

Spectral editing of two-dimensional magic-angle-spinning solid-state NMR spectra for protein resonance assignment and structure determination

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Abstract Several techniques for spectral editing of 2D ^{13}C – ^{13}C correlation NMR of proteins are introduced. They greatly reduce the spectral overlap for five common amino acid types, thus simplifying spectral assignment and conformational analysis. The carboxyl (COO) signals of glutamate and aspartate are selected by suppressing the overlapping amide N–CO peaks through ^{13}C – ^{15}N dipolar dephasing. The sidechain methine (CH) signals of valine, leucine, and isoleucine are separated from the overlapping methylene (CH_2) signals of long-chain amino acids using a multiple-quantum dipolar transfer technique. Both the COO and CH selection methods take advantage of improved dipolar dephasing by asymmetric rotational-echo double resonance (REDOR), where every other π -pulse is shifted from the center of a rotor period t_r by about $0.15 t_r$. This asymmetry produces a deeper minimum in the REDOR dephasing curve and enables complete suppression of the undesired signals of immobile segments. Residual signals of mobile sidechains are positively identified by dynamics editing using recoupled ^{13}C – ^1H dipolar dephasing. In all three experiments, the signals of carbons within a three-bond distance from the selected carbons are detected in the second spectral dimension via ^{13}C spin exchange. The efficiencies of these spectral editing techniques range from 60 % for the COO and dynamic selection experiments to 25 % for the CH selection experiment, and are demonstrated on well-characterized model proteins GB1 and ubiquitin.

Keywords Spectral editing · REDOR · CH selection · Protein secondary structure

Introduction

Recent progress in magic-angle-spinning (MAS) solid-state NMR (SSNMR) spectroscopy has made it possible to determine the three-dimensional structure of insoluble proteins such as membrane proteins (Hong et al. 2012; Luca et al. 2003) and amyloid fibrils (Tycko 2011). Central to this structure determination is the resolution of protein ^{13}C and ^{15}N peaks in multidimensional MAS correlation spectra and the assignment of these peaks to specific amino acid residues (Luca et al. 2003). Within the repertoire of multidimensional NMR correlation experiments for resonance assignment, the 2D ^{13}C – ^{13}C correlation experiment based on ^1H -driven ^{13}C spin diffusion (PDSD) is especially widely used, since it is simple to perform and allows the assessment of protein homogeneity, sensitivity, and secondary structure. The 2D ^{13}C PDSD experiment is also the basis for distance extraction, since cross-peak intensities measured as a function of mixing time provide semi-quantitative distance information (Castellani et al. 2002).

While small proteins (<100 residues) with microcrystalline structural order can give completely resolved 2D PDSD spectra (Castellani et al. 2002; Franks et al. 2005a, b; Igumenova et al. 2004; Pauli et al. 2001), non-crystalline proteins usually exhibit ^{13}C linewidths of greater than 1 ppm (Su and Hong 2011). As a result, their 2D ^{13}C correlation spectra often show significant resonance overlap. While 3D and 4D experiments can be used to reduce resonance overlap, the high dimensionality severely reduces the sensitivity compared to 2D experiments; moreover, analyses of 3D and 4D spectra are much more time-consuming. For proteins

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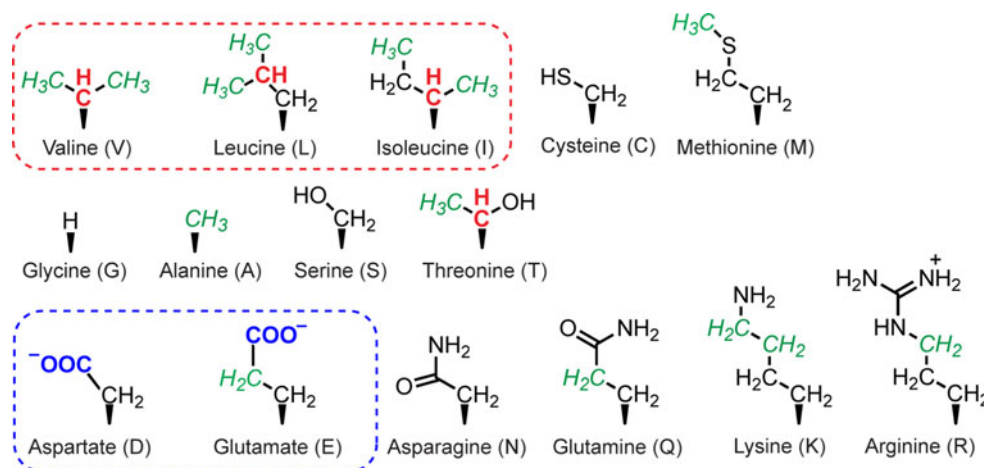
with linewidths exceeding 1 ppm, even 3D spectra often do not fully resolve all signals. Therefore, there are significant incentives to develop simple and efficient strategies to increase the information content of 2D NMR spectra and enable resonance assignment in the presence of line broadening.

