined by means of negative
87 optical lithography. The sample was then etched by means of DRIE (PlasmaLab System 100, Oxford
88 Instr.), with a final height of the pillars of about $10\ \mu\text{m}$. Finally, the functionalization of the surface with
89 hydrophobic material was achieved by the deposition of Perfluorodecyltrichlorosilane (FDTS) in a
90 Molecular Vapor Deposition System (MVD100E, Applied MST). FDTS in the presence of water reacts

91 with OH groups which are exposed at the surface of the oxidized Si, creating a monolayer of hydrophobic
92 fluoro-carbon chain molecules bound to the sample surface. The functionalizing FDTs was dispensed
93 together with water vapor in a vacuum pumped chamber of an MVD commercial system (MVD100E,
94 Applied MST).

95 **2. Lysozyme fibrils preparation**

96 Lysozyme amyloid fibrils were produced starting from a solution of HEWL powder (Sigma) [10
97 mg/ml] in MilliQ water based on [21]. Acidification of the solution to pH~2.0 was obtained by adding 9%
98 (v/v) of HCl [1N] in the solution. The mix was then immersed in a water bath at 60 °C for 120 h, until the
99 formation of suitably long amyloid fibrils, as verified by AFM. No purification of the amyloid fibers was
100 performed. The solution was then diluted ten times before depositing a 10 µl drop over the super-
101 hydrophobic substrate with a hypodermic syringe microneedle. The drop was, thereafter, dried at room
102 temperature and 50% humidity for 3 hours.

103

104 **3. Sample characterization by AFM and SEM**

105 The formation of amyloid fibrils at least 12 µm long (since this is the exact distance between the
106 pillars edge in radial direction) was monitored by AFM. A 30 µl drop was withdrawn every 12 hours and
107 spotted on a freshly cleaved mica sheet. It was let adsorb for 10 minutes and then rinsed thoroughly with
108 MilliQ water. Finally it was N₂ dried and ready for measurement. JPK Nanowizard III mounted on
109 inverted Olympus IX73 microscope was used for the measurement at relative humidity below 15%.
110 XSC11 AFM probes (MikroMasch, Nanoworld AG) with nominal resonance frequency of about 150 kHz
111 and nominal force constant of 7 N/m were run in tapping mode for the topography measurement.

112 After drop-casting the solution over the pillars top, we let it dry for 3 hours, then the deposition of the
113 fibers was verified by a Quanta 200 FEG Scanning Electron Microscope (FEI) at 5 kV beam voltage and

114 64 pA current. Before imaging, the sample was sputter coated with 2 nm of Iridium, to avoid charging
115 effects.

116

117 **4. Raman spectroscopy of lysozyme amyloid fibrils**

118 Laser micro-Raman spectroscopy was performed in confocal back scattering geometry by exciting the
119 sample with 532 nm linearly polarized laser wavelength (Coherent Compass Sapphire Laser, 75 mW) at 4
120 mW power on a WiTec Raman spectrometer (Alpha300 RA) with Andor CCD detector (DU970N) cooled
121 at -65 °C and a 100x objective (Zeiss, EC EPIPLAN NEOFLUAR, 0.9 NA). Measurements of the
122 lysozyme powder dissolved in MilliQ water and lysozyme fibrils in acidic solution were performed after
123 depositing the sample on a CaF₂ substrate to minimize signal background. The range of pillars height and
124 spacing useful for effective deposition is chosen to be bigger than the focal length and lateral dimension
125 of the focus of the objective. In this way the excitation volume is optimized to minimize the background
126 from the silicon substrate. For the measurement of the fiber anisotropy, the sample was oriented at 0°
127 (parallel) or 90° (perpendicular) with respect to the polarization of the excitation laser. Data analysis was
128 carried out as follows: at least ten measurements per sample were acquired, baseline subtracted in the
129 range 1100-1800 cm⁻¹ (by grade 5 polynomial fitting) and averaged. Afterwards all the spectra were
130 normalized to the Raman peak intensity at 1450 cm⁻¹ to compare data. Amide I spectral region was fitted
131 to a band described by a mixture of Lorentzian and Gaussian functions.

