Local Structure in Spider Dragline Silk Investigated by Two-Dimensional Spin-Diffusion Nuclear Magnetic Resonance

J. Kümmerlen,* J. D. van Beek,† F. Vollrath,‡ and B. H. Meier*‡

NSR-Center for Molecular Structure, Design and Synthesis, Laboratory of Physical Chemistry, University of Nijmegen, Toernooiveld, 6525 ED Nijmegen, The Netherlands, and Department of Zoology, University of Aarhus, Universitetsparken B135, DK 8000 Aarhus C, Denmark

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ABSTRACT: The local structure of dragline silk from the spider Nephila madagascariensis is investigated by solid-state nuclear magnetic resonance. Two-dimensional (2D) spin-diffusion experiments show that the alanine-rich domains of the protein form β-sheet structures in agreement with one-dimensional NMR results from a different species of the genus Nephila (Simons, A.; Ray, E.; Jelinski, L. W. Macromolecules 1994, 27, 5335) but at variance with diffraction results. The microstructure of the glycine-rich domains is found to be ordered. The simplest model that explains the experimental findings is a 31-helical structure. Random coils, planar β-sheets, and α-helical conformations are not found in significant amounts in the glycine-rich domains. This observation may help to explain the extraordinary mechanical properties of this silk, because 31-helices can form interhelix hydrogen bonds.

(1) Introduction

Spider dragline silk is a remarkable biopolymer: its mechanical properties are a unique combination of high tensile strength and high elasticity. In contrast to the synthesis of man-made high-performance materials (e.g., steel or Kevlar) spider silk is synthesized at ambient temperature and pressure. The polypeptides that form the spider’s dragline are produced in a set of glands (the major ampullate) and channeled through a duct to the spigot. It seems that the solvent (water) is extracted in the duct, and that the silk goes into a liquid crystalline phase. At the end of the duct the silk is drawn through a valve where it apparently forms the fiber, now insoluble in all but the most aggressive solvents; remaining free water evaporates quickly in the air. The extraordinary properties of the spider’s dragline silk make it attractive to think about technically produced analogues with well-defined amino-acid sequences (see ref 10 and references therein). To do so, it is essential to understand the macroscopic properties of dragline silk in terms of its microscopic structure including the amino-acid sequence of the protein (the primary structure), the (local) folding of the strands (its secondary structure), the packing arrangement (crystalline or amorphous regions) in the solid fiber, and possible superstructures. The observations that water can act as a plasticizer for some of the silks produced by spiders and that wetting of the dragline silk can lead to a phenomenon called supercontraction are a clear indication that the knowledge of the amino-acid sequence alone is not sufficient to understand the properties of silk but that detailed knowledge of the secondary structure, the packing, and the interplay with resident and ambient water is required.

Solid-state NMR can probe the local structure in disordered systems where the lack of translational long-range order makes the application of diffraction technology difficult. The most useful NMR interaction for structural studies is the magnetic dipole interaction because of its simple and quantitative relationship to the atomic structure of the material. In the simplest case, e.g., for an isolated spin pair, the dipolar interaction manifests itself as a line splitting from which the internuclear distance can directly be evaluated. If the spectral resolution is insufficient, the investigation of the dynamics of the dipolar-induced polarization transfer between the nuclei can be used to determine the distance. Combined with two-dimensional (2D) NMR spectroscopy it can be used to determine the relative orientation of neighboring molecular segments in crystalline and amorphous materials. The resulting 2D spin-diffusion correlation maps directly relate to the local geometry if the purely spectral contributions are corrected for, or eliminated by, suitable pulse schemes. Here we apply these methods that use two-particle NMR interactions to dragline silk. Solid-state NMR methods that employ one particle interactions, in particular the chemical shift, have already been applied to dragline silk. In a recent study, Simmons et al. used the empirical dependence of the isotropic chemical shift of the Cα and Cβ resonances on conformation to obtain information about the secondary structure of alanine in the dragline. The spider silk most investigated is the dragline forcibly silked from the neotropical golden silk spider Nephila clavipes. The molecular weight of dragline was determined to be on the order of 200–350 kDa. The primary structure of the major ampullate gland protein (dragline) is still not known completely. The results of Lewis suggest that N. clavipes dragline silk is composed from two different proteins designated as spidroin I and spidroin II. In an independent study Mello et al. have confirmed the existence of spidroin I. The primary structure of spidroin I contains (Gly–Gly–X)m segments (where X = Gin, Ala, Tyr, Ser, or Leu and m = 3–6 if minor sequence errors are tolerated) and Aln segments (with n = 4–7). In a simplified view, silk may be looked at as a block copolymer with glycine-
rich and alanine-rich domains. Diffraction results clearly indicate that dragline silk is a heterogeneous material with "amorphous" and "crystalline" domains. It is generally accepted that the "crystalline" domains consist of protein segments that adopt an ordered β-sheet conformation. Controversy has, however, arisen whether these β-sheets are formed by the alanine-rich or the glycine-rich segments of the protein. The structure of the "amorphous" domains is largely unknown. It was suggested that they are formed from irregularly packed α-helical protein segments or random coils, while other authors find no evidence for such structures.

