

Probing the urea dependence of residual structure in denatured human α -lactalbumin

Victoria A. Higman · Heike I. Rösner ·
Raffaella Ugolini · Lesley H. Greene ·
Christina Redfield · Lorna J. Smith

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Abstract Backbone ^{15}N relaxation parameters and ^{15}N – ^1H residual dipolar couplings (RDCs) have been measured for a variant of human α -lactalbumin (α -LA) in 4, 6, 8 and 10 M urea. In the α -LA variant, the eight cysteine residues in the protein have been replaced by alanines (all-Ala α -LA). This protein is a partially folded molten globule at pH 2 and has been shown previously to unfold in a stepwise non-cooperative manner on the addition of urea. ^{15}N R_2 values in some regions of all-Ala α -LA show significant exchange broadening which is reduced as the urea

concentration is increased. Experimental RDC data are compared with RDCs predicted from a statistical coil model and with bulkiness, average area buried upon folding and hydrophobicity profiles in order to identify regions of non-random structure. Residues in the regions corresponding to the B, D and C-terminal 3_{10} helices in native α -LA show R_2 values and RDC data consistent with some non-random structural propensities even at high urea concentrations. Indeed, for residues 101–106 the residual structure persists in 10 M urea and the RDC data suggest that this might include the formation of a turn-like structure. The data presented here allow a detailed characterization of the non-cooperative unfolding of all-Ala α -LA at higher concentrations of denaturant and complement previous studies which focused on structural features of the molten globule which is populated at lower concentrations of denaturant.

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V. A. Higman · R. Ugolini · L. H. Greene · L. J. Smith (✉)
Department of Chemistry, Inorganic Chemistry Laboratory,
University of Oxford, South Parks Road, Oxford, UK
e-mail: lorna.smith@chem.ox.ac.uk

H. I. Rösner · C. Redfield (✉)
Department of Biochemistry, University of Oxford,
South Parks Road, Oxford, UK
e-mail: christina.redfield@bioch.ox.ac.uk

Present Address:

V. A. Higman
Leibniz-Institut für Molekulare Pharmakologie,
Robert-Rösle-Str. 10, Berlin, Germany

Present Address:

H. I. Rösner
Department of Biology, University of Copenhagen,
Ole Maaløes Vej 5, 2200 Copenhagen, Denmark

Present Address:

L. H. Greene
Department of Chemistry and Biochemistry,
Old Dominion University, 4541 Hampton Blvd., Norfolk,
VA 23529-0126, USA

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Introduction

While structural biology generally concerns itself with the study of folded states of proteins, the study of denatured, unfolded or partially-folded states is equally important. Understanding the structure and dynamics of these states can help to elucidate mechanisms and processes which occur during protein folding (Arai and Kuwajima 2000; Dinner et al. 2000; Ptitsyn 1995). Non-native states of proteins have also been recognized to be involved in a range of disease states (Dobson 2001; Kelly 2002). In

addition, it has been shown that many proteins and protein domains are unstructured or only partially structured under physiological conditions (Wright and Dyson 1999). Non-native protein states are not characterized by a single stable three-dimensional structure. Instead they consist of ensembles of interconverting conformers. NMR is well suited to the study of such non-native protein states (Dyson and Wright 2001; Shortle 1996). Among the NMR techniques and parameters which have proved to be useful in defining conformational ensembles are chemical shift deviations (Eliezer et al. 2000; Ramboarina and Redfield 2003; Spera and Bax 1991; Wishart and Sykes 1994), $^3J_{\text{NH}\alpha}$ coupling constants (Smith et al. 1996), short-range ^1H – ^1H NOEs (Schwalbe et al. 1997), paramagnetic relaxation enhancement (Gillespie and Shortle 1997a, b), ^{15}N relaxation measurements (Dyson and Wright 2001; Klein-Seetharaman et al. 2002; Schwalbe et al. 1997) and unfolding with urea (Schulman et al. 1997; Wang et al. 1995; Wang and Shortle 1995). Together with mutagenesis (Klein-Seetharaman et al. 2002; Nishimura et al. 2005; Song et al. 1998; Wu and Kim 1998) these have enabled the localization of short-range secondary structure propensities and long-range non-native hydrophobic clusters within unfolded and partially-folded proteins.

More recently, residual dipolar couplings (RDCs) have emerged as an important tool for characterizing residual structure in unfolded or partially-folded proteins (Meier et al. 2008). Initial studies by Shortle and co-workers on denatured staphylococcal nuclease, eglin c and short unstructured peptides showed these to have comparatively uniform non-zero RDC values (Ackerman and Shortle 2002a, b; Ohnishi et al. 2004; Ohnishi and Shortle 2003; Shortle and Ackerman 2001). Predictions using random flight and valence chains show this to be the expected behavior of a fully-denatured polypeptide chain (Louhivuori et al. 2004, 2003; Obolensky et al. 2007). Improved fits between experimental and predicted RDCs are obtained when using structural models derived from Φ/Ψ propensities of coil regions in folded proteins (Bernadó et al. 2005b; Jha et al. 2005). Deviations from these models can be used to identify regions of persistent residual structure. This approach has been used to characterize the populations of residual helical structure in ribonuclease S-peptide (Alexandrescu and Kammerer 2003), ACBP (Fieber et al. 2004), apomyoglobin (Mohana-Borges et al. 2004) and Sendai virus nucleoprotein (Jensen et al. 2008), to probe persistent long-range contacts in α -synuclein (Bernadó et al. 2005a) but lacking in β -synuclein (Bertoncini et al. 2007), to identify residues involved in persistent hydrophobic clusters or close to proline residues in a nucleocapsid-binding domain (Bernadó et al. 2005b), apomyoglobin (Mohana-Borges et al. 2004) and the A-state of foldon (Meier et al. 2004) and to locate turn

conformations populated in denatured ubiquitin (Meier et al. 2007) and natively unfolded Tau protein (Mukrasch et al. 2007). In addition, variations in RDC values with temperature have been used to identify melting hotspots during the unfolding of the protein GB1 (Ding et al. 2004) and to characterize the unfolding of the β -hairpin in the A-state of foldon (Meier et al. 2004).

