Stabilization of small increments of β-sheet in aqueous solution is critical for fundamental research on protein folding preferences8–11 and for development of biologically active small molecules.8–11 Two different strategies for promoting double-stranded antiparallel β-sheet are found among natural and synthetic peptides: side-chain linkage, usually involving cysteine residues,10,11 and backbone linkage.12 Here, we report a direct comparison between side-chain disulfide cyclization and backbone cyclization for generation of reference compounds that can be used to determine β-sheet populations of flexible peptides. NMR data suggest that both cyclization modes in our peptide system stabilize a very highly populated native-like antiparallel β-sheet conformation, although backbone cyclization appears to confer somewhat greater conformational stability. These reference compounds allow quantitative evaluation of sequence-stability relationships among linear peptides that adopt double-stranded β-sheet (“β-hairpin”) conformations in aqueous solution.

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Cyclic peptides 1 and 2 (Figure 1) are potential models for the fully folded state of linear peptide 3. Both 1 and 2 contain the 12-residue segment of 3, with two additional residues to allow ring closure. Our previous studies of 3 and related linear peptides have shown that the d-Pro-Gly segment is a very strong inducer of β-sheet formation in adjacent strands,13,14 and we therefore used a second d-Pro-Gly segment to achieve backbone cyclization in 1.15 For 2, the disulfide-linked cysteines were placed at a “non-hydrogen-bonded” pairing because this juxtaposition is common among proteins.14 We have previously concluded from NMR data that a single β-sheet conformation is significantly populated for linear peptide 3 in aqueous solution.21 Like other short peptides, however, 3 equilibrates rapidly on the NMR time scale between the unfolded and β-sheet conformations. A high population in 1 or 2 of a β-sheet conformation analogous to that observed for 3 would allow determination of the β-sheet population of 3 from conformationally averaged NMR data.15

Two-dimensional NMR data for 1 and 2 in aqueous solution revealed numerous NOEs between residues that are not adjacent in sequence.16 The NMR data for 1 and 2 were used for NOE-restrained structure determinations with the program DYANA (Figure 1).17 For each peptide, all NOEs were consistent with a single family of conformations in which the two five-residue strand segments, Arg-2 to Val-6 and Orn-9 to Gln-13, display a high population in β-sheet conformations. A high population in Figure 2 of a β-sheet conformation analogous to that observed for 3 would allow determination of the β-sheet population of 3 from conformationally averaged NMR data.15

Figure 1. Comparison of NOE-restrained dynamics simulations (DYANA17) for cyclic peptides 1 and 2. The 20 best structures were compared for each peptide; RMSD among the 20 structures (backbone only) was 0.75 ± 0.23 Å for 1 and 1.40 ± 0.39 Å for 2. The 20-structure average for 1 showed a 1.20 Å RMSD relative to the 20-structure average for 2, indicating that the two cyclization modes promote similar conformations. Amidic H/D exchange data (100 mM sodium deuteroacetate buffer, pH 3.8 (uncorrected), 4°C) provided further evidence that cyclic peptides adopt a regular β-sheet conformation; the exchange data also suggested that backbone cyclization induces a somewhat more stable folded state than does disulfide cyclization. Retardation of amide H/D exchange is generally interpreted to indicate a steric and/or electronic impediment to chemical catalysis of exchange, e.g., from intramolecular hydrogen bonding.18 For 1, there was a regular alternation of pseudo-first-order rate constants along the strands: each residue expected to orient its N–H away from the opposite strand showed a rate constant >10–3 min–1, and each residue expected to be engaged in interstrand hydrogen...
bonding showed a rate constant $< 10^{-3}$ min$^{-1}$ (for two residues the rate constant was $< 10^{-5}$ min$^{-1}$). For 2, a similar distinction was seen, but the differences among rate constants were less pronounced for 2 than for 1. The backbone amide protons of Arg-2 and Gln-13 of 2 exchanged relatively rapidly, suggesting that the disulfide linkage is less effective than a β-sheet linkage at stabilizing local β-sheet structure.