Spectral editing has long been used to simplify and assign the SSNMR spectra of unlabeled organic compounds, whose sensitivity is too low for multidimensional ^{13}C correlation spectroscopy. 1D ^{13}C MAS spectra have been edited based on the number of attached protons per carbon (CH_n , $n = 0, 1, 2$) (De Vita and Frydman 2001; Frey and Opella 1984; Lesage et al. 1998; Mao and Schmidt-Rohr 2005; Sakellariou et al. 2001; Schmidt-Rohr and Mao 2002a, b; Wu et al. 1994; Wu and Zilm 1993), bonding to a heteronucleus such as nitrogen (Fang and Schmidt-Rohr 2009; Schmidt-Rohr and Mao 2002a, b), chemical shift anisotropy (Mao and Schmidt-Rohr 2004), and molecular motion, to give selective 1D ^{13}C spectra of different chemical functional groups. Based on spectral editing, the chemical composition of a variety of organic materials has been determined (Fang and Schmidt-Rohr 2009; Keeler and Maciel 2000; Mao et al. 2007a, b; Schmidt-Rohr et al. 2004). In comparison, spectral editing of 2D and 3D correlation spectra of ^{13}C , ^{15}N -enriched proteins has so far been restricted to the detection of the sidechain signals of several types of amino acids (Jehle et al. 2006a, b; Reggie et al. 2011). The potential for protein spectral editing is illustrated in Fig. 1: various amino acids with distinct functional groups exhibit similar chemical shifts. For instance, the ^{13}C chemical shifts of several methylene (CH_2) groups in Lys, Asp, and Arg are similar to the ^{13}C chemical shifts of the methine (CH) groups in Val, Leu and Ile, while backbone and sidechain carbonyl groups bonded to nitrogen share similar chemical shifts as the sidechain COO groups of Asp and Glu.

In this work, we demonstrate three spectral editing techniques that reduce such resonance overlap and facilitate assignment of 2D ^{13}C PDS spectra of ^{13}C , ^{15}N -labeled proteins. First, ^{15}N - ^{13}C dipolar dephasing is used to select the signals of the COO -containing Glu and Asp residues. Carboxyl sidechains are important in hydrogen bonding and ionic interactions such as those between proteins and biomineral surfaces (Goobes et al. 2006; Hoang et al. 2003). Despite their functional importance, an examination of the BMRB protein database shows that less than 5 % of Glu and Asp residues have assigned sidechain COO chemical shifts, indicating that standard multidimensional NMR techniques do not easily provide this information. Second, we select CH signals by suppressing the CH_2 signals using multiple-quantum recoupled dipolar dephasing (Mao and Schmidt-Rohr 2005). This method reveals the CH signals of Ile, Val, and Leu, which are found in fairly crowded but mutually exclusive regions of the 2D ^{13}C PDS spectra. Third, we demonstrate the selection of mobile residues' signals by ^{13}C - ^1H dipolar recoupled gated ^1H decoupling. Protein dynamics often contain important information about protein function (Hong 2006; Hong et al. 2012; Huster et al. 2001; Lorieau et al. 2008). While scalar-coupling based polarization transfer schemes can detect nearly isotropically mobile residues in proteins (Etzkorn et al. 2007), residues with anisotropic dynamics do not survive these solution-NMR-derived pulse sequences well, thus dipolar-based experiments to selectively detect dynamic residues are useful.

Common to these techniques is the novel use of asymmetrically positioned 180° pulses in the REDOR sequence (Gullion and Schaefer 1989), which enables complete dephasing of undesired signals. We demonstrate this asymmetric REDOR technique using numerical simulations and experiments. The spectral editing experiments presented here are tailored to MAS frequencies of ~ 10 kHz, which are higher than the typical MAS rates used for spectral editing of

Fig. 1 Sidechain structures of aliphatic amino acid residues whose functional groups are spectrally edited here. *Red* Sidechain CH groups that are selected by the dipolar DEPT experiment. *Green italics* Dynamic groups that may be selected by the gated ^1H decoupling experiment. *Blue* Carboxyl groups that are selected by the ^{13}C - ^{15}N REDOR experiment



unlabeled organic compounds (4–6 kHz), in order to better suppress ^{13}C – ^{13}C dipolar couplings and obtain higher spectral resolution for ^{13}C -labeled proteins.

Experimental methods

Sample preparation

Uniformly ^{13}C , ^{15}N -labeled immunoglobulin-binding domain $\beta 1$ of the streptococcal protein G (GB1) was generously provided by Professor Chad Rienstra. Microcrystalline order was achieved by precipitating the protein using polyethylene glycol (PEG) (Franks et al. 2005a, b; Gallagher et al. 1994). The 56-residue GB1 has sufficient amino acid diversity for demonstrating COO, CH, and dynamics editing of the 2D spectra, and has excellent spectral resolution to enable complete resonance assignment. The sample was packed into a 2.5-mm MAS rotor by centrifugation. Bulk mother liquor was removed from the rotor by pipette. The protein was kept above 0 °C in all NMR experiments to avoid perturbation of the microcrystallinity by ice formation. Uniformly ^{13}C , ^{15}N -labeled ubiquitin in lyophilized powder form was obtained from Sigma-Aldrich and used without recrystallization.

Solid-state NMR experiments

SSNMR experiments on GB1 were carried out on a wide-bore Bruker AVANCE II-600 spectrometer (14.1 T) while ubiquitin spectra were measured on a wide-bore Bruker 400 MHz NMR spectrometer. 2.5-mm triple-resonance MAS probes were used and the samples were spun at a frequency of 10 kHz. Typical radio-frequency (rf) field strengths were 62.5 kHz for ^{13}C , 50 kHz for ^{15}N and 100 kHz for ^1H . ^{13}C chemical shifts were externally referenced to the carbonyl peak of α -Gly at 176.465 ppm, which is consistent with the adamantane CH_2 chemical shift of 38.48 ppm (Morcombe and Zilm 2003), both reported on the neat TMS scale. On this scale, the measured ^{13}C chemical shifts of GB1 are smaller by 2.4 ppm than the DSS-based values reported by Rienstra and coworkers (Franks et al. 2005a, b; Morcombe and Zilm 2003). This difference is verified to match most of the previously reported chemical shifts. The ^1H – ^{13}C cross-polarization (CP) contact time was 1 ms. A ^{13}C spin diffusion mixing time of 20 ms was used to obtain primarily intra-residue cross peaks. ^1H irradiation at a field strength that matches the MAS frequency (DARR) was applied during the mixing time to facilitate ^{13}C spin diffusion (Takegoshi et al. 2001). For the ^{13}C – ^{15}N REDOR period, ^{15}N inversion was achieved by composite $90^\circ_{\phi-90}$ $180^\circ_{\phi}90^\circ_{\phi-90}$ pulses to reduce the effects of flip-angle

errors (Sinha et al. 2004). ^{13}C , ^{15}N -enriched Gln, which contains both a COO and an N–CO group, was used to optimize the COO selection experiment. The main parameters of optimization to achieve complete suppression of the undesired peaks were the asymmetric timing of the ^{15}N π pulse within each rotor period in the COO selection experiment, the position and duration of the gated ^1H decoupling in the dynamic selection experiment, and the multiple-quantum (MQ) excitation time in the CH selection experiment.