132 **Results and Discussion**

133 **1. Amyloid fibers on SHS**

134 Lysozyme amyloid fibrils suspended between pillars of a super-hydrophobic surface bear four major
135 characteristics that make their study by Raman spectroscopy unique: they are made of single/few isolated

136 molecules, free from the presence of a substrate, clean and not contaminated by debris, and highly
137 oriented along one given direction. This effect is obtained by the shear flow generated in the receding
138 meniscus of the drop which contains the reaction solution (including fibrils, oligomers and native
139 enzyme) and is evaporating over the micro-patterned SHS, and it was first demonstrated by De Angelis *et*
140 *al.* [1]. In this work, cylindrical micro-pillars arranged in a circular pattern were used. Pillars inter-
141 distance was 12 μm . The result of the dehydration process can be appreciated in Fig. 1: in Fig. 1a, four
142 different drying areas emerge as a result of the evaporation of the drop maintained in quasi-spherical
143 configuration by the SHS. The arrows indicate the drying direction, radial with respect to the center of the
144 drop. Along the drying pattern, an area with salt residues is evident on the right bottom corner, preceding
145 the area where the fibers are effectively extended hanging from the pillars edge (green line); this is then
146 followed by a zone with a larger and amorphous residue on the pillars top (starting at the red line)
147 followed by the edge of the collapsed drop (delimited by the blue line), where most of salt and fibers
148 monomers, not long enough to extend across the pillars, are accumulated. This qualitative result partially
149 supports the hypothesis that the SHS has a sieving effect on the drying solution. Fig. 1b-c are a sequence
150 of zoom-in pictures better elucidating the formation of the fiber bundle hanging out from the top of a
151 pillar in the part of Fig. 1a colored in green. The image of pillar top shows a darker area, where it is
152 clearly visible the semicircular drying periphery of the solution bearing radial direction, indicated by the
153 red arrows (Fig. 1b). A bundle of fibers is then formed, by coalescence of the fibers aided by the
154 centripetal drying process (Fig. 1c). Following this organization process the fibers align over the
155 interested area bridging consecutive pillars at a pillar-to-pillar distance of 12 μm . In the case shown in
156 Fig. 2c, the generated fiber was quite thick, while in other cases thinner fibers were extended. For
157 instance, in Fig. 2a, the fibers pinned at the edge of the pillar top have diameters of about 11 nm. This
158 value includes the 2 nm of Iridium coating, such that the actual thickness is slightly reduced. The
159 population distribution is then quite wide with fibers measuring also several tens of nanometers, mainly

160 depending on the distance from the center, such that a concentration effect is evident (the closer to the
161 drop center, the more concentrated the solution, the thicker the fiber).

162 Amyloid fibrils can aggregate in different fashion to the limit, however, of the minimum size of the
163 single amyloid fibril (called protofibril) at about 3 nm in diameter [22]. This effect of aggregation of
164 multiple fibrils due to concentration can be better explained after considering the metrology of the single
165 fibers as measured by the AFM. Fig. 2b is the AFM measurement topography performed at 120 h solution
166 incubation time, after a drop of the mixture was spotted and dried over a freshly cleaved mica sheet:
167 amyloid fibrils dimensions are 18 ± 3 nm of width, 3.4 ± 0.6 nm of height and a length of 19 ± 7 μ m. The
168 width originates actually from the profile measurement convolved with the tip shape, because it was not
169 possible to deconvolve reasonably the signal. Thus, the measured height was taken into consideration.
170 However it has to be pointed out that slight deformation of the fiber might depend on its adhesion to the
171 mica surface, on the shrinking of the molecule due to dehydration and on the effect of the probe tip
172 applying a force while performing the topography measurement. Moreover, the amyloid fibers have
173 different shapes themselves, where the protofibril can organize into fibrils having from cylindrical to
174 ribbon shape. Because the metrology of the amyloid fiber is beyond the scope of this work, we rely on the
175 literature that reports an average radius of the amyloid protofibril, rather common among the different
176 originating proteins, of about 3 nm, and we assume that the height of the fiber might be similar to this
177 value. Given these premises, we can thus state that the height average measured by the AFM on the fibrils
178 population is in agreement with the literature [23-25]. Amyloid fibers are composed of multiple
179 protofilaments that are intertwined together into a twisted ultrastructure with regular periodicity. In these
180 AFM measurements, two populations were found bearing each a twisting pitch of 76.5 ± 1.2 and $96.2 \pm$
181 1.7 nm. In conclusion, by comparing the AFM and the SEM measurement, we can argue that the thinnest
182 fibers extended on the pillars top could likely be 2 to 4 protofibrils intertwined together, given the SEM
183 diameter estimation.

