

Predicting the conformational preferences of proteins using a physics-based free energy method

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Abstract

The calculation of free energy differences is of central importance in the simulation of biochemical systems. It is particularly difficult to calculate between pairs of macromolecular conformations as well as a computationally expensive task with existing methods. In this work, the confinement approach is used to calculate absolute free energies of biomolecular systems. This method provides two main advantages: it does not require a reaction coordinate or transition path and it is fast to compute. The calculated free energy can be decomposed into a per residue contribution in an approximate way. Per residue free energy allows us to identify the reason behind conformational preferences in biomolecules. Through out the article we show its use in different challenging modeling problems. In particular, we show its use in predicting the conformational preference of chameleon sequences (sequences with high sequence identity and different folds). This sequence dependent conformational preferences and per residue free energy decomposition set the stage for the use of this method in protein design.

Introduction

In some problems of protein science, you want to know the relative stability of a protein's conformation A compared to its conformation B . We call this the *difference free energy*. For example, in all allosteric mechanisms, a protein often adopts one conformation when a ligand is bound and another conformation when no ligand is bound¹. To understand the mechanism requires knowledge of the relative free energies of the protein conformations. Another example is in protein structure prediction, for example as practiced in the Critical Assessment of Structure Prediction (CASP) competition². If you are using some computational model and have predicted two putative native structures, A and B , you want to compute which has the lower free energy, in order to know which is the more native-like. In a third example, you may want to know whether mutating a few amino acids in a protein could cause the protein to switch from one stable conformation to another, because that could have important consequences for biological mechanisms and disease. And fourth, sometimes binding a ligand induces a protein from one conformation to another. A quantitative understanding of such *induced-fit* situations requires knowledge of the difference free energy of the protein in the two states. In all these cases, it would be useful to have a computational method that can be used efficiently with atomically detailed physical forcefield models to compute the free energy differences between different given protein conformations.

One widely explored strategy is to use molecular dynamics simulations along some putative reaction coordinate pathway from conformation A to B ³⁻¹⁰. The free energy along this reaction coordinate can then be determined using methods such as umbrella sampling^{11,12} and weighted histogram analysis method (WHAM)¹³. Such approaches have several limitations. First, it is necessary to know an efficient reaction pathway from A to B . If conformations A and B are quite different, then it can be challenging to find such paths. Second, these methods are computationally slow. To get an accurate estimate of the total free energy difference $\Delta G = G_B - G_A$ requires accurate determinations of the many small free energy differences $A \rightarrow 1 \rightarrow 2 \dots \rightarrow B$, so each step requires substantial amounts of sampling. And third, these methods are prone to large errors, because the pathway error is a sum of many errors among the many steps.

The calculation of protein conformational free energy has been successfully attempted by a number of groups¹⁴⁻²³. Some of these methods like the reference system method¹⁵, deactivated morphing¹⁶, the confinement method¹⁹⁻²¹ take an alternative strategy for computing difference free energies that does not require knowing a pathway from *A* to *B*.

Here, we adopt the confinement method of Tyka et al¹⁹ and Cecchini et al.²⁰, which is based on the thermodynamic cycle shown in Figure 1. We start with an ensemble of related conformations, *A*. We then restrict *A* to a much “tighter” ensemble *A** by applying position restraints in a series of MD simulations. Similarly, a second ensemble *B* is restricted to *B**. We then compute the free energy between *A** and *B** using either normal mode analysis^{24,25} or the quasiharmonic method^{26,27}. This step takes into account both the remaining conformational entropy in each ensemble and the remaining enthalpy difference. In this way, we can calculate the free energy difference between the two end states without needing to define a physical path or reaction coordinate connecting the two states. The confinement method shares some similarities with the ‘confine-and-release’ method for computing ligand binding affinities²⁸⁻³⁰.

[Figure 1 about here.]

Previously, the confinement method has been validated on small model peptides^{19,20}. Here, we show that the confinement method can also be applied successfully to larger systems, such as the series of chameleon proteins designed by Orban and co-workers³¹⁻³⁵, which can switch between two completely different folds with only slight changes in sequence. We also show that our computed difference free energies are useful in evaluating the quality of CASP target protein predictions. This could be of significant value for ultimately improving the energetics in protein-structure models, not just the structures. And, finally, we show that we can approximately decompose difference free energies into individual amino acid components. This offers the opportunity for diagnosing the structural basis for difference free energies, which can be useful for interpreting biological mechanisms.