Numerical simulations

REDOR curves were simulated using SPINEVOLUTION (Veshtort and Griffin 2006). ^{13}C and ^{15}N pulse lengths were finite and matched the values used in the experiments, which were 8 μs for ^{13}C and 10 μs for ^{15}N . Composite ^{15}N $90^\circ_{\phi-90}$ $180^\circ_{\phi}90^\circ_{\phi-90}$ pulses were also used in the simulations. Where ^1H spins were included, 60–250 kHz continuous-wave decoupling was applied. The MAS rotation period was 100 μs , corresponding to the 10 kHz MAS frequency of the experiments. Asymmetric REDOR refocuses isotropic chemical shift as well as the chemical shift anisotropy, thus we ignored chemical shift in the simulations. Relaxation was also ignored. Powder averaging used a REPULSION scheme containing 700 sets of (α, β) angles for the crystallite orientations (Bak and Nielsen 1997).

Results and discussion

Principle of spectral editing by asymmetric REDOR

Figure 2 shows the pulse sequences of COO, CH, and dynamics edited 2D ^{13}C – ^{13}C exchange experiments. In all three experiments, suppression of the undesired signals is achieved by recoupled heteronuclear dipolar interactions using a variant of REDOR (Gullion and Schaefer 1989), with two π -pulses per rotor period (t_r).

In spectral editing, it is crucial to dephase the undesired peaks, which are often larger than the signals of interest, to $<2\%$. The standard REDOR pulse sequence, where the π -pulses are spaced by $t_r/2$, generally does not permit this level of peak suppression. While the minimum of the ideal REDOR dephasing curve is -3% , in practice the minimum is often raised to greater than 10% , which precludes complete suppression of the undesired signals. The raised minimum can result from inaccurate flip angles of the recoupling pulses, whose effects are significantly but not completely suppressed by composite pulses (Sinha et al. 2004). A second cause is insufficient ^1H decoupling during the inversion pulses. Figure 3a displays the simulated REDOR dephasing curves for a 1.0-kHz dipolar coupling

Fig. 2 Spectral editing pulse sequences for ^{13}C , ^{15}N -labeled proteins. **a** COO selection by ^{15}N - ^{13}C asymmetric REDOR. **b** CH selection by dipolar DEPT. **c** Dynamic residue selection by ^1H - ^{13}C recoupled gated decoupling. For (a) and (c), standard phase cycles involving inversion of the ^1H 90° excitation pulse with respect to spin lock, CYCLOPS of the ^{13}C pulses, and EXORCYCLE of the ^{13}C 180° pulse, are used, with corresponding changes of the receiver phase. For the DEPT experiment (b), the full phase cycle involves CYCLOPS permutation of the basic element shown here. All pulse programs can be found at <http://www.public.iastate.edu/~nmrksr/>

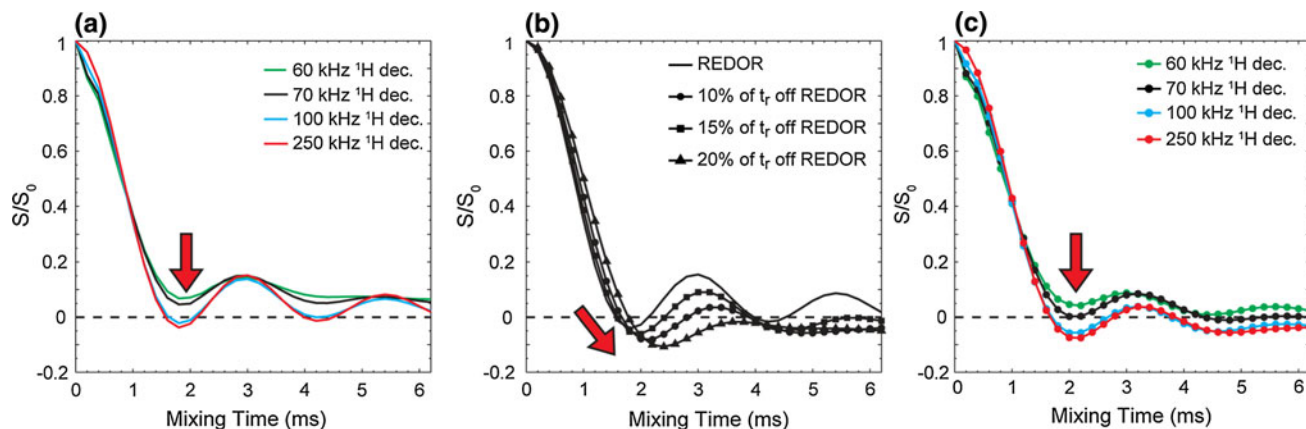
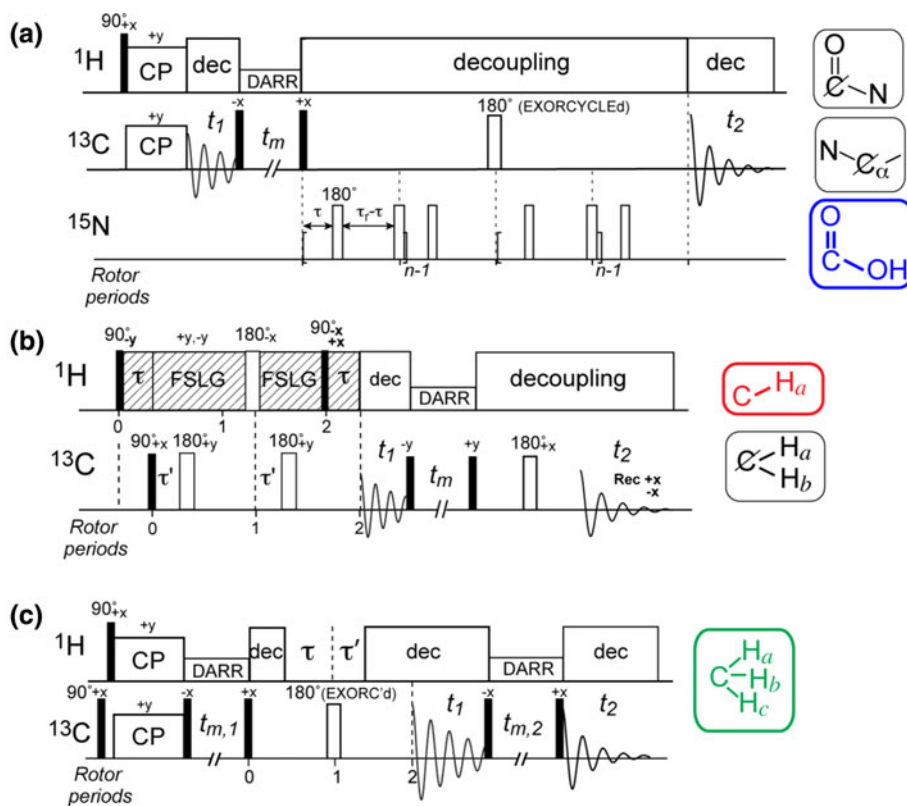