In this paper we use ^{15}N relaxation and RDC measurements to study the stepwise unfolding in urea of a molten globule formed by a variant of human α -lactalbumin (α -LA) (Fig. 1). Non-native states of α -lactalbumin have been studied extensively using a wide range of experimental and theoretical approaches (Kuwaitjima 1996; Ramboarina and Redfield 2003, 2008; Redfield et al. 1999; Schulman et al. 1997, 1995; Smith et al. 2005; Song et al. 1998; Wu and Kim 1998). α -LA forms a molten globule, the so-called A-state, at low pH (Ptitsyn 1995). Heteronuclear NMR methods have shown that this A-state unfolds in a very non-cooperative manner on the addition of urea (Schulman et al. 1997). A core containing residues from the native-state A, B, D and C-terminal 3_{10} helices in α -LA persists even in 10 M urea. α -LA contains four disulfide bonds. The unfolding of a variant of α -LA, in which the eight cysteine residues have been replaced by alanines (all-Ala α -LA), has been characterized using the same approach (Redfield et al. 1999). Interestingly, this variant shows a similar persistent core and non-cooperative unfolding on the addition of urea. However, in all-Ala α -LA the core containing residues from the A, B, C, D and C-terminal 3_{10} helices only persists up to concentrations of 3 M urea. Unfolding is virtually complete in 7 M urea, as indicated by the observation of all expected peaks in the ^{15}N – ^1H HSQC spectrum; circular dichroism measurements at 222 nm show that almost all helical structure is lost in 7 M urea (Redfield et al. 1999). In this paper, we investigate the all-Ala α -LA variant by measuring

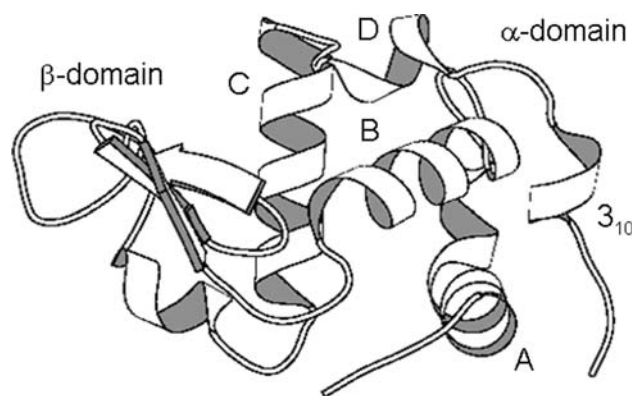


Fig. 1 Schematic representation of the native structure of human α -lactalbumin. The α - and β -domains and the five α -domain helices are labelled. The diagram was generated using MOLSCRIPT (Kraulis 1991)

^{15}N relaxation and RDC data for the protein at pH 2 in 4, 6, 8 and 10 M urea. By combining these data with ^{15}N chemical shifts, results from urea unfolding experiments and studies of small peptides we have been able to derive a comprehensive picture of the backbone dynamics and residual structure present during protein unfolding.

Materials and methods

^{15}N -labelled all-Ala α -LA was expressed and purified as described elsewhere for related proteins (Peng et al. 1995; Schulman et al. 1995). Samples for the ^{15}N relaxation and RDC measurements were prepared at a protein concentration of 0.4 mM in 4, 6, 8 and 10 M urea at pH 2. 6% polyacrylamide gels were used to align the samples for RDC measurements. These were prepared in a similar manner to that described by Sass et al. (2000). The isotropic data were collected first using Shigemi microcells. The dried gels were then added to the samples, and allowed to swell before being compressed from an initial length of 3 cm to a final length of 2.1 cm. In this manner it was possible to ensure that the urea concentration remained the same for both datasets. Repeat experiments were conducted in order to estimate the error. Average RDC values are reported and the error was found to be ~ 0.5 Hz at all urea concentrations. Despite this proportionately fairly large error, the magnitude and sign of RDCs was reproduced extremely well in separate datasets, even when using different samples.

NMR experiments were performed on a home-built NMR spectrometer, located in the Department of Biochemistry, operating at a ^1H frequency of 599.48 MHz. ^{15}N – ^1H RDCs were measured at 20°C using an HSQC experiment which incorporated an S^3E pulse-sequence element (Meissner et al. 1997). 128 and 1,024 complex points and sweep widths of 1,272.26 and 6,756.76 Hz were collected in F_1 (^{15}N) and F_2 (^1H), respectively. The data were processed using Felix 2.3 (Accelrys) and zero-filled to give final digital resolutions of 0.3 Hz/pt and 3.3 Hz/pt in F_1 and F_2 , respectively. ^{15}N R_1 and R_2 relaxation rates and the $\{^1\text{H}\}$ – ^{15}N NOE were measured using standard pulse sequences at 20°C, with 128 and 1,024 complex points and sweep widths of 1,152.07 and 6,514.66 Hz in F_1 (^{15}N) and F_2 (^1H), respectively. The R_1 and R_2 measurements used a series of 9 to 12 experiments with relaxation delays ranging from 20 to 2,500 ms and from 8.6 to 1,200 ms, respectively. The $\{^1\text{H}\}$ – ^{15}N NOE is reported as the ratio of peak heights in spectra collected with and without saturation of the ^1H spectrum by a non-selective 90° pulse every 10 ms for a period of 4 s. Errors in the measurement of peak heights were estimated from baseline noise. Fitting of R_1 and R_2 rates and reduced spectral density mapping

(Bracken et al. 1999; Farrow et al. 1995) were carried out using in-house programs written in Fortran77.