184 **2. Raman measurement of amyloid fibers on SHS**

185 Lysozyme is an enzyme protein with antibacterial role and is commonly found in human secretions,
186 macrophage and egg white. Due to its immunity importance and availability, lysozyme 3D structure and
187 sequence have been among the first to be resolved. It has also been a model molecule in the early ages of
188 Raman spectroscopy applied to biomedicine due to the presence of the main aminoacids in its sequence
189 and of the S-S bond. Some major peaks characterize the laser Raman spectroscopy spectrum of a protein
190 in the visible excitation range, related to specific aminoacids (e.g. Phenylalanin $\sim 1003\text{ cm}^{-1}$; Tyrosine ~ 830 ,
191 850 cm^{-1} ; Tryptophan $\sim 763, 980, 1556\text{ cm}^{-1}$), to specific covalent bonds (S-S $\sim 510\text{ cm}^{-1}$; C-H (def.) ~ 1450
192 cm^{-1} and C-N $\sim 1130\text{ cm}^{-1}$) and to secondary structure (Amide III $\sim 1240\text{ cm}^{-1}$ and Amide I $\sim 1650\text{ cm}^{-1}$)
193 [15]. Our interest particularly focuses on the secondary structure fingerprints, given that a globular
194 protein, such as HEWL, will change its three dimensional structure by re-organizing the population
195 distribution of its secondary structures in the conversion to amyloid protein. The Amide I band (Raman
196 shift: $1620\text{-}1690\text{ cm}^{-1}$), mainly due to the C=O stretching of the peptide carbonyl group, will be used in
197 this work to describe the protein secondary structure in the visible excitation range. After the discovery
198 that every protein can turn into amyloid fibril, lysozyme was readily used to study the onset of these
199 pathological aggregates. The lysozyme powder is processed to obtain the amyloid fibrils, usually in
200 denaturing conditions and high temperature. The result is the destabilization of the protein tertiary
201 structure into an open globular feature which then rearranges into sheets of β -strands, since this type of
202 aggregation is energetically favored [26]. The structure of these assemblies has been somewhat
203 controversial; however, a general pattern common too many proteins is nowadays accepted, where the
204 core of the amyloid fiber is made up of β -sheets running perpendicular to the fiber and packed together by
205 side-chains interactions [27]. Aside from this core, other secondary structures such as α -helices, random
206 coils, loops, and other unordered structures are detected, depending on the sequence of the protein.
207 Besides, proteins showing the tendency to form amyloid fibrils always possess a susceptible aminoacid

208 sequence which is prone to alteration [28, 29]. It has to be pointed out that different preparation
209 conditions can give rise to amyloid polymorphs originating from the same protein, such that slight
210 variability exists in the fibrils structure and related analysis outcomes. All of these characteristic
211 secondary structures have been assigned specific Raman shifts in the Amide I band of the Raman
212 spectrum. As a rule of thumb, the Amide I band will be mostly populated by the β -sheet assigned peak,
213 thus having a center at $\sim 1672\text{ cm}^{-1}$ [16, 17, 30]. In this work, by suspending bundles made of few fibrils
214 between pillars, we expect a purified Raman signal unmistakably elucidating the secondary structure
215 typical of the amyloid molecule.