Fig. 3 Simulated ^{15}N - ^{13}C REDOR dephasing curves for a 1.0-kHz ^{13}C - ^{15}N dipolar coupling (corresponding to a ^{13}C - ^{15}N distance of 1.458 Å) to achieve complete suppression of the ^{15}N -bonded ^{13}C signals. **a** Effects of ^1H decoupling field strength on regular symmetric REDOR dephasing. The weaker the ^1H decoupling field, the higher the minimum S/S_0 . Zero crossing cannot be achieved at decoupling fields weaker than 70 kHz. A six-spin system containing one ^{13}C , one ^{15}N , and four protons was used, with a geometry that reflects the amino acid backbone plus two $\text{H}\beta$ protons. **b** Effects of the

π -pulse position within the rotor period on REDOR dephasing. Ideal ^1H decoupling is assumed by excluding protons in the simulations. The more asymmetric the π pulse position, the more negative the REDOR minimum, but at the expense of slowing down the REDOR oscillation. **c** Effects of ^1H decoupling field on the dephasing of 15% asymmetric REDOR. Zero crossing can be reached even under moderate ^1H decoupling. All simulations used composite ^{15}N π pulses with finite pulse lengths that are identical to the experimental pulse lengths

under moderate ^1H decoupling fields, where the minimum intensity is raised from the ideal value. For example, under 70-kHz ^1H decoupling, the REDOR minimum is +5%. It has been shown empirically (Sinha et al. 2004) that the ratio of the ^1H and ^{15}N pulse field strengths must exceed

three for the minimum of the REDOR curve to approach zero.

To lower the REDOR minimum to zero and completely suppress undesired signals at practical ^1H decoupling field strengths, we take advantage of asymmetric REDOR

(Gullion and Schaefer 1989). Here, the π pulses that are usually applied at the center of a rotor period are shifted by a fraction of t_r , typically $0.15 t_r$ (Fig. 2a). The effect of this asymmetry on the depth of the dephasing minimum can be understood qualitatively by considering the fact that the displacement of every other pulse moves the pulse sequence towards the quasi-static limit, where the smaller spacing between the recoupling pulses becomes so short that sample rotation during this period can be neglected (Bax et al. 1983; Tycko and Dabbagh 1990). The time signal of the spin-pair dipolar dephasing in the static limit is well known: it is the Fourier transform of the Pake pattern, and has the desired deep minimum below zero due to the oscillation associated with the doublet peaks of the Pake pattern (Schmidt-Rohr and Spiess 1994).

For a quantitative analysis, Fig. 3b shows simulated REDOR dephasing curves under ideal ^1H decoupling for varying positions of the π pulse, specified in terms of the fraction of t_r by which the pulse within the rotor period is shifted from the center of each rotor period. With 10–20 % asymmetry, the REDOR minimum ranges from -6 to -11 %, thus establishing zero-crossing points for complete suppression of the undesired signals. The secondary maxima, which have significant intensities in standard REDOR, also decrease in intensity, and the time regime with low intensities broaden, both facilitating peak suppression. When a ^1H decoupling field of realistic strength is applied, the asymmetric-REDOR dephasing curves retain the downward shift of the minimum intensity (Fig. 3c). For

example, under 70 kHz ^1H decoupling, the intensity of the first minimum is zero for a pulse shift of 10–15 % t_r , compared to the minimum intensity of +5 % when standard symmetric REDOR is used.