The conformational rigidity implied for 1 by the H/D exchange data suggests that this cyclic peptide is an excellent reference compound for quantitative NMR-based determination of β-sheet populations in flexible peptides such as 3–7. We analyze the flexible peptides in terms of a two-state conformational equilibrium, unfolded (U) vs folded (F), where the latter is the β-sheet. $\Delta G^\circ$ can be estimated from any conformationally sensitive proton chemical shift via eq 1, where $\delta_{\text{obs}}$ is the chemical shift of the proton in question within the equilibrating peptide, $\delta_U$ is the chemical shift of that proton in the fully unfolded state, and $\delta_F$ is the chemical shift of that proton in the fully folded state. We have previously shown that 8, the β-Pro diastereomer of 3, is completely unfolded in water; therefore, we obtained $\delta_U$ values from 8. Results presented above show that 1 is a good source of $\delta_F$ values.

Protons on the α-carbons of amino acid residues are very sensitive to backbone conformation; residues in β-sheets display downfield shifts relative to the random coil state. We used Val-3, Orn-8 (or Lys-8), and Ile-10 to estimate $K_{\text{eq}}$ for β-sheet formation by linear peptides 3–7 in aqueous solution, based on eq 1. These three residues were selected because all occur at hydrogen-bonded positions, which means that $H_\text{B}$ is oriented away from the opposite strand in the folded state. This $H_\text{B}$ orientation should minimize the effect of any differences in side-chain packing between 1 and the folded state of the linear peptides (because of Tyr ring current effects, such packing differences could influence $\delta_{\text{obs}}$ values). $\Delta G^\circ$ for β-sheet formation was calculated from $K_{\text{eq}}$ and $\Delta G^\circ$ values (Table 1) were calculated for 4–7 by comparison with n-proline peptide 3 (systematic error is presumably minimized in the $\Delta G^\circ$ values).

The reasonably good agreement among the β-sheet population values deduced for Val-3, Orn-8 (or Lys-8), and Ile-10 of 3, 4, and 7 (Table 1) supports our hypothesis that these peptides display a two-state folding equilibrium. The small variations in population values within each peptide might indicate real differences in the folded population at each of the indicator residues or systematic error arising from the reference peptides. In particular, it is possible that cyclic peptide 1 and the folded state of 3 have different ensembles of side-chain orientations, even though 1 is an excellent model for the backbone of 3 in the folded state.

### Table 1

<table>
<thead>
<tr>
<th></th>
<th>Val-3</th>
<th>Orn-8</th>
<th>Ile-10</th>
<th>Pop. (%) of the β-Hairpin State for Each Linear Peptide and $\Delta G^\circ$ Values (kcal/mol) for Each Mutant Relative to Peptide 3$^a$</th>
<th>$\Delta G^\circ$</th>
<th>$\Delta G^\circ$ av $\Delta G^\circ$</th>
</tr>
</thead>
<tbody>
<tr>
<td>peptide</td>
<td>pop. (+2)</td>
<td>$\Delta G^\circ$</td>
<td>pop. (+3)</td>
<td>$\Delta G^\circ$</td>
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</tr>
<tr>
<td>3</td>
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<tr>
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<td>62</td>
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<td>59</td>
<td>+0.01</td>
</tr>
</tbody>
</table>

$^a$ The population of the β-hairpin state for each linear peptide was calculated at each indicator residue from $\delta_{\text{obs}}$ data as $\delta_{\text{obs}} - \delta_U/(\delta_F - \delta_U) \times 100$. $\Delta G^\circ$ for each mutant relative to peptide 3 was calculated in the usual way ($K_{\text{eq}}$ for β-hairpin formation from eq 1; $\Delta G^\circ = -RT \ln K_{\text{eq}}$). $\Delta G^\circ$ values for 3–6 reflect error propagation from the ±0.01 ppm uncertainty in the $\delta_{\text{obs}}$ measurements. In peptide 7, residue 8 is lysine rather than ornithine.

Communications to the Editor


(20) Val-5 also occurs in a hydrogen-bonded position but cannot be used because this residue is followed by proline in reference peptides 1 and 8 but not in equilibrating peptides 4–6. Proline profoundly influences the backbone conformation of the preceding residue: Wilmot, C. M.; Thornton, J. J. Mol. Biol. 1988, 205, 221.


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Supporting Information Available: Resonance assignments and NOEs for peptides 1 and 2 (PDF). This material is available free of charge via the Internet at http://pubs.acs.org.