An ensemble of 5,000 all-Ala α -LA structures was generated using the Jha et al. algorithm (<http://unfolded.chicago.edu/>) (Jha et al. 2005). RDCs were predicted for this ensemble using the program PALES (Zweckstetter and Bax 2000). Each structure was aligned separately and the resulting RDCs were then averaged over the ensemble and scaled so that the mean predicted RDC value was the same as that observed experimentally. The average area buried upon folding (AABUF) (Rose et al. 1985) and hydrophobicity (Abraham and Leo 1987) profiles for all-Ala α -LA were calculated using the ExPASy web server (<http://expasy.org>) weighted over a 7-residue window. The bulkiness profile for all-Ala α -LA was calculated as described by Cho et al. (2007).

Results

^{15}N relaxation data for all-Ala α -LA

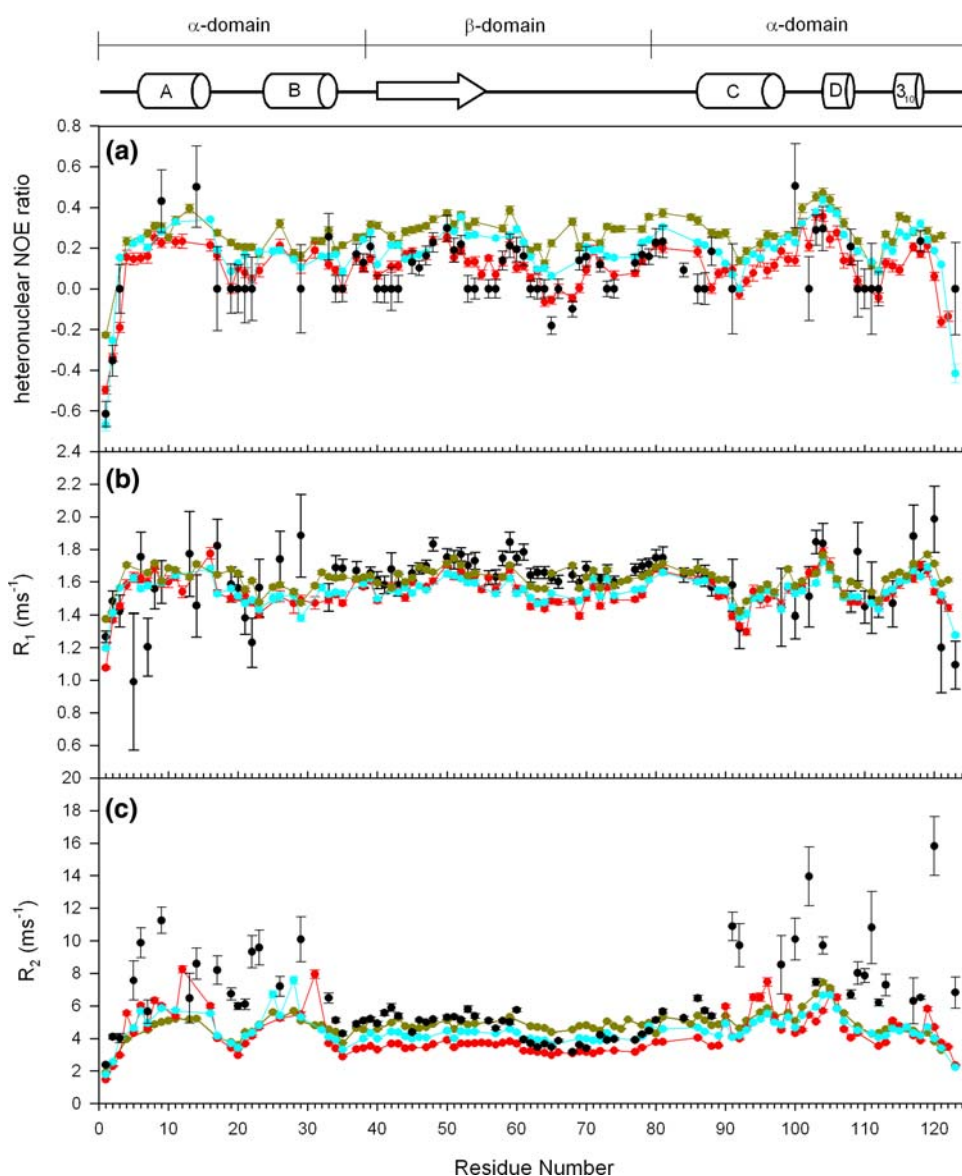
R_1 and R_2 relaxation rates and the $\{^1\text{H}\}$ – ^{15}N heteronuclear NOE ratio measured in 4, 6, 8 and 10 M urea are plotted as a function of the all-Ala α -LA sequence in Fig. 2 (the full data are provided as Electronic supplementary material). The relaxation data at high urea concentrations are similar to those reported for other denatured proteins (Schwalbe et al. 1997; Schwarzsinger et al. 2002). The R_2 relaxation rates, generally the most informative of the three parameters for denatured proteins, are smaller at the termini and rise to a plateau in the central region of the polypeptide chain. This is in agreement with predictions from a simple model of an unfolded polypeptide chain with unrestrained segmental motion in which the ^{15}N relaxation properties of a given amide group are not influenced to any significant extent by the identity of neighboring residues in the chain (Schwalbe et al. 1997). Deviations from this pattern are largest at 4 and 6 M urea and decrease as the urea concentration is increased. In 10 M urea, average R_2 values of 4.89 and 4.81 are observed for the α -domain (residues 1–39 and 82–123) and the β -domain (residues 40–81), respectively. Elevated R_2 relaxation rates are observed in the α -domain of the protein, particularly in 4 M urea. A sharp decrease in R_2 is observed around residue 20, particularly in 6, 8, and 10 M urea. The sequence from 17–20 is GYGG and the high conformational freedom of glycine residues presumably enables this region to be relatively more flexible at all urea concentrations. In addition, residues 17–22 correspond to a loop region between the native A and B helices (Fig. 1). This loop region has been observed to unfold at a lower urea concentration than the flanking helices (Redfield et al. 1999). A less pronounced decrease