In this report, we present an investigation of dragline silk that uses the magnetic dipolar interaction. The silk was collected from the species Nephila madagascariensis whose relatively large body size facilitates the collection of comparatively large samples. Local ordering is found and interpreted for both the alanine- and glycine-rich domains of the protein.

(2) Materials and Methods

Silk was reeled from N. madagascariensis in the standard way at a speed of 26 cm/min. The spiders were kept at a low diet (of Tenebrio mealworms) supplemented with daily doses (starting a week before silking) of either [1-13C]alanine or [1-13C]glycine in an aqueous solution. The isotopic enrichment of the amino acids exceeded 98%, and in the resulting silk, 60% enrichment was found.

NMR spectra were obtained on Bruker MSL 300, MSL 400 and AM 500 spectrometers. Approximately 45 mg of sample was used. Radio-frequency-field strengths of typically 50 kHz have been used on carbon and proton channels. Cross-polarization contact times of 2 ms were used unless specifically differently in the text; the recycle delay between scans was set to 4 s.

The 2D 13C proton-driven polarization-transfer experiments were recorded on static samples using the pulse sequence CP-t1-π/2-τm-π/2-t2 with proton decoupling and quadrature detection in both dimensions. During the mixing time, no rf was applied. For some experiments, a one-dimensional version of the spin-diffusion experiment, CP-TOM-π/2-τm-π/2-t2, as suggested by Yang et al., was used; maximum τm have been set to be equal to a multiple of the rotor period by means of an external synchronization apparatus.

The solid-state NMR data will be interpreted by assuming that no significant transfer of the 13C label to other amino acids takes place by the metabolism of the spider. The solid-state spectra of silk containing 13C-labeled alanine and 13C-labeled glycine, presented below, make it immediately evident that no significant amounts of 13C are transferred to any aliphatic amino acids. The labeling of the silk corresponds to a chemical shift of 171.4 ppm. The observed isotopic chemical shifts are compiled in Table 1. To obtain the carboxylic chemical shifts of glycine and alanine and to get simplified spectra containing only carbonyl resonances, 13C-labeled alanine and 13C-labeled glycine (Gly)-labeled samples were prepared. This

3 Results and Discussion

3.1 One-Dimensional 13C CP/MAS Spectra. The natural abundance 13C CP/MAS spectrum of N. madagascariensis dragline silk is depicted in Figure 1 along with a tentative assignment of the individual resonances, obtained by comparison with the previously assigned spectrum from the silk produced by the silkworm Bombyx mori. As expected, the spectrum of the dragline silk is similar to the one obtained by Simmons et al. for dragline silk from N. clavipes. The aliphatic resonances of the three most abundant amino acids: glycine, alanine, and glutamine, are fairly well resolved. The Cα and Cβ resonances of serine are partially hidden due to overlap with the Cα resonance of alanine. The carboxylic resonances of all amino acids overlap and lead to a single slightly asymmetric resonance at approximately 172.3 ppm. The observed isotropic chemical shifts are compiled in Table 1. To obtain the carboxylic chemical shifts of glycine and alanine and to get simplified spectra containing only carboxylic resonances, [1-13C]alanine (Ala)- and [1-13C]glycine (Gly)-labeled samples were prepared. This
coincide, within experimental error, with the values published by Simmons et al. if we use the glycine C=O resonance with the larger chemical shift value. In addition we have obtained information about the C=O chemical shift tensors of alanine and glycine. The alanine isotropic chemical shifts of 172.5, 48.9, and 20.5 ppm for the C=O, Cα, and Cβ resonances, respectively, are clearly compatible with a β-sheet conformation. The α-helical conformation is unlikely to be a major component, while a β-helical conformation, for which a model study yielded 172.1, 48.7, and 17.4 ppm,49 could, in view of the distorted line shape of the resonance that is most dependent on the structural difference between these two forms (Cα), be a major constituent. For the glycine-rich parts of the polypeptide, structural conclusions from the chemical shifts are difficult to draw. Unfortunately, the resonance frequency of the Cα carbon of glycine does not allow a structural conclusion, in a fashion similar to that in the case of the alanine Cα resonance. Saito et al.46 have noted (by comparison of the 13C CP/MAS spectra of polyglycine in the form I and form II structure) that this frequency is rather insensitive to conformational changes. From the isotropic values as well as from the anisotropy and asymmetry of the chemical shift tensors of the carboxylic resonances, none of the possible conformations (α, β, 3i, ω) at the glycine position can be excluded.47,48