Results and Discussion

The confinement method succeeds at some basic consistency checks

First, we validated that our implementation of the confinement method produces results compatible with previous calculations reported in the literature. The method has previously been applied to a 16 amino acid residue β -hairpin from protein G, known as BHP²⁰. We calculated the free energy difference between the native conformation—called bhp1, which has a two-stranded β -sheet—and a non-native conformation—called bhp3, which has a three-stranded β -sheet. Our confinement calculation shows that bhp1 is more stable by 1.7 kcal/mol, consistent with 4 μ s equilibrium simulations showing that bhp1 is more favorable configuration by 1.8 kcal/mol, and in agreement with previous calculations²⁰.

Second, we looked at 6 target proteins from the CASP9 experiment². For each target, we examined up to 5 submitted models. We computed the difference free energy between the true native and the best model. Figure 4 and Table 1 show that, in 5 out of 6 cases, the confinement method assigns a lower free energy to the experimentally determined structure than to any of the decoys. Other well-known discriminators can also successfully tell native structures from computer generated models^{36;37}; it is just an independent useful validation that the confinement calculations make sense.

[Table 1 about here.]

Should we comment on T0560? Why is it giving such a high value with such high gdt? It's not the point to make here, but it is what jumps at me. I think we have made it clear that crystal structures are often more stable due to salt-bridge effects. Also, we have shown in other cases that sidechains orientation may play important roles inspite of close GDT-TS. You will see free energy difference between two computer generated model is not so high. So, I believe sidechain may play some role here.

The confinement method correctly predicts the structures of chameleon sequences

We also tested the confinement method predictions for difference free energies on the chameleon sequences of Orban et al.³¹⁻³⁵. These are instances in which two highly similar sequences fold into remarkably different structures. Orban and co-workers have designed a protein-G-like sequence of 56-residues that is marginally stable in one of two possible folds. By mutating key residues in this sequence they are able to stabilize one fold or the other (see Figure 2). We refer to the $4\beta + \alpha$ structure as the β conformation, and we refer to the 3α structure as the α conformation. We denote sequences that prefer the α fold as GA and sequences that prefer β as GB. One pair of sequences (GA88/GB88) is 88 percent identical in sequence, differing in seven positions. Another pair (GA95/GB95) is 95 percent identical, differing in three positions. Accurately predicting the structural preferences of these structures presents a serious challenge for computational methods³⁸.

[Figure 2 about here.]

We initially approached this problem by making a model of each sequence with the same backbone structure as its partner chameleon sequence. For example, we took the sequence of GA88 and built a model with the same overall structure as GB88. We then used the confinement method to assess the free energy difference between the experimentally determined structure of GA88 and the model (with the GA88 sequence and the GB88 structure). The confinement method was able to predict the conformational preferences correctly for all four sequences (data not shown). This is, however, not surprising as it is often easy to distinguish computational models from native structures^{36,39}. To avoid this potential problem, we instead computed relative free energy of two different computer generated models for each sequence. One model is based on the α structure and the other on the β structure (see Supporting Information for details on the modeling procedure). This is a much more realistic test of the confinement method's ability to accurately calculate the difference free energy.

Figure 2 shows that the confinement method identifies the correct structure for all four sequences. One hypothesis is that such structural transitions re-

quire states with diminished stability³⁴. It is believed that if the free energy of the native state and the alternative state are within a range of around 5 kcal/mol, then it is possible for the native state to be destabilized relative to the alternative state with only small changes in sequence³²⁻³⁴. The calculated free energy differences range from around 3.5 to 5.0 kcal/mol, which is consistent with this hypothesis. In a more recent study³⁵, the amino acid residue at position 45 (Tyr for β and Leu for α) was found to be important for switching between α and β conformation. This inspired us to introduce another mutation at this position, Y45A, which we refer to as GA98. Our calculations predict that this mutation shifts the equilibrium to the α conformation, which is now more stable than the β conformation by 3.8 kcal/mol. Although this result has not yet been confirmed experimentally, it is consistent with the previously observed effect of Y45L.

Per-residue free energy calculations can identify the mechanistic detail behind conformational preferences

To better understand the mechanism behind these chameleon proteins, we decomposed the calculated free energy into per-residue contributions in an approximate way¹⁹. We decompose each confinement step ($\Delta G_{A,A^*}$, $\Delta G_{B,B^*}$) into per-residue contributions. We can also decompose the remaining enthalpy in the confined ensemble into per-residue contributions. However, the method is approximate because we do not include the residual conformational entropy from the normal mode or quasiharmonic analysis steps. Computing per-residue contributions helps us to identify important residues that stabilize a particular conformation. The per residue free energy, $\Delta\Delta G(\beta - \alpha)$ is shown in Figure 3.