COO selection by N–CO suppression

Figure 4 demonstrates COO selection of 2D ^{13}C PDS spectra by asymmetric ^{13}C – ^{15}N REDOR. We used 15 % asymmetry and a total REDOR period of 2.2 ms to completely suppress the peaks of immobile N–CO and N–C α carbons. The REDOR period can be placed either before the t_1 period to suppress the C–N signals in the ω_1 dimension (Fig. 4b) or between the t_1 and t_2 periods to suppress the signals in the ω_2 dimension (Fig. 4c). Since the 2D ^{13}C PDS spectrum usually shows higher cross peak intensities on the upper left side ($\omega_1 < \omega_2$) than the lower right side ($\omega_1 > \omega_2$) of the diagonal due to the higher CP efficiency of protonated carbons, we focus on the ω_2 -edited CN-free spectrum.

Figure 5 shows the expanded carbonyl-aliphatic region of the full versus ω_2 -COO-selected 2D spectra. In the full spectrum (Fig. 5a), the cross peaks of Asp and Glu COO groups partially overlap with the NCO peaks of Asn and Gln sidechains, and the COO–C α cross peaks of Glu and Asp partially overlap with the Ala CO–C α cross peaks. With a 2.2-ms ^{13}C – ^{15}N REDOR dephasing time, all cross peaks to the backbone NCO and most of the sidechain Asn and Gln signals are suppressed, while all cross peaks of Asp and Glu

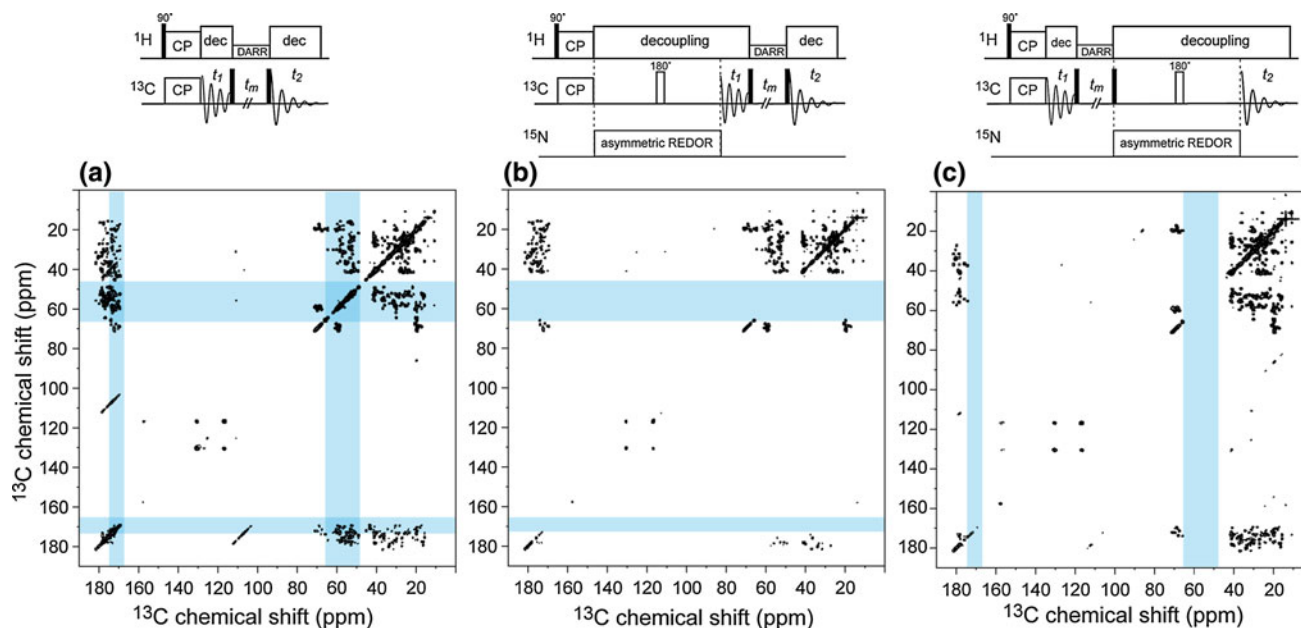


Fig. 4 COO selection of the 2D ^{13}C – ^{13}C correlation spectra by ^{15}N – ^{13}C dipolar dephasing. **a** Unedited 2D PDS spectrum of GB1. **b** ω_1 -edited 2D spectrum, where the C α and N–CO signals are removed in the ω_1 dimension. **c** ω_2 -edited 2D spectrum, where the C α

and N–CO signals are suppressed in the ω_2 dimension. The pulse sequence for each spectrum is shown at the top to clarify the distinction between the ω_1 and ω_2 -edited spectra. Shaded blue bars indicate the region where N–C α and N–CO signals are suppressed