in R_2 is observed in the vicinity of Gly35 at the end of the B helix. The differing plateau values for R_2 at the three highest urea concentrations reflect a viscosity effect; as the urea concentration is raised, the viscosity of the sample increases, the overall molecular tumbling is slowed and the R_2 relaxation rate is raised. Interestingly, the data collected at 4 M urea do not follow this trend and the R_2 values are higher than expected. Furthermore, residues 40–60 form a higher plateau than residues 61–80 which could be the result of differential exchange broadening in the β -domain in 4 M urea. This would correlate with previous NMR studies of all-Ala α -LA which have shown that the protein is not completely unfolded in 4 M urea (Redfield et al. 1999) and a recent study of backbone dynamics of the all-Ala α -LA molten globule which has shown differences in fast timescale dynamics within the β -domain; residues

40–60 show more restricted mobility than residues 61–80 (Ramboarina and Redfield 2008).

^{15}N relaxation data for folded proteins are usually analyzed using the model-free formalism of Lipari and Szabo; using this approach order parameters and internal correlation times can be extracted for individual residues (Lipari and Szabo 1982a, b). This approach is not appropriate for unfolded proteins because such systems cannot be described by an overall rotational correlation time. Instead, reduced spectral density mapping is often used to describe backbone motions in unfolded proteins (Farrow et al. 1997; Le Duff et al. 2006; Schwarzingner et al. 2002; Shojania and O'Neil 2006; Wirmer et al. 2006b) and this approach has been adopted here for all-Ala α -LA (Fig. 3). Both $J(0.87\omega_{\text{H}})$ and $J(\omega_{\text{N}})$ show little variation with amino acid sequence and urea concentration. Some deviations in these

Fig. 2 ^{15}N relaxation data for all-Ala α -LA plotted as a function of the amino acid sequence. **a** $\{^1\text{H}\}$ - ^{15}N NOE as $I_{\text{sat}}/I_{\text{nonsat}}$, **b** R_1 and **c** R_2 values for all-Ala α -LA in 4 M (black), 6 M (red), 8 M (cyan) and 10 M (gold) urea are shown. All data were collected at 600 MHz and 20°C. The secondary structure and domain organization found in native α -lactalbumin is summarized above



spectral densities are observed for α -domain residues in 4 M urea but these are associated with large errors due to the very weak peak intensities. $J(0)$, which is most sensitive to R_2 , shows a significant dependence on the urea concentration for residues located in the α -domain of the protein. The increased $J(0)$ values in the α -domain observed as the urea concentration is lowered reflect the increasing contributions from exchange processes on the millisecond to microsecond time scale as the protein becomes more structured.

^{15}N chemical shift data for all-Ala α -LA

^{15}N chemical shifts measured in 6, 8 and 10 M urea are in good agreement with the sequence-corrected shifts

predicted by Braun et al. (1994); correlation coefficients of 0.98 are obtained for comparison of predicted and experimental ^{15}N shifts in both 6 and 10 M urea. Nevertheless, differences in ^{15}N chemical shifts are observed as a function of urea concentration in the spectra of all-Ala α -LA (Redfield et al. 1999). The difference between the ^{15}N chemical shifts measured in 10 and 6 M urea is plotted as a function of sequence in Fig. 4a. Chemical shift differences as large as 1 ppm are observed. The largest differences are observed in the α -domain in regions of sequence that correspond to the native A, B, D and C-terminal 3_{10} helices. These are the regions of the protein that showed the largest exchange contributions to R_2 at lower urea concentrations. The chemical shifts for these regions in 6 M urea are shifted upfield of the 10 M values, consistent with the

Fig. 3 Reduced spectral density mapping for all-Ala α -LA at **a** 0 MHz ($J(0)$), **b** 60 MHz ($J(\omega_N)$) and **c** 522 MHz ($J(0.87\omega_H)$) in 4 M (black), 6 M (red), 8 M (cyan) and 10 M (gold) urea. The secondary structure and domain organization found in native α -lactalbumin is summarized above