[Figure 3 about here.]

Although the overall free energy difference between the two structures is small (within 3–5 kcal/mol), individual residues can show marked preferences for being in either α or β conformation. Such differences can be understood by looking in detail at the local environment for those residues. For example, the region around residue 7 forms a random coil in the α structure, whereas it is forming a beta sheet structure in the β structure. These

residues strongly favour the locally well packed and hydrogen bonded environment found in the β sheet. Overall, favoring α or β structure is a delicate balance, where the relative global free energy difference is small and the contributions of different per residue tendencies balance out. It is therefore very likely that by changing key residue the global fold preferences can be changed.

Three residues are mutated between GA95 and GB95, at positions 20, 30, and 45. In GA95, these residues (L20, I30 and L45) stabilize the α structure (compare the upper and lower panels on Figure 3). In GB95, two of these residues (F30 and Y45) favor the β structure, because they have large solvent exposed surface areas in the α structure, but are more buried in the β structure. Additionally Y45 forms a hydrogen bond with D47 in the β structure. On the other hand, residue A20 from GB95 still favors the α structure, although less strongly than L20 in GA95.

The α and β sequences are nearly identical, so there are some common features observed for all sequences. The experimental observations classified the protein into two parts: Amino acids 9–51 are fully structured in both folds, whereas residues 1–8 and 52–56 are unstructured in the α fold, but form β -strands in the β fold. Most of the amino acid residues in the region 1–9 have negative per-residue free energies, which means that these residues favor the β structure. Roles of some other important residues in stabilizing either the α or β conformation are summarized in Figure S1.

In addition to the direct effects of the mutation, there are also indirect effects due to small perturbations in the environment around the mutations. For example, the L20A mutation causes a slight repacking around residue 20. This causes large changes in the per-residue free energies of nearby residues T25 and A26. However, the changes for these two residues have opposite signs and nearly cancel.

These per-residue free energy decompositions provide a great deal of insight into the driving forces behind protein folding and conformational change. We believe that such calculations may also be useful for protein design—designing proteins with specific structures and functions.

The confinement method is a useful tool for structure prediction

We have tested the ability of the confinement method to act as a “meta-predictor” for structure prediction. Here, the task is to correctly identify the most accurate models out of a set of “decoys” generated by different methods during the CASP experiment. CASP is a blind test in which different groups apply methods to predict the 3-dimensional structure of proteins from their sequences. Each group is allowed to submit five possible structures, which they are supposed to rank from best to worst.

We have performed two experiments centered on CASP. In the first experiment, we tried to rank-order predictions for several targets. For each target, the predictions were either produced by a single group—presumably using the same method for each prediction, or were produced by several different groups—using different methods. The goal of this experiment is to determine if the physics-based confinement method can correctly identify more native-like structures as having lower free energies. Our second experiment was to see if the confinement method can identify structures that are missed by other meta-prediction servers. Most successful meta-prediction servers are based on the idea of consensus: if many different prediction methods produce similar results, then that is probably a correct prediction^{40;41}. This is often a powerful heuristic, but it can miss cases where there is a very good result that is only predicted by one method. We chose several cases where such structures were missed by the best meta-predictors in CASP and assessed if the confinement method can correctly identify these accurate models.

As is common in the CASP experiment, we assess our results in terms of Global Distance Test Total Score (GDT-TS)⁴², which is a C α based measure of structural accuracy. It can be understood roughly as the percentage of residues that are correctly positioned in the model (range 0 to 100, higher is better). We did not have enough computer power to analyze every model, so the initial models for the test were chosen from a selection of different server groups that have done well in past CASP events.

Overall, the confinement method performs well on our CASP tests. Figure 4 shows that in almost every case, the native structure has the lowest free energy and the best model has the next lowest free energy. The confinement

method appears to be a useful tool for ranking structure predictions.

[Figure 4 about here.]

The confinement method can correctly rank-order structure predictions

First, we examined the ability of the confinement method to rank different models for a target that has been generated by a single group using the same methodology for all predictions. We examined two targets: T0559 and T0560 (see Table 2).

[Table 2 about here.]