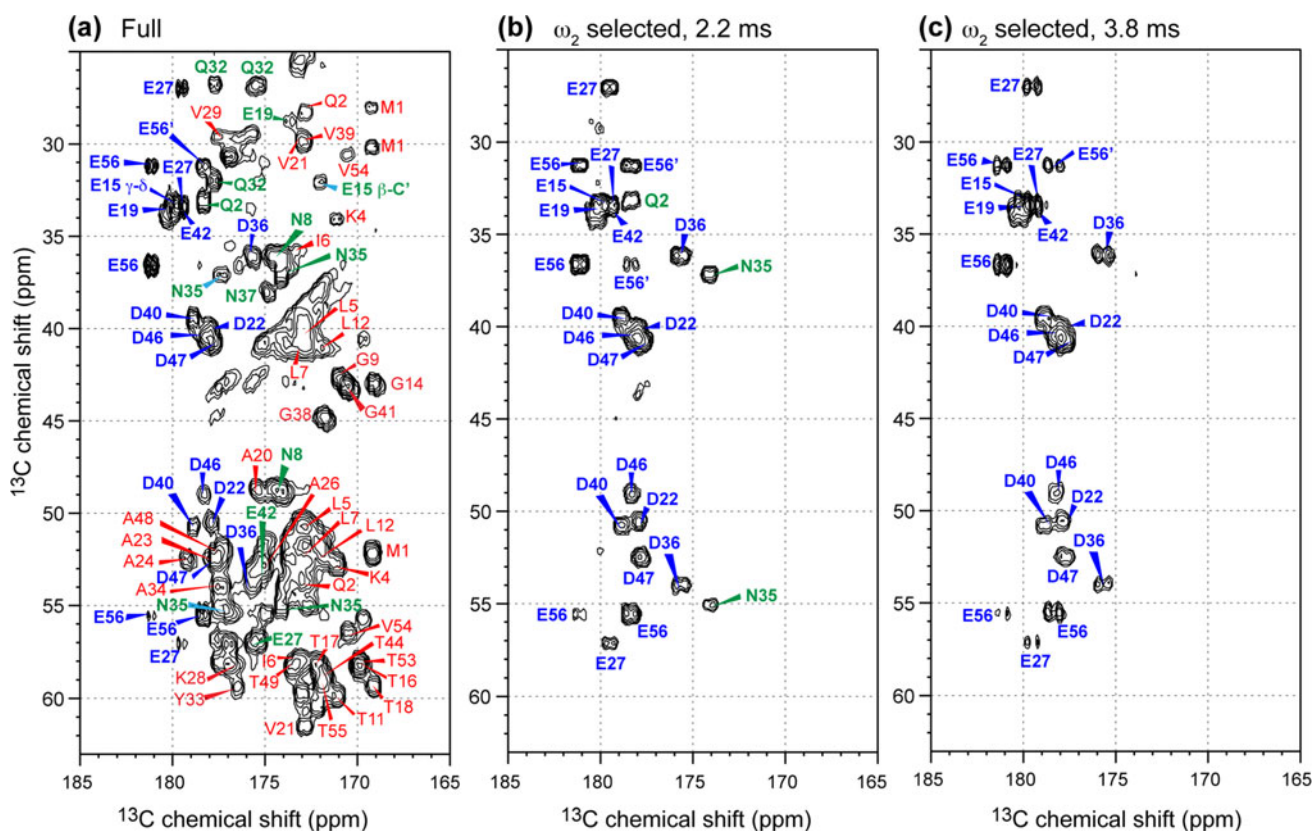


Fig. 5 Expanded carbonyl region of the 2D COO-selected spectra. **a** Unfiltered spectrum. The COO signals of Glu (E) and Asp (D) are assigned in *blue*, the sidechain and backbone N–CO signals of Gln (Q) and Asn (N) are assigned in *green*, and the backbone N–CO signals of all other amino acids are assigned in *red*. The backbone COO peaks of the terminal E56 are denoted as E56'. **b** ω_2 -COO

selected spectrum with a REDOR period of 2.2 ms. **c** ω_2 -COO-selected spectrum with a REDOR period of 3.8 ms. The signals of dynamic Q2 and N35 (*green*), which survive the 2.2 ms filter, are suppressed by the longer REDOR time. Spectra (**b**, **c**) were measured with $\tau = 40 \mu\text{s}$ in Fig. 2a, which shifts the position of the ^{15}N π pulse within the rotor period by 0.1 t_r before the center of the rotor period

sidechain COO and the signal of the terminal COO of the protein backbone (E56) survive (Fig. 5b). The only signals that should not be present are those of N35 and Q2 sidechains; presumably they remain due to partial motional averaging of the ^{13}C – ^{15}N dipolar coupling. Using a longer REDOR dephasing time of 3.8 ms, we were able to suppress these signals (Fig. 5c) while retaining the Glu and Asp peaks, giving a cleanly edited COO-only spectrum.

The 2D PDSO spectra were measured using a spin diffusion mixing time of 20 ms, which allowed the observation of intra-residue cross peaks within two or three bonds from the COO carbons. The peak positions of Asp (D) are mostly in a rectangle between 175 and 179 ppm in the ω_2 dimension and 35 and 55 ppm in the ω_1 dimension. These signals are relatively distinct from the Glu (E) peaks, which have larger COO (ω_2) frequencies and thus appear more to the left. Thus, type assignment of Glu and Asp residues can be made based on the peak positions in the COO-selected 2D spectra.

The frequencies of the $C\beta$ and $C\alpha$ peaks in each vertical slice of the ω_2 -COO-selected spectrum contain valuable information about the secondary structure of the Glu and

Asp residues. For instance, the peak positions for D36, with a relatively small $C\beta$ and relatively large $C\alpha$ chemical shift, are characteristic of an α -helix (Wishart et al. 1992). The same is true for E27, which exhibits peaks at the top and bottom of the spectrum. On the other hand, the chemical shifts of D46 suggest a β -sheet conformation. These conclusions are in agreement with the structure of GB1 and were previously obtained from 3D NMR analysis (Franks et al. 2008). We point them out here to demonstrate how spectrally edited 2D spectra can provide a low-resolution view of the protein secondary structure, in a manner that would be compatible with significant line broadening. Since the COO-selected 2D spectrum is sparse, longer spin diffusion mixing times can be used to detect inter-residue contacts from the Glu and Asp sidechains to provide valuable tertiary structure constraints.

The sensitivity of this COO selection experiment is very high, about 60 % of the unedited experiment. The intensity reduction mainly results from ^{13}C T_2 relaxation and evolution under ^{13}C – ^{13}C J-coupling during the REDOR mixing period.