216 In Fig. 3 the Raman spectra in the range $1500\text{-}1800\text{ cm}^{-1}$ of the lysozyme in MilliQ water (Lys-MQ),
217 the lysozyme in the acidic solution after 120 h of incubation in hot bath (Lys-HCl) and the lysozyme fiber
218 bridged across pillars (Fibril_Pill) are plotted. Focusing on the Amide I band in the range $1620\text{-}1690\text{ cm}^{-1}$,
219 an upshift of the Amide I peak is readily recognizable when moving from globular protein to amyloid
220 fiber, where the peak apex moves from 1662 cm^{-1} to 1672 cm^{-1} . In particular, the curve Lys-MQ has a
221 peak at 1662 cm^{-1} ; the Lys-HCl has a peak centered at 1667 cm^{-1} ; the Fibril_Pill bears a remarkably
222 narrow peak at $\sim 1672\text{ cm}^{-1}$. This shift is already assigned in the literature to the transition from globular
223 protein to amyloid fibril, where the peak at 1672 cm^{-1} , might be assigned to the β -sheet predominant
224 component of the Amide I region [17]. The measurement in which the laser polarization is parallel to the
225 fiber is giving the better results in terms of the intensity of the peak attributed to the β -sheet secondary
226 structure, as compared to the spectrum obtained when the fiber is oriented perpendicularly to it: in fact,
227 the peak intensity increases nearly three-folds (Fig. 4). This result shows that the β -sheets are
228 preferentially oriented perpendicular to the fiber axis, as previously demonstrated by Sereda *et al.* [19], in
229 a polarized Raman study performed on insulin amyloid fibers aligned by the coffee ring stain effect.
230 However in their work, there was a need for previous purification of the fibers and the analysis was not
231 restricted to few molecules. Here we obtained the similar enhanced effect on the Raman spectrum, but

232 with the great advantage to provide with our method single bundles composed of few molecules that,
233 while purified, are oriented according to the patterned surface. In fact, the drop-casted solution is
234 populated by several species (globular enzymes, amyloid seeds, etc.), as it is shown in the AFM
235 topography image in Fig. 2b. These species are largely removed from the extended fibers after the drop
236 dries over the pillars, thanks to the sieving effect of the drying process on the SHS.

237 A deeper analysis of the results obtained from Raman spectroscopy on oriented fibers and electron
238 microscopy characterization of these samples is beyond the scope of this publication and will be reported
239 elsewhere in more detail.

240 **Conclusions**

241 In this work we have demonstrated a method to generate suspended amyloid fibrils highly
242 oriented in a pre-defined direction by the use of a SHS. This method provides fibrils free of any
243 contaminant and residual, decoupled from any supporting substrate, ready for further Raman or electron
244 microscopy analysis. The sample is obtained by the shear flow generated at the rim of a drop, casted over
245 a micro-patterned SHS. The receding meniscus combined with the shear flow and the pinning of the
246 fibrils on pillars edge is responsible for the alignment and purification of the amyloid fibrils. In particular,
247 we were able to suspend single/few amyloid fibrils of different diameters down to about 11 nm, as
248 verified by the SEM imaging. The micro-Raman analysis of the sample, as expected, revealed a
249 progressive upshift of the Amide I band main peak (from 1662 to 1667 to 1672 cm^{-1}) when the transition
250 from pure-lysozyme to lysozyme-amyloid-fibrils solution (mix of fibers and oligomers) and to lysozyme
251 amyloid fibrils suspended on pillars was considered. This testifies for the increased contribution of the β -
252 sheet secondary structure typical in amyloid fibrils. The distinctive anisotropic property of the amyloid
253 fiber was also revealed with an enhanced Raman signal peaked at $\sim 1672 \text{ cm}^{-1}$ after orienting the stretched
254 fibrils parallel to the polarization of the incident laser. In conclusion, we would remark that this method

255 could be considered as well suited to characterize any type of anisotropic fiber crystals. We foresee the
256 extension of the method for studying other protein molecules not only by Raman spectroscopy but also by
257 transmission electron microscopies, both in diffraction and direct imaging configuration and x-ray
258 crystallography. In fact, a single preparation technique, such as the dehydration on SHS device, provides
259 a sample available for characterization under three complementary techniques, simplifying and reducing
260 processing times. Future and alternative applications include, but are not limited to, the generation of
261 patterned surfaces when bio-compatible material is desirable, thanks not only to the availability of the
262 protein molecule (and eventual new synthesis of simpler sequences), but also to the possibility of
263 chemical-biological modification of the starting material.