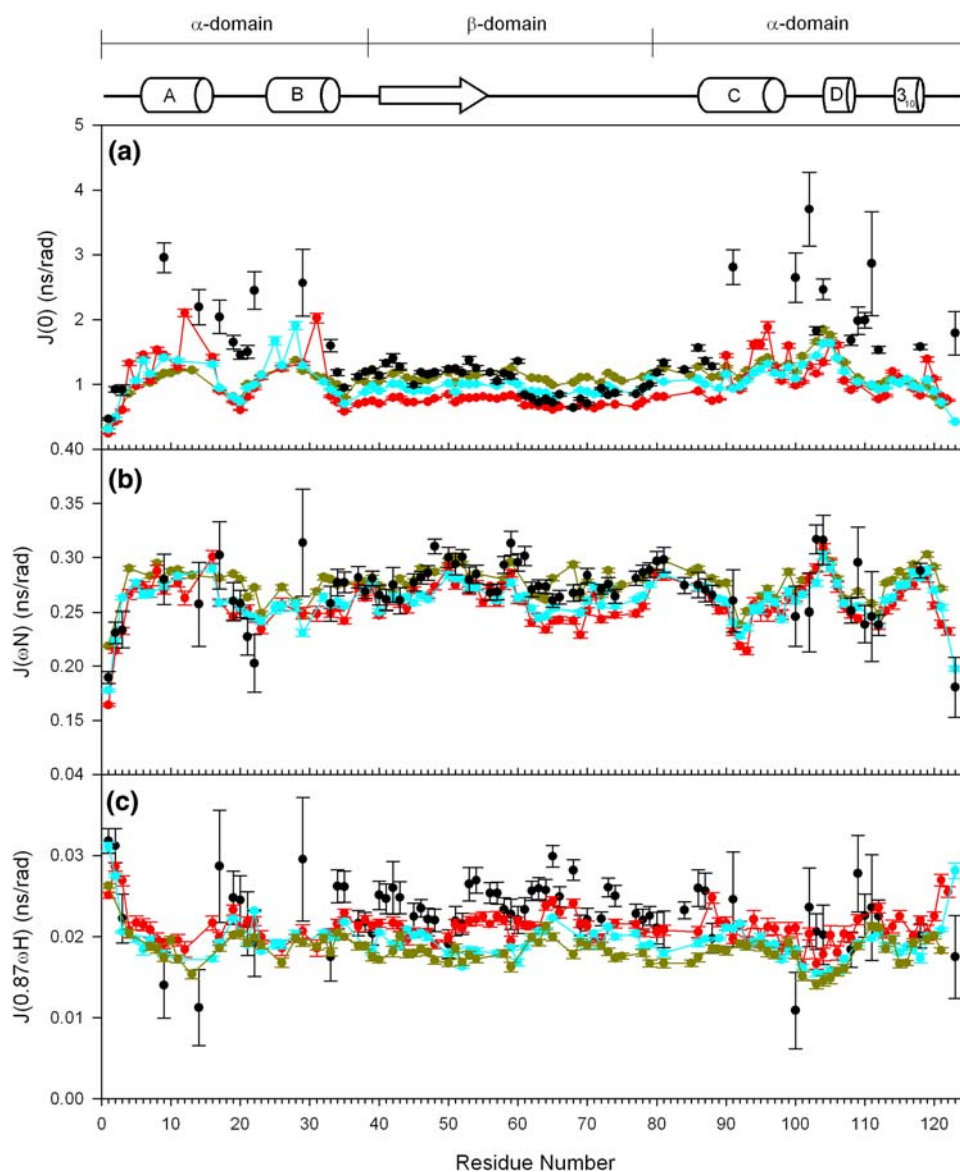
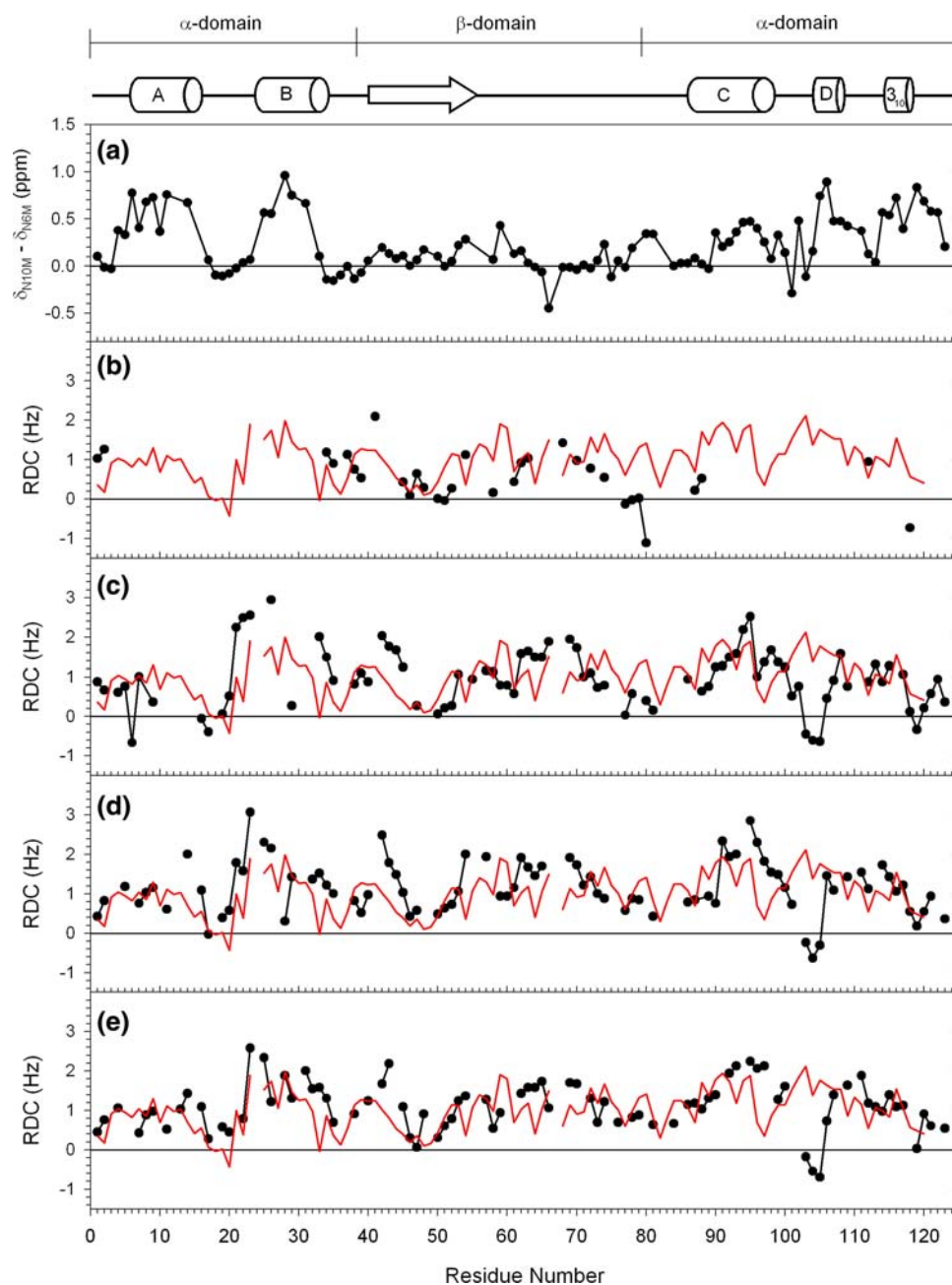