CH and dynamic selection experiments

To select the CH signals in 2D spectra, we used the dipolar analog of the DEPT (distortionless enhancement by polarization transfer) experiment, adapted for 10-kHz MAS (Fig. 2b) (Schmidt-Rohr and Mao 2002a, b). In this experiment, ^1H magnetization H_x is converted to two-spin coherence C_zH_y by a short evolution period (τ) under C–H dipolar coupling. A 90° x-pulse on ^{13}C converts this coherence to C_yH_y , a combination of double- and zero-quantum coherence. For methine (CH) groups, this two-spin coherence survives the subsequent dephasing period with heteronuclear C–H dipolar recoupling, since it commutes with the heteronuclear dipolar Hamiltonian $\omega_{CH}2C_zH_z$, while the additional proton in CH_2 groups destroys this coherence by its dipolar coupling to the carbon. Using a ^1H 90° x-pulse and a second heteronuclear evolution period τ identical to the first, the surviving two-spin coherence of CH groups is reconverted to observable ^{13}C magnetization $-C_x$. Thus, the MQ excitation and reconversion period, in addition to removing the CH_2 signals, also serves to transfer polarization from ^1H to ^{13}C .

During the excitation, evolution, and reconversion of the MQ coherence, ^1H – ^1H homonuclear dipolar couplings are suppressed by the FSLG sequence (Bielecki et al. 1989). Because of the dependence of the dipolar couplings on the rotor orientation, the MQ reconversion period and excitation period must be separated by an integer multiple of t_r , except when the excitation period is half a t_r , in which case the two periods can be separated by multiples of $t_r/2$. The ^1H and ^{13}C π pulses during the MQ evolution period between the ^{13}C 90° pulse and the second ^1H 90° pulse prevent the refocusing of the heteronuclear dipolar evolution by MAS. In addition, they refocus most of the ^1H and all of the ^{13}C chemical shift evolution. The π pulses are spaced in the asymmetric REDOR fashion, with a shift of every other pulse by $\sim 20\%$ of t_r .

Figure 6a, b compares the full 2D PDS spectrum with the CH-edited spectrum, in which the CH signals of Val, Leu and Ile (in red) are selected and resolved from the overlapping CH_2 resonances. The selection particularly allowed the identification of the Val $C\beta$ – $C\alpha$ cross peaks, which overlap significantly with the Lys, Glu and Gln cross peaks. Leu $C\gamma$ – $C\beta$ cross peaks also benefitted from editing and became resolved from various Lys peaks. The CH signals of these three amino acids fall into three bands with little overlap (34–28 ppm from TMS for Val, 26–23 ppm for Leu, and 42–32 ppm for Ile), showing highly characteristic cross-peak patterns. Thus, the CH selection allows amino acid type assignment simply from the observed peak positions in most cases. The sensitivity of the CH-edited spectrum is about 25 % of the full spectrum. This is comparable to the sensitivity of most homonuclear double-

quantum filtered 2D experiments and heteronuclear 3D NCC experiments.

As before for Glu and Asp, the resolved $C\alpha$ and $C\beta$ chemical shifts of Val, Leu and Ile from the edited 2D spectrum provide information on the backbone conformation. For instance, V29 with its high $C\alpha$ and low $C\beta$ shifts is clearly in an α -helix, while V54 at the other end of the $C\alpha$ chemical shift range is expected to be in a β -sheet. The chemical shifts of I6 show that it is clearly not in an α -helix, but suggest a β -sheet. None of the three Leu residues in the protein (L5, L7, and L12) show α -helical characteristics, indicating that the α -helix does not include residues 5 through 12. These conclusions are consistent with the previous 3D NMR analysis (Franks et al. 2005b).

The CH edited spectrum in Fig. 6b also retains undesired signals of mobile CH_2 groups (in green italics) of some long aliphatic sidechains, in particular Lys residues. These signals of dynamic residues could be reduced significantly by using a short MQ excitation and reconversion time of $\sim 30\ \mu\text{s}$. Fortunately, there is no significant spin exchange from sites far out in mobile sidechains to backbone $C\alpha$ or CO. Therefore, the $C\alpha$ – $C\beta$ or $C\alpha$ – $C\gamma$ cross peaks of Ile, Val, and Leu can be determined unambiguously. In addition, the cross peaks of Lys $C\epsilon$ and $C\delta$ are mostly resolved from the CH-selected $C\beta$ – $C\gamma$ signals of Ile and Val. Alternatively, CH selection at low temperature where motion is suppressed can remove the CH_2 signals. Scalar-coupling based spectral editing may also provide an alternative approach to select CH signals independent of motion (Sakellariou et al. 2001).

When studying proteins at physiological temperature, one can identify the dynamic residues using a dynamics selection experiment (Fig. 2c). In this experiment, a 60- μs period without ^1H decoupling is combined with a ^{13}C π pulse that recouples the ^1H – ^{13}C dipolar interaction. The polarization of the dynamic moieties was enhanced by combining direct polarization and cross polarization, with a 90° ^{13}C pulse before CP. After CP, a 90° pulse flips the ^{13}C magnetization to the z-direction and a ~ 50 ms mixing period was applied to transfer the magnetization from the well-polarized rigid protonated carbons to the mobile segments and the non-protonated carbons. This combined polarization approach is general and can be incorporated whenever the signals of both mobile and rigid residues are of interest. It complements scalar-coupling based polarization transfer schemes that detect only residues with nearly isotropic mobility in proteins (Etzkorn et al. 2007).

Figure 6c shows the dynamics edited spectrum, where the signals of all methyl groups and the methylene signals of several Lys and Glu residues are detected. The spectrum indicates which peaks in the CH-selected spectrum result from mobile CH_2 and CH_3 groups. The efficiency is about 50 % for methyl groups and highly mobile methylene groups.

most prominently exhibits the various cross peaks of K10, which is uniquely identified based on its $C\beta$ chemical shift. These peaks indicate that not only $C\epsilon$ but also $C\delta$ and $C\gamma$ of K10 undergo large-amplitude motion. This is consistent with the position of K10 in the loop between the β_1 and β_2 strands (Franks et al. 2008). The lack of mobility of K31 may be attributed to its position in the central α -helix, but the absence of dynamics for K50 is somewhat surprising, since K50 is located in the loop between the β_4 and β_5 strands (Franks et al. 2008).