The signal intensities in a CP-MAS spectrum must be interpreted with care because they depend not only on the relative abundance of the amino acids but also on the cross-polarization efficiency. The 13C signal intensity as a function of the cross-polarization time is often characterized by an exponential buildup $I(t) = (1 - \exp(-t/T)}$ multiplied by a decaying exponential with a time constant $T_1$, that describes the decay of the sum-polarization in the rotating frame.50 The time constants $T_{1S}$ and $T_{1P}$ are both rather sensitive measures for dynamical processes with correlation times in the millisecond range. For a heterogeneous sample like silk, multi-exponential buildup and decay of these line intensities may be expected. For N. madagascariensis dragline silk, however, the experimental results are, within experimental error, described by a single exponential for buildup and decay of each resonance. The parameters describing the CP dynamics at room temperature are compiled in Table 2 for all resolved resonances. As expected, $T_{1S}$ is shortest for the carbons with directly bound protons. No large differences are observed for the proton $T_{1P}$ relaxation times of the different amino acids. These results show that the potential heterogeneity of the structure is not clearly reflected in the dynamics of the cross-polarization.

The observation of almost uniform CP dynamics with $T_{1P} > T_{1S}$ allows us to use the relative intensities of the individual resonances in the CP/MAS spectrum with 2 ms contact time as a rather faithful measure for the amino acid composition. Through spectral deconvolution of the aliphatic resonances in the 13C CP/MAS spectrum of N. madagascariensis dragline silk (see

Figure 2. 13C CP/MAS spectra of (a) 1-13C glycine-labeled N. madagascariensis dragline silk and (b) 1-13C alanine-labeled dragline silk. A total of 240 transients were acquired at a carbon resonance frequency of 75.47 MHz and a MAS-spinning frequency of 2 kHz. (c) Expanded centerband of the carboxylic resonances of a wet 1-13C glycine-labeled spider dragline silk sample. The expanded centerband (marked with an arrow) of spectrum (a) is given in (d) for comparison.
Figure 1) an estimate for the amino acid composition can be given. The relative abundance of the amino acid was found to be approximately 45%, 30%, 10%, 8%, and 4% for glycine, alanine, glutamine, serine, and tyrosine, respectively. A comparison with the amino-acid composition data of *N. clavipes* dragline silk reveals no significant differences between these two members of the genus *Nephila*.

(3.2) Two-Dimensional CSA-Tensor Correlation through Spin-Diffusion. To obtain insight into the local structure in the alanine- and glycine-rich segments, two-dimensional proton-driven spin-diffusion experiments in static samples were performed. These experiments yield the relative orientation of chemical-shift tensors of nuclei in spatial proximity. "Spatial proximity" is defined such that polarization transfer between the nuclei can take place within the mixing time of the 2D experiment. In general, the spin-diffusion rate constant $W_{ij}$ between two spins $i$ and $j$ depends not only on their internuclear distance $r_{ij}$ but also on the angle $\theta_{ij}$ between the internuclear vector and the external magnetic field direction and, through the intensity of the zero-quantum spectrum at frequency zero $F_{ij}(0)$, on the chemical shift difference of the two spins $i$ and $j$:

$$W_{ij} = \frac{\pi \hbar^2}{2 (4\pi)^2} \sum_{x,y,z} (x^2 \cos^2 \theta_{ij} - 1)^2 F_{ij}(0)$$  (1)

$F_{ij}(0)$ can be estimated from the experimental zero-quantum spectrum. Averaged over the chemical shift differences present in the sample, we have found a value of $F(0) \approx 5 \times 10^{-4}$ s and can therefore estimate that distances as large as 6–7 Å can be bridged by spin diffusion for a mixing time of 10 s.