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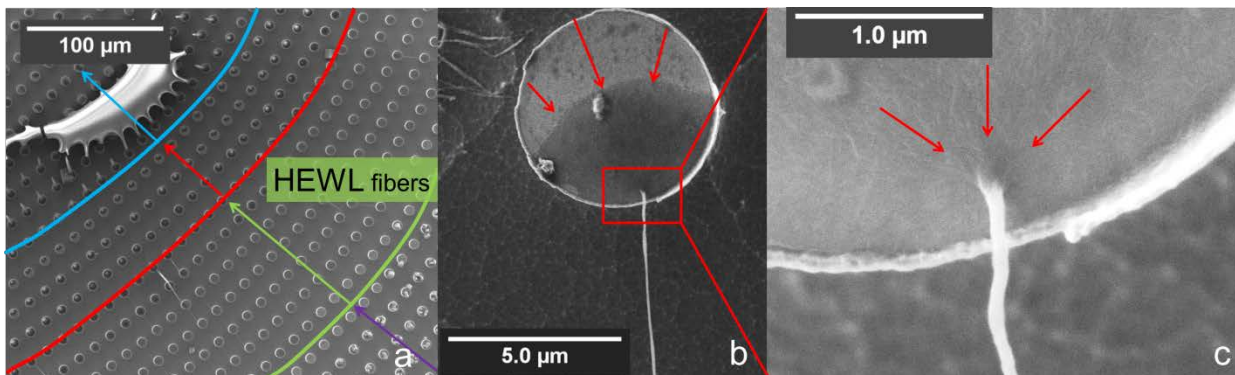
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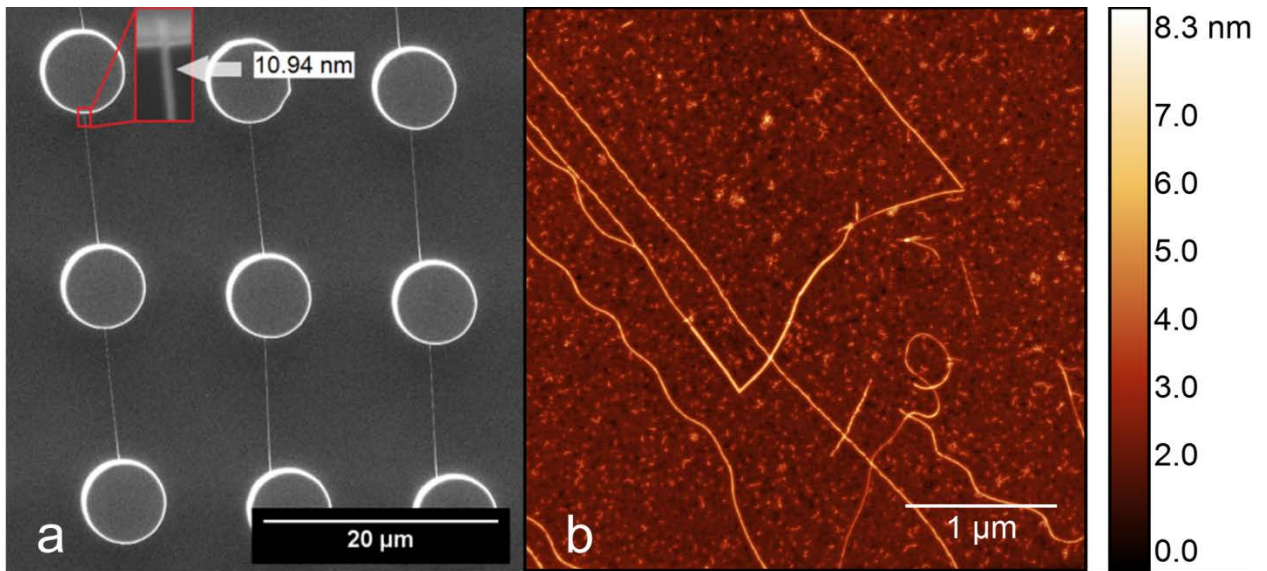
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350 **Figures and figure captions**