Fig. 4 **a** Difference in ^{15}N chemical shifts for all-Ala α -LA at 6 and 10 M urea at 20°C [$\delta_{15\text{N}}(10\text{ M}) - \delta_{15\text{N}}(6\text{ M})$]. **b–e** Experimental RDCs recorded for all-Ala α -LA in compressed polyacrylamide gels at 20°C (black) are compared with predicted RDCs for all-Ala α -LA based on a coil-model (red). Measurements were made in **b** 4 M, **c** 6 M, **d** 8 M and **e** 10 M urea. The secondary structure and domain organization found in native α -lactalbumin is summarized above



presence of residual helical propensity at the lower urea concentration (Wishart and Sykes 1994). For residues 17–23, which are located between the native A and B helices, much smaller chemical shift differences are observed; this is consistent with the unfolding of this loop at a lower concentration of urea. Smaller chemical shift differences are observed in the β -domain but the pattern is not uniform; somewhat larger differences are observed for residues 40–60 than observed for 61–80. The larger shift differences observed for residues 53, 54 and 59 are consistent with the recent observation of restricted backbone dynamics for residues in this region of the all-Ala α -LA

molten globule (Ramboarina and Redfield 2008). A large negative shift difference is observed for Val66; this residue precedes a proline in the sequence.

Experimental RDCs for denatured all-Ala α -LA

^{15}N – ^1H RDCs were measured for all-Ala α -LA in compressed polyacrylamide gels in the presence of 4, 6, 8 and 10 M urea at pH 2 and 20°C. The chemical shifts observed in the compressed gels are identical to those in isotropic solution, suggesting that the structural ensemble of all-Ala α -LA is not perturbed in the polyacrylamide gel medium.

The RDC values are plotted as a function of amino acid sequence in Fig. 4b–e (Tables listing the RDC values are available as Electronic supplementary material). The RDC values are positive for all four datasets with average values of 0.55, 0.95, 1.17 and 1.11 Hz in 4, 6, 8 and 10 M urea, respectively. The observation of positive non-zero RDC values for denatured all-Ala α -LA in compressed polyacrylamide gels is in agreement with predictions for random flight and valence chains (Louhivuori et al. 2004, 2003) and data reported for other denatured proteins (Fieber et al. 2004; Meier et al. 2004; Mohana-Borges et al. 2004). Deviations from uniform values across the sequence may be due to nearest neighbor effects (Jha et al. 2005) and the inherent Φ/Ψ preferences of this amino acid sequence (Bernadó et al. 2005b; Louhivuori et al. 2004). Alternatively it may be indicative of specific long-range interactions favoured by the polypeptide chain (Bernadó et al. 2005a). This will be explored in the next section using a coil model for all-Ala α -LA. The absence of sustained stretches of negative RDC values indicates that no persistent helical structure is present (Alexandrescu and Kammerer 2003; Fieber et al. 2004), which is consistent with circular dichroism data for all-Ala α -LA in urea solutions (Redfield et al. 1999).

Although the pattern of RDC values for the protein at different urea concentrations is similar overall, there are some significant differences. For instance in 6 M urea negative values are observed for Asp16 and Gly17. In 8 M urea the value for Asp16 is positive and in 10 M urea both residues have positive values. By contrast, negative RDC values are observed for Tyr 103, Trp 104 and Leu 105 in 6, 8 and 10 M urea. In 4 M urea, RDC values for the majority of α -domain residues could not be measured because the peaks are too broad. The correlation coefficients between the larger data sets are 0.70 (6 and 8 M urea), 0.64 (8 and 10 M urea) and 0.46 (6 and 10 M urea); this is considerably lower than found for staphylococcal nuclease in water and at different urea concentrations (Shortle and Ackerman 2001) or for ACBP at 2.5 and 4 M GuHCl (Fieber et al. 2004). This suggests that although the protein is largely unfolded at the higher urea concentrations used, significant changes take place in the structural ensemble of all-Ala α -LA as the urea concentration is increased from 4 to 10 M; this is consistent with the previous observation of stepwise non-cooperative unfolding of all-Ala α -LA in urea (Redfield et al. 1999).

Modelling RDCs for denatured all-Ala α -LA

Two all-atom models have been used to predict RDC values for completely unfolded proteins (Bernadó et al. 2005b; Jha et al. 2005). Here we have used an ensemble of 5,000 structures generated using the Jha et al. coil algorithm including nearest-neighbor effects (Jha et al. 2005)

and have calculated average predicted RDCs for this ensemble as described in the “Materials and methods” section. Deviations from these predicted values should indicate regions of residual structure within all-Ala α -LA. Figure 4b–e show comparisons of the experimental RDCs with those predicted from the statistical coil model. The correlation coefficients between experimental and calculated RDCs are 0.06, 0.17 and 0.25 at 6, 8 and 10 M urea, respectively. It is likely that residues with a negative experimental RDC value are involved in residual structure; after removing these values the correlation coefficients increase to 0.16, 0.32 and 0.50. The improvement of the correlation between the experimental and calculated values at higher urea concentration reflects the increasingly unfolded conformational ensemble for all-Ala α -LA. The correlation coefficient of 0.50 obtained for the 10 M urea RDC data is significantly lower than that reported by Jha et al. for apomyoglobin (0.71) and ubiquitin (0.70) but higher than the values for eglin c (0.30) and Δ 131 Δ staphylococcal nuclease (0.14) (Jha et al. 2005).