As discussed in detail by Robyr et al., 28 2D spin-diffusion spectra can most easily be interpreted if the polarization transfer progresses to completion and a quasi-equilibrium state is reached where the spectrum is independent of the mixing time. In a completely ordered system, quasi-equilibrium is obtained if the slowest component of the polarization transfer establishes an equilibrium between neighboring symmetry-equivalent units. For a disordered system, quasi-equilibrium is reached if the spin diffusion has proceeded farther than the correlation length of the local order. If no quasi-equilibrium spectrum can be obtained, the interpretation of the data becomes quite involved and is best done in the initial-rate regime. For our silk samples, such an analysis is not possible because signal-to-noise considerations prevented us from obtaining initial-rate spectra. It is therefore important to work in a regime where the quasi-equilibrium assumption is a reasonable one. Obviously, a heterogeneous, partially disordered material like spider silk will never allow us to obtain true quasi-equilibrium spectra.

The analysis of the quasi-equilibrium spectra yields the Euler angles between the principal axis systems of the CSA tensors of the involved nuclei. To obtain information about the molecular fragments involved, the orientation in the CSA tensor in the molecular coordinate system must be known. From the study of model compounds, it has been found that, consistently, the orientation of the $^{13}$C CSA tensor in peptides is as follows: $\delta_{33}/\delta_{22}$ is perpendicular to the plane spanned by the O=O=N fragment, and the principal axis associated with the intermediate component of the CSA tensor $\delta_{33}$ is approximately the same for both samples. The orientation of the principal axis associated with the most shielded CSA tensor $\delta_{22}$ is almost colinear with the C=C vector. Experimental determinations of the angle between show the principal axis associated with $\delta_{22}$, and the bond direction falls, for proteins, into the range 0–12°.

The proton-driven 2D spin-diffusion spectra of $[1-^{13}$C]glycine- and $[1-^{13}$C]alanine-labeled silk at $T = 150 K$ and 10 s mixing time are given in Figures 3 and 4, respectively. The qualitative difference between the two spectra is quite striking: the spectrum of $[1-^{13}$C]glycine (Figure 3) shows a rather broad exchange pattern, while virtually no off-diagonal intensity is observed for the alanine carboxylic carbons (Figure 4). The degree of labeling is approximately the same for both samples. This can easily be seen from a comparison of the intensity of the labeled C=O resonances with the natural abundance signals in Figure 2a,b.

The absence of cross-peaks in alanine indicates that either the spin diffusion between alanines is very slow because they are well separated in space or the polypeptide regions are highly ordered with all three principal axes of carboxylic CSA tensors being either parallel or antiparallel. The zero-quantum line shape can be assumed to be of the same order of magnitude for both amino acids. 17 If we assume that the primary structure of *N. madagascariensis* silk is similar to that of *N. clavipes*, then the first explanation is highly improbable because its primary structure consists of alanine rich segments. We conclude therefore that close-by $[1-^{13}$C]alanine CSA tensors have approximately colinear principal axis systems.

**Figure 3.** (a) proton-driven spin-diffusion spectrum of $[1-^{13}$C]glycine-labeled *N. madagascariensis* dragline silk at $T = 150 K$. A mixing time of 10 s was used. The spectrum was acquired with 128 transients per data point in $t_1$; 96 spectra have been recorded in the $F_1$ domain. The data matrix of 96 × 128 points was zero-filled to 256 × 256. An exponential filter function was used prior to the double Fourier transformation. The resulting 2D spectra were symmetrized about the diagonal to enhance signal to noise. (b) Contour plot of the same data. The total signal intensity of all spectra is normalized to 1000 (for a digital resolution of 1.76 points per ppm) and equidistant contour levels are set at $\pm 0.025, 0.05, 0.075, ....$
A structure where most alanines are incorporated into an ordered β-sheet structure explains these experimental findings. Taking into account the experimental line shape, we find that the spectrum of Figure 4 confines all three Euler angles to be 0 ± 10 or 180 ± 10°.

The spectrum from the glycine-labeled samples exhibits a nonisotropic exchange pattern with a two-dimensional line shape strongly deviating from the 2D product of the one-dimensional line shape function given in Figure 5a. Therefore, the local packing must be ordered. Because not much is known about the interchain packing and the resulting interchain distances, we shall consider models with both intra- and interchain interactions to explain the experimental spectra.