The dynamics of the M1 and Q2 sidechains may be attributed to increased mobility near the N terminus of the protein. In Fig. 6b, we observe a relatively weak $C\beta$ – $C\alpha$ cross peak of V21, located in the loop between the β_2 and β_3 strands, which is also due to dynamics as discussed in (Franks et al. 2008).

Application of spectral editing to disordered proteins with broad lines

To demonstrate how spectral editing can facilitate secondary structure analysis of disordered proteins with broad linewidths, we measured the 2D ^{13}C PDSM spectra of uniformly ^{13}C -labeled ubiquitin. The sample is a lyophilized powder that was not subject to crystallization and thus exhibits ^{13}C linewidths of 1–2 ppm. The ^{13}C chemical shifts of microcrystalline ubiquitin have been almost fully assigned (Igumenova et al. 2004), thus allowing us to verify our conclusions.

Figure 7 shows the full (a) and CH-selected (b) 2D spectra: the cross peak patterns are significantly simplified after CH selection. The selected signals of the seven Ile, four Val, and eight Leu residues in ubiquitin can be analyzed in terms of β -sheet, α -helix, and loop conformations. All three amino-acid types show strong $C\alpha/C\beta$ cross peaks for multiple residues at positions typical of β -sheet (e.g., Ile: 58/39 ppm, TMS scale; Val: 57/34 ppm). Much smaller peaks of one or two residues each are observed at positions characteristic of α -helices (Ile: 64/34 ppm, Val: 65/28 ppm). Various $C\beta$ – $C\gamma$ cross peaks of Ile were observed around a $C\beta$ chemical shift of 33 ppm, which is typically associated with the α -helical conformation, but not all these have $C\alpha$ chemical shifts typical of α -helices. This suggests Ile in a loop or at the end of an α -helix, which is confirmed by a distinct $C\beta$ – $C\delta$ cross peak (marked ‘loop’) with an unusually low $C\beta$ but also a low $C\alpha$ chemical shift, incompatible with α -helix or β -sheet. On the basis of these cross peak positions, one can propose that the protein is significantly β -sheet but also contains some α -helical segments and loops; indeed, ubiquitin has $\sim 35\%$ β -sheet and $\sim 25\%$ α -helical residues (Heinig and Frishman 2004; Kabsch and Sander 1983).

This type of secondary-structure information can often be gleaned from the resolved $C\alpha$ – $C\beta$ cross peaks of Ala,

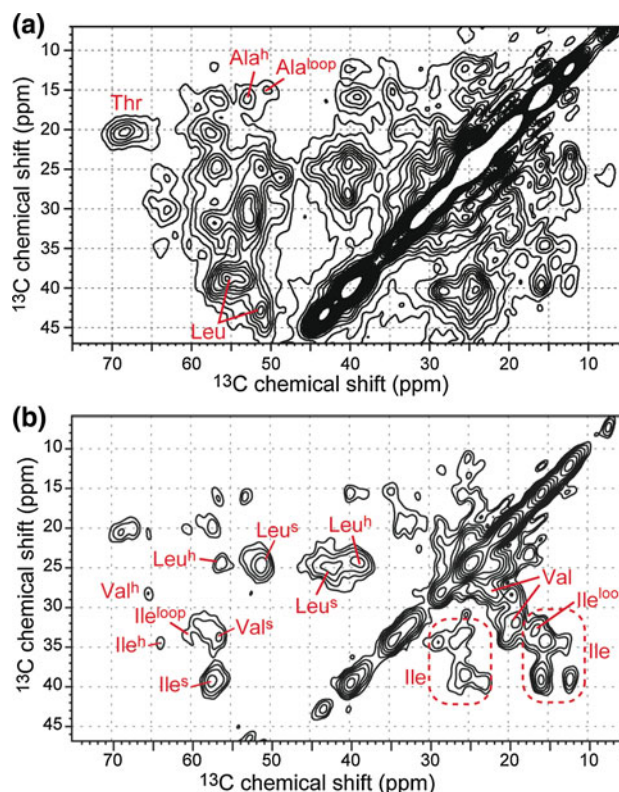


Fig. 7 2D ^{13}C PDSM spectra of lyophilized ubiquitin **a** without editing and **b** with CH selection. The CH selection is applied to the ω_1 dimension of the 2D spectrum. Amino acid types are annotated along with the identified secondary structures. Helix: h; sheet: s, and loop. The spectra were measured at room temperature on a 400 MHz SSNMR spectrometer

but the two Ala residues in ubiquitin show no β -sheet character, only an α -helical and a loop conformation (Fig. 7a). Only in some of the Thr $C\alpha$ – $C\beta$ peaks far off the diagonal (not shown) does the unedited spectrum give a clear indication of a high β -sheet content.

Next, one can look for segments in the amino acid sequence of ubiquitin containing one Ala, Ile, Val, and Leu residue each, as candidates for α -helical stretches. This search yielded the segment Val26–Ile30 (VKAKI), which is indeed part of an α -helix in ubiquitin, as well as Leu43–Ala46 (LIFA), which is not α -helical (Igumenova et al. 2004). To distinguish between these two cases, sequential resonance assignment is necessary.

In summary, the CH-selected 2D spectrum yields secondary-structure information and helps with the identification of secondary-structure elements even in the presence of significant line broadening.

Potential spectral editing in 3D NMR

The COO selection experiment can be incorporated into 3D NCOX and NCAX experiments, to permit assignment

of residues close to Asp and Glu, with minimum resonance assignment. The pulse sequence module should be implemented in the ω_3 ^{13}C dimension, to remove non-E/D signals. The dynamics editing experiment can also be incorporated into 3D NCC experiments, provided that long mixing times are used to transfer the backbone CO or C α magnetization to the mobile sidechain ends.

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