Differences between the experimental data and the predicted RDCs from the coil model can be broadly placed into two categories. The first of these includes residues 25–35 (the B helix in native α -LA) and the C-terminal residues (3₁₀-helix in native α -LA): the calculated RDC values deviate significantly from the experimental values measured in 6 M and to some extent in 8 M urea, but match those from 10 M urea well. This is accompanied by R_2 values in these parts of the sequence which are elevated in 4 M urea but decrease to the plateau value in 10 M urea. These residues are evidently involved in residual structure at lower urea concentration, but become unstructured at high urea concentrations. A second group is formed by residues 101–106 (D helix in native α -LA) which do not correlate well with the predicted RDC values at any urea concentration, and show somewhat elevated R_2 values even in 10 M urea. This indicates that these residues are to some extent structured, even at this high urea concentration.

Other discrepancies between the experimental and predicted RDCs can be attributed to certain amino acid types and sequence characteristics. Higher than expected RDCs are observed around the two proline residues (Pro24 and Pro67), for example, which suggests that the coil model for all-Ala α -LA does not sufficiently account for the way in which proline rigidifies the polypeptide chain. Wells et al. made a similar observation that their initial statistical coil model did not adequately model the conformational behaviour around proline residues (Wells et al. 2008). The low RDC values for the highly mobile GYGG sequence between residues 17 and 20, on the other hand, are well predicted by the model. Interestingly, the tryptophan residues (Trp60, Trp104 and Trp118) give rise to regions of lower RDC values than predicted and in many cases to

negative values. While sustained stretches of RDCs with the opposite sign have been shown to be characteristic of helical conformations, it has been shown that changes in sign spanning just a couple of residues can arise from turn conformations (Mukrasch et al. 2007). Model peptides containing residues 101–111 and 111–120 from α -LA were shown to adopt turn-like structures for residues 103–106 and residues 115–118, respectively (Demarest et al. 1999; Demarest and Raleigh 2000). Thus, Trp104 and Trp118 may be inducing turns in the all-Ala α -LA structure even in the presence of denaturant. To investigate the possibility of long-range contacts, different ensemble sub-sets of the random-coil ensemble were considered (Bernadó et al. 2005a), but no evidence of long-range contacts was found for all-Ala α -LA which is consistent with previous studies (Wirmer et al. 2006a).

Discussion

Previous studies have elucidated the pattern of non-cooperative unfolding of the α -LA molten globule as the concentration of urea is increased (Schulman et al. 1997). Chemical shifts were used to monitor this unfolding event by exploiting the observation that in the molten globule state the NMR resonances are very broad and cannot in general be resolved. This is due to fluctuations on a millisecond to microsecond time scale in the populated conformational ensemble. On unfolding, however, sharp well-resolved resonances appear in the NMR spectrum. The urea-induced unfolding is, therefore, followed by monitoring the appearance of these sharp resonances in the spectra as the concentration of urea is increased. The results of these studies for all-Ala α -LA show that the residues that are most resistant to urea-induced unfolding are those in the regions corresponding to helices A, B, C, D and the C-terminal 3_{10} helix of native α -LA (Redfield et al. 1999). The HSQC cross peaks of many of the residues in these regions are only observed when the protein is exposed to urea concentrations of 4 M or higher. The observed pattern of the appearance of peaks in HSQC spectra indicates that unfolding of the all-Ala α -LA molten globule in urea is a non-cooperative process.

Here we have extended the investigation of all-Ala α -LA by measuring ^{15}N relaxation data and RDC values at urea concentrations of 4, 6, 8 and 10 M. It is now possible to characterize the unfolding of all-Ala α -LA in greater detail and, in particular, to follow the unfolding of residual structure occurring at higher urea concentrations. Both the ^{15}N relaxation and RDC data show that the region corresponding to the native D helix unfolds last and is involved in non-random interactions even in 10 M urea. The native A and B helices are the next most stable parts of the

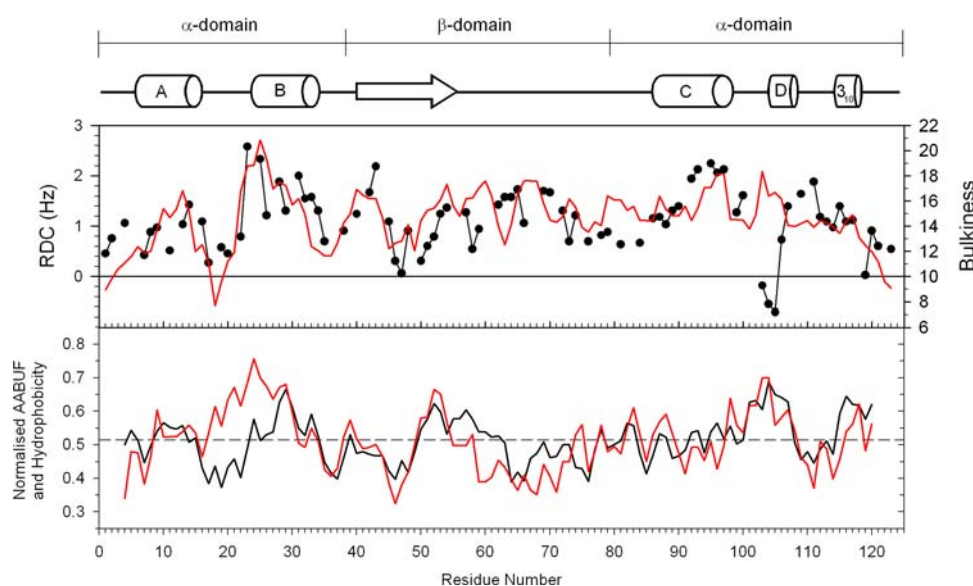
structure, with millisecond time scale motions present in 8 M urea as well as RDCs indicative of non-random coil conformations. The regions of all-Ala α -LA corresponding to the C and C-terminal 3_{10} helices of native α -LA show non-random interactions up to a urea concentration of 6 M.