Let us first consider intrachain spin diffusion only. We assume that the spin diffusion has equalized the polarization over many (>10) carbon atoms within one chain. This is a reasonable assumption because the rate constant obtained from eq 1 is much faster than the inverse mixing time. This is true not only for 13C in amino acids adjacent in the primary sequence (where an average rate constant of about 5 s⁻¹ is predicted) but also for 13C in amino acids separated by two in the primary sequence. A random coil arrangement would then lead to a spectrum closely resembling the amorphous spectrum of Figure 5a and can be excluded. β-sheet structures can also be excluded because they would lead again to an almost diagonal 2D spectrum. We shall, in the following, test the compatibility of two other plausible structures, the α-helix and 3i-helix, with the experimental data of Figure 3.

For an α-helical structure (see Figure 6) all C=O bond directions are approximately parallel to the helical chain axis. Correspondingly, the principal axis directions associated with the Cartesian α₁ components of all 13C CSA tensors coincide approximately with the helix axis. The relative tensor orientation between two neighboring carboxylic CSA tensors is therefore described by a rotation around the y axis of the principal axis system of the CSA tensor, corresponding to a set of Euler angles (0°, 100°, 0°). From a simulated poly(Gly-Gly-X) α-helix (with X = Ala), obtained with the program Quanta, Euler angles of (65.4°, 97.7°, 346°) are extracted. The quasi-equilibrium spectrum corresponds to a two-site exchange spectrum between all 13C in an isolated helix. The spectrum clearly disagrees with the experimental data of Figure 3.

Next we consider a 3i-helical arrangement (see Figure 6). Such a structure is found, for example, for the form II crystal structures of poly-Gly, poly(t-Ala—Gly—Gly), poly(t-Ala—Gly—Gly), and poly(Gly—β-Ala). In a (Gly—Gly—X)m-3i-helical structure only two symmetrically inequivalent [1-13C]Gly sites exist and the quasi-equilibrium spectrum corresponds to a two-site exchange spectrum.

From a model (Gly—Gly—Ala)ₙ polypeptide produced by the program Quanta the Euler angles that relate to the experimental spectrum of Figure 5a are extracted. The quasi-equilibrium spectrum of a (Gly—Gly—X)₁ α-helix (see Figure 3) is found, for example, for the form II crystal structures of poly-Gly, poly(t-Ala—Gly—Gly), poly(t-Ala—Gly—Gly), and poly(Gly—β-Ala). In a (Gly—Gly—X)m-3i-helical structure only two symmetrically inequivalent [1-13C]Gly sites exist and the quasi-equilibrium spectrum corresponds to a two-site exchange spectrum.

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the O=C—N fragments of each glycine residue were found to be \( \alpha = 114^\circ \), \( \beta = 78^\circ \), and \( \gamma = 346^\circ \). A simulated quasi-equilibrium 2D spin-diffusion spectrum assuming spin exchange between glycine sites related by these Euler angles is depicted in Figure 5c. For all simulations, the GAMMA programming environment was used.\(^6\) The total signal intensity of all spectra in this paper is normalized by the same procedure described in the legend of Figure 3. Therefore the contour levels in all figures can directly be compared. Qualitatively, the simulated spectrum resembles the experimental spectrum of Figure 3.