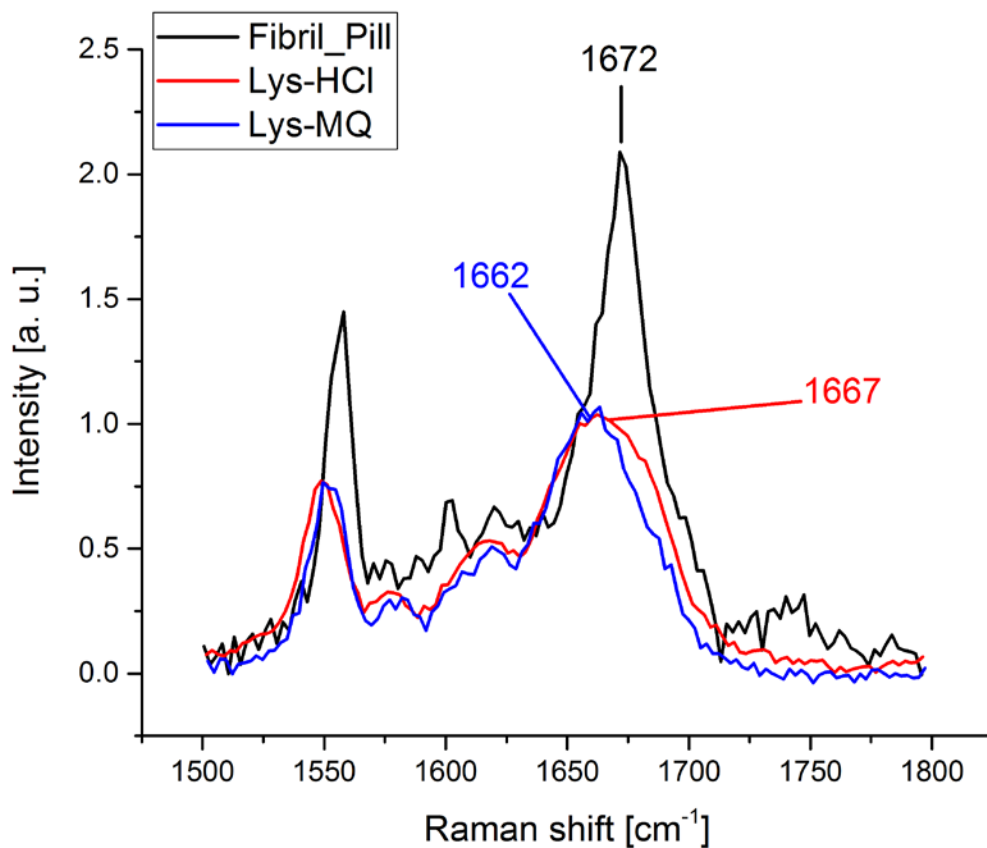


352 Fig. 1: SEM micrographs with cartoons elucidating the drying mechanism of the drop solution
353 spotted over the SHS. (a) A view over a large area which includes the side of the evaporated drop (top
354 left) and several drying areas evidenced by different marker colors. The green tag indicates the area where
355 the HEWL fibers are stretched over the pillars' top. The arrows indicate the drying direction; (b) a pillar
356 top with red arrows evidencing the drying shape and direction; (c) the area in the red square in (b) imaged
357 at higher magnification to show the coalescence of the fibers to one point (indicated by the red arrows)
358 where they are pinned out of the pillar top. The images were contrast and brightness corrected for better
359 visualization of the amyloid fibers.