Other evidence for persistent non-random structure in the C-terminal region of the protein comes from studies of short peptide fragments (residues 101–120) taken from the α -LA sequence (Demarest et al. 1998, 1999, 2001; Demarest and Raleigh 2000). These have identified two hydrophobic clusters involving residues 101–104 and 115–119, respectively. The cluster involving Ile101, Tyr103 and Trp104 has been shown to be intact in 8 M urea and based on the ^{15}N relaxation and RDC data we propose that this remains intact even in 10 M urea in all-Ala α -LA. The negative RDC values in this region are consistent with the formation of a turn-like structure by these residues (Mukrasch et al. 2007). It has been shown that ubiquitin forms a hairpin turn at urea concentrations as high as 8 M (Meier et al. 2007).

It has been a long-term goal to be able to predict non-random structure directly from the polypeptide sequence. It has been shown previously that amino-acid bulkiness averaged over a 5-residue window along the protein sequence is able to predict RDCs for unfolded polypeptides (Cho et al. 2007). The experimental RDC data for all-Ala α -LA in 10 M urea are compared with predictions from the bulkiness profile for the all-Ala α -LA sequence in Fig. 5a. A reasonable correlation is obtained for most regions of the sequence with the exception of residues 103–106. The trends in the bulkiness profile along the sequence are similar to those in the predicted RDC values from the statistical coil model although the bulkiness profile is smoother as the values are averaged over a five-residue window. The good agreement between the measured RDCs and the bulkiness profile is not surprising when high concentrations of denaturant are present, since the effect of electrostatic and hydrophobic forces within the polypeptide chain will be minimized by the denaturant. The only physicochemical property of the amino acids which remains a dominant factor is their relative size (or bulk).

As the concentration of denaturant is lowered, electrostatic and hydrophobic forces will start to come into play. In studies of apomyoglobin, the average area buried upon folding (AABUF) has been suggested to provide a good indication of those regions within the polypeptide chain which might be involved in the initial non-random interactions which then initiate the collapse and folding of the protein (Nishimura et al. 2005; Rose et al. 1985). An alternative parameter which has been investigated in this context is hydrophobicity (Wirmer et al. 2006a). Hydrophobicity scales generally disregard amino acid size; lysine and arginine, for instance, are assigned low values on the

Fig. 5 a Experimental RDCs recorded for all-Ala α -LA in 10 M urea (black) are compared with the bulkiness profile (red) calculated as described by Cho et al. (2007). **b** Normalised values for all Ala α -LA of the average area buried upon unfolding (AABUF) (black) (Rose et al. 1985) are compared with the hydrophobicity using the Abraham & Leo scale (red) (Abraham and Leo 1987). All values were averages over 7-residue windows. The dashed grey line shows the mean value of AABUF and hydrophobicity



basis that they are charged. Significantly higher values are attributed to amino acids such as alanine, which although small is entirely hydrophobic (Abraham and Leo 1987). AABUF, on the other hand, takes size into account and assigns lower values to Ala than Arg or Lys (Rose et al. 1985).

The AABUF and hydrophobicity profiles for all-Ala α -LA are compared in Fig. 5b; these two parameters show similar trends in most regions of the sequence. The regions corresponding to the native B, D and C-terminal 3_{10} helices, which unfold at the highest urea concentrations, are predicted well by both parameters. The R_2 data and ^{15}N chemical shift differences between 6 and 10 M urea in the region of residues 15–25 indicate significant backbone mobility and a lack of residual structure, particularly for residues 17–20; this is better reflected by the AABUF than the hydrophobicity values.

Surprisingly, comparison of the experimental data and the AABUF and hydrophobicity values for the A-helix region and the first half of the β -domain shows less agreement. The experimental data suggest that the A-helix region is involved in residual structure at urea concentrations of 6–8 M urea, but the AABUF and hydrophobicity values in this region are only slightly larger than the average values across the whole sequence. It may be the high degree of helical propensity rather than the hydrophobicity in this region of the sequence which is driving the formation of residual structure. The experimental R_2 values and ^{15}N chemical shifts suggest that the β -domain is largely unfolded at urea concentrations of 6 M and higher. Reasonably good agreement is observed between the experimental RDCs and both the values predicted from the statistical coil model (Fig. 4b–e) and

the bulkiness profile (Fig. 5a) confirming that the β -domain is largely unfolded in 6 M urea. The AABUF and hydrophobicity profiles on the other hand show elevated values within the β -domain, in particular for residues 50–60. The higher values for residues $\sim$ 50–60 compared to residues $\sim$ 61–70 mirror the slightly elevated R_2 values which are observed for residues 40–60 compared to residues 61–80 in 4 M urea and correlate with a recent study of backbone dynamics of the all-Ala α -LA molten globule that has shown more restricted mobility for residues 40–60 than for 61–80 (Ramboarina and Redfield 2008). This suggests that some non-random behavior may be present in the β -domain at lower urea concentrations but that this structure does not persist at higher urea concentrations as it does in the α -domain. The correlation between measured NMR parameters and the AABUF profile although reasonable for all-Ala α -LA, does not appear to be as clear as that reported previously for apomyoglobin (Nishimura et al. 2005).

In conclusion, the measurement of ^{15}N relaxation data and RDCs for all-Ala α -LA at several concentrations of urea has provided a more detailed understanding of the stepwise non-cooperative unfolding of this protein at higher urea concentrations. This has identified the presence of non-random interactions at urea concentrations as high as 10 M which arise from conformational sampling of turn structures. The presence of such transient structure was not evident from the previous HSQC-based urea titration study which focused largely on the loss of stable secondary structure and long-range hydrophobic interactions in the molten globule which gave rise to significant peak broadening at lower urea concentrations (Redfield et al. 1999). The application of both approaches can provide a more

complete picture of the loss of structure that occurs as a molten globule unfolds in urea.

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