To improve the description of the experimental spectrum, we fit the data in Figure 3 by a quasi-equilibrium spin-diffusion spectrum between two magnetically non-equivalent sites. Starting from the spectrum in Figure 5c the Euler angles are optimized by means of a nonlinear least mean squares fit.\(^6\) Free parameters of the fit are the set of Euler angles (\( \alpha, \beta, \gamma \)) and the line width. We assume the line shape to be Lorentzian with a constant full width at half-height over the 2D spectrum. Furthermore, we allowed for a superposed signal with an isotropic powder exchange spectrum to account for spin diffusion between chains. The fraction of the isotropic exchange spectrum yields a fifth parameter for the fit. The chemical shift parameters were set to the values determined from the 1D spectra. The best fit, shown in Figure 7b, reproduces the off-diagonal features of the experimental 2D spin-diffusion spectrum (Figure 7a) remarkably well. The fitted Euler angles are \( \alpha = 101 \pm 13^\circ \), \( \beta = 62 \pm 14^\circ \), and \( \gamma = 333 \pm 10^\circ \) (error estimated at 90% confidential limit) and an admixture of 14 ± 5% signal intensity from an isotropic exchange pattern. The absolute value of the difference between the experimental spectrum and the best fit amounts to 12% of the integrated intensity of the experimental spectrum. If we do not allow for an isotropic powder pattern as part of the model, the difference between the calculated and experimental spectrum rises to 23%. The difference spectrum is depicted in Figure 7c and gives no indication that the quasi-equilibrium spectrum is distorted through the influence of the factor \( F_\gamma(0) \) which would lead to a monotonic decrease of the cross-peak intensity with increasing distance from the main diagonal of the spectrum. The main difference intensity appears along the diagonal itself, particularly at the sharp features of the tensor. The pronounced intensity at the sharp features indicates non-Lorentzian line shapes and the remaining intensity over the entire diagonal implies that complete polarization transfer between all glycines in the sample has not been established. The remaining diagonal signal may be attributed to glycines that are located outside the (Gly—Gly—X)* segments\(^37\) and therefore more remote from the next Gly neighbor. This, as well as the relatively small intensity of the diagonal ridge and the absence of a decay of the experimental cross-peak intensity (compared to the simulation) perpendicular to the diagonal of the spectrum, is a good \textit{a posteriori} justification of the assumption of quasi-equilibrium conditions. We have repeated the fit with starting values that correspond to an \( \alpha \)-helical conformation and have found that it converges to the same 3\( \beta \)-helical structure described above.

The fitted Euler angles deviate by about 15° from the idealized (Gly—Gly—X)* structure. These deviations can be explained by structural distortions and/or the deviation of the principal-axis directions of the CSA tensor.

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Figure 6. (a) Conformation of an \( \alpha \)-helix. (b) Conformation of a \( 3_1 \)-helix. The C=O bond direction (indicated by filled bond symbols) is approximately along the helical axis direction in the \( \alpha \)-helix and almost perpendicular to it in the \( 3_1 \)-helix. In contrast to the \( \alpha \)-helical structures which are stabilized by intrachain hydrogen bonds, \( 3_1 \)-helices are stabilized by interchain hydrogen bonds. The figures were created with the program MOLSCRIPT.\(^6\)

Figure 7. (a) Experimental spectrum \( ^1\)\( ^3\)C glycine (see also Figure 3), replotted for direct comparison with (b)-(e). (b) Best fit of the experimental spectrum by a two-site exchange model with an amorphous background, as described in the text. The conformation represents, within reasonable expectations (see text), a \( 3_1 \)-helical structure. (c) Difference between the experimental spectrum (a) and fit (b). (d) Best fit by a superposition of an amorphous and a diagonal spectrum (see text). (e) Difference between (a) and (d). Contour levels are chosen as described in the legend of Figure 3.
from the idealized values. Therefore, the spin-diffusion data are in good agreement with a 3_1-helical structure formed by the (Gly–Gly–X)_n segments of the protein.

The overall agreement between the certainly over-simplified model and the experimental spectrum may, considering the heterogeneous nature of the dragline silk as well as the possible experimental imperfections, be fortuitously good. We shall now shortly investigate possible other models that explain the experimental spectrum but assume significant interchain spin diffusion. Obviously, the random-chain model spectra of Figure 5a is not changed by taking into account interchain diffusion. All the other model spectra also approach the spectrum of Figure 5a if interchain diffusion between completely disordered chains ("spaghetti model") is assumed. Because this prediction is in contradiction with the experiment, we consider next uniaxially ordered bundles of chains. For such a model bundle, quasi-equilibrium spectra for 2_1-helices, a-helices, and 3_1-helices, respectively, are shown in Figure 5a–f. No satisfactory agreement with the experimental data is found.

As a further step, we may consider more complex models which describe heterogeneous situations. We therefore consider an amorphous spectrum (from a random-coil domain) superposed with a diagonal spectrum from isolated glycines. Using the line width and the relative amount of diagonal spectrum as free parameters, a least squares fit to the experimental spectrum was performed. The best fit yields 31 ± 4% isolated glycine, the resulting spectrum is shown in Figure 7d, and the difference between experiment and fit is given in Figure 7e. The agreement is not satisfactory and leaves 25% of the spectral intensity unexplained. Furthermore, we have observed that the carboxylic signal of a uniaxially oriented bundle of fibers shows a significant deviation from the powder pattern obtained from an unoriented sample. On the basis of these two arguments, we exclude amorphous random coils as a major component of the structure. A large number of further heterogeneous structures could be considered. We shall refrain, in the present context, from doing so because the simple 3_1-helical model shows already a good agreement with the experiment.