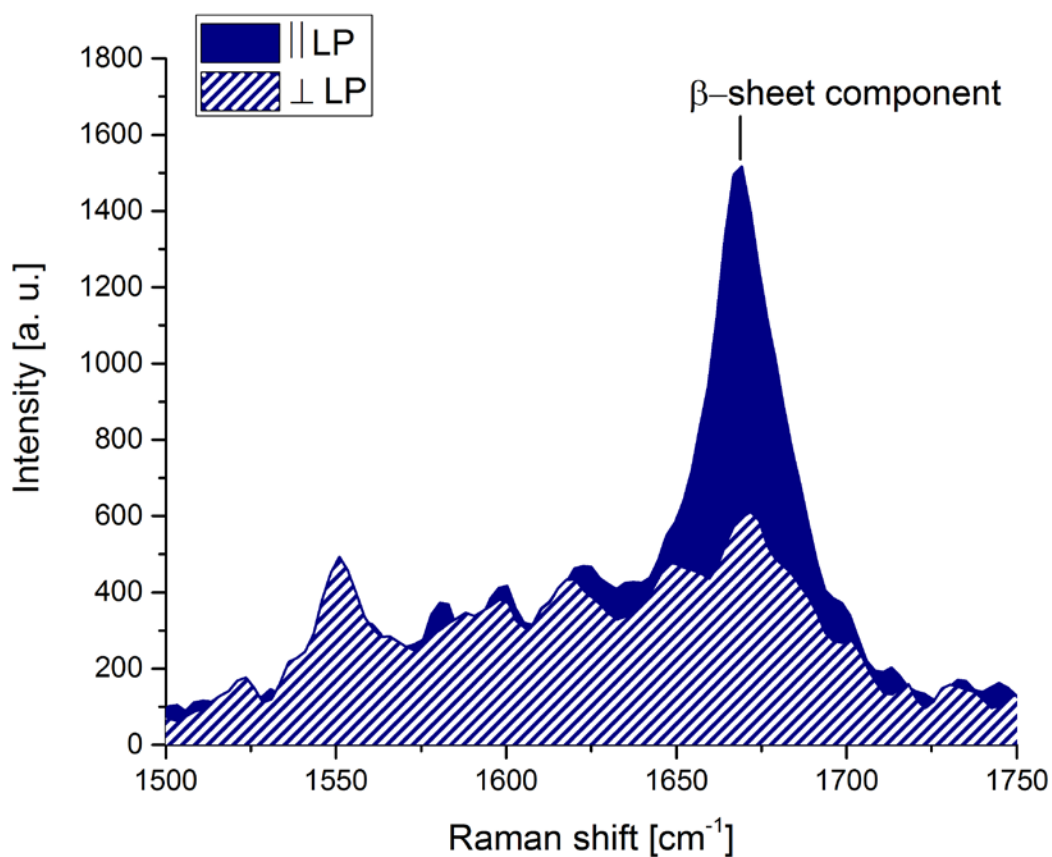
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361 Fig. 2: (a) SEM micrograph of the suspended fibers over super-hydrophobic pillars. Minimum
362 diameter measured was about 11 nm (inset). (b) AFM topography of lysozyme amyloid fibrils deposited
363 on mica sheet after 120 hours of incubation. The height of the fibrils is about 3 nm.



364

365 Fig. 3: Raman spectrum and peak assignment of the Amide I region for the HEWL in MilliQ water,
 366 named Lys-MQ, in blue, HEWL in acidic solution, named Lys-HCl, in red and the HEWL fibers bridging
 367 the pillars top, named Fibril_Pill, in black. Raman spectra were normalized to the $\sim 1450 \text{ cm}^{-1}$ peak. The
 368 assigned Raman peak was obtained by fitting Voigt function to a single peak in the Amide I band.



369

370 Fig. 4: Raman spectra of the amyloid fibers extended over the SHS with parallel (||LP, filled blue
 371 spectrum) and perpendicular (⊥LP, striped blue spectrum) orientation with respect to the exciting laser
 372 polarization (LP) in the region 1500 – 1800 cm⁻¹: the peak of the β-sheet component at 1672 cm⁻¹ is
 373 remarkably enhanced in the case of the laser polarization parallel to the fiber direction.