It should be pointed out that the off-diagonal peaks in the spectrum of Figure 3 could also be caused by chemical exchange instead of spin exchange or by the simultaneous action of both processes. Usually, the temperature dependence of the cross-peak intensity allows for a distinction of the two effects: the spin diffusion is largely temperature independent while an Arrhenius-type activation is expected for the chemical exchange. While the full information available by spin-diffusion spectroscopy is contained only in two-dimensional static spectra, selective 1D experiments under magic-angle spinning conditions can be used to extract partial information using shorter total measurement times, although a direct comparison between spin-diffusion rate constants in static and rotating samples is difficult. The one-dimensional analog of the exchange experiment proposed by Yang et al. was chosen to follow the polarization-transfer process as a function of temperature and to extend the spin-diffusion measurements to longer mixing times. The one-dimensional spectrum obtained in this experiment does not contain the entire spinning-sideband manifold but only the centerband if no exchange (spin diffusion or chemical exchange) takes place in the mixing time of the experiment. If, however, one of the processes is active, the special magnetization state prepared by the TOSS sequence is disturbed and spinning sidebands are reintroduced into the spectrum. A series of spectra recorded with different mixing times allows us therefore to extract information similar to that from the 2D spin-diffusion spectrum with the difference that only a limited number of sampling points are available. Figure 8 plots the TOSS centerband and sideband intensities in 13C spectra of the [1-13C]glycine-labeled sample obtained with the 1D experiment as a function of mixing time; the sample temperature was 220 K. It can easily be seen that on an increase of the mixing time the intensity of the centerband decreases exponentially (with a time constant of 6.3 ± 0.5 s) and spinning sideband intensity is reintroduced monotonically. For the long mixing times used in the experiments magnetization losses due to T1 relaxation are considerable and the total intensities of the spectra at different mixing times have been normalized in Figure 8. These data are a further indication that the assumption of an established quasi-equilibrium after 10 s of mixing, used for the interpretation of the 2D spectra, is a reasonable one.

Figure 9 depicts the resulting 1D TOSS spectra of the [1-13C]glycine-labeled sample using a mixing time of 10 s for temperatures of 220 and 293 K. The temperature dependence of the spectra and, therefore, of the spin-diffusion rate constant is negligible and leads us to the conclusion that contributions from chemical exchange to the spectrum of Figure 3 (recorded at an even lower temperature) can be excluded.

(4) Conclusions

We have shown that the secondary structures of the alanine-rich and glycine-rich segments of spider dragline silk can be characterized by means of proton-driven 13C 2D spin-diffusion experiments. Our results are in good agreement with the following structural model for N. madagascariensis dragline: The polyalanine seg-
ments adopt a highly ordered \( \beta \)-sheet structure whereas the glycine-rich segments form \( \alpha \)-helical structures. Considering the relatively short length of these segments and the resulting constraints on the packing scheme of the solid, these idealized structures are astonishingly well represented in the experimental spectra. Obviously, there may be more complicated structural models for the glycine-rich domains that can also explain the experimental data. \( \beta \)-Sheets and \( \alpha \)-helices can, however, be excluded on the basis of our data.

It is obvious that an explanation of the mechanical properties of spider dragline does not only demand the knowledge of the secondary structure of the different domains in the protein. The local packing, i.e. the size of the \( \beta \)-sheet structures and the packing of the \( \alpha \)-helices, is required for a complete understanding of the macroscopic properties. Nevertheless, one might speculate that the extremely strong mechanical properties of silk may be related to the presence of \( \alpha \)-helical structures because they are able to form inter-helix hydrogen bonds cross-linking the helices. A detailed analysis of the packing schemes of the helices among themselves requires multiple labeled samples and experiments with macroscopically aligned fiber samples. Work along these lines is in progress in our laboratory.

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(55) Quanta4.0 is a commercial Program from Molecular Simulations Inc., Burlington, MA.
(61) Program MINUIT was used. This is part of the PACKLIB program package. It was obtained under the conditions of the CERN Program Library/Division CN, CERN 1211 Geneva, Switzerland.