The first test case is target T0559. The best predictor group for this 69 amino acid target was “BAKER-ROSETTASERVER”. To save computer time, we excluded two models that were very similar to other models that we did include. For this target, the difference free energy computed by the confinement method can be used to accurately rank-order all of the models and the native experimental structure (Figure 5).

[Figure 5 about here.]

We performed a similar calculation for target T0560 with two models from the group “Splicer”. The remaining three models were discarded as they were too similar to the rest of the models. Again, we can correctly identify the native state and our calculated free energy based ranking matches well with GDT-TS (Figures S2 in the supporting information).

Next, we tested the ability of the confinement method to rank models for target T0540 that were produced by different prediction groups. The top models from groups “LTB” (Model 1) and “Mufold” (Model 2) were chosen for analysis. Once again, the free energy based ranking correlates well with the GDT-TS based score (Figure 6).

[Figure 6 about here.]

if the drop down bars/point are a different color it will be easier to see when the crystal is on top of the axis I will wait for Sarina's figure. I have changed the caption of figure 5 to say why some region are colored brown.

The per-residue free energy is sensitive to small changes in protein conformation:

In a previous section, we discussed how the per-residue free energy can reveal mechanistic detail behind the conformational preference of a chameleon sequence. In this section, our aim is to apply the same method and try to understand if the per-residue free energy can help us identify residues which stabilize or destabilize a particular region of a protein. In this case, the two conformations have similar folds with only small changes in localized regions. We chose CASP target T0569 and compared the experimental NMR structure with the best predicted model (GDT-TS=78; predicted by the "Mufold" group). The confinement method predicts that the experimental structure is more stable by 20 kcal/mol (Figure S2).

The confinement method clearly identifies two hydrophobic residues V59 and I61, which destabilize the predicted model with respect to the experimental structure (Figure 7 and Figure S3). The side chains of these hydrophobic residues are oriented towards the protein hydrophobic core in the native NMR structure but oriented towards the exterior of the protein and are solvent exposed in the model. These residues are part of a beta-sheet in the experimental structure, but because of their side chain orientation the corresponding beta-sheet becomes disordered in the predicted model (Figure 7 and Figure S4). There is also a large difference around K76, which forms a salt-bridge with D11 in the predicted model, but not in the experimental structure. This suggests that salt-bridge interactions are too favorable for the combination of force field and implicit solvent model we use, which has been a problem noted in the past⁴⁵.

[Figure 7 about here.]

The confinement method occasionally produces incorrect results:

Despite success for most of the studied systems, there are few failures, specially when the GDT-TS scores of the compared structures are very close. One example is Target T0538, where we compared the experimental structure with three models (Model 1: "PconsR"—GDT-TS=96; Model 2: "Shell"—GDT-TS=90; Model 3: "FOLDIT"—GDT-TS=86). Contrary to our expectation, the confinement method predicts that computer generated Model 1 is more stable than the crystal structure (Figure S5). Per-residue free energy calculations (not shown) show that despite only small variations at the backbone level, the side chains are oriented in very different ways (Figure 8), giving rise to large differences in the stabilization of certain residues. In particular, some of the differences arise from different salt bridge patterns and certain flexible polar residues exposed to the surface. This unexpected result shows that the confinement method is very sensitive to local interactions (including side chain reorientation) and may indicate issues with the forcefield and implicit solvent models used in the calculation.

[Figure 8 about here.]

Can the confinement method perform better quality assessment in protein structure prediction?

A part of the CASP experiment is dedicated to the quality assessment (QA) of predicted models⁴⁰. Here, predictors were asked to score each model on a scale (known as qmode) from 0 to 1, with higher values corresponding to better models⁴⁰. Most of the groups use consensus strategies in such experiments. Here, we chose two computer generated models from CASP target T0538, where the top performing group "MUFOLD-WQA"⁴¹ failed. One of the models was from "PconsR" (GDT-TS = 83, qmode 1 = 0.5865) and another from "MULTICOM-NOVEL" (GDT-TS = 83, qmode 1 = 0.5865). We choose these two models as the model by PconsR was the most accurate predicted model for this target, whereas the other model was predicted best by QA test of MUFOLD-WQA. The calculated free energy

using confinement method indicate that the model by PconsR is more stable by 3.9Kcal/Mol which supports the GDT_TS trend. Although consensus prediction strategies do well specially when dealing with large numbers of structures, they can miss a good answer that is only predicted by a few methods. At least in this case, the confinement method is able to capture this structure that was otherwise missed.

Conclusion

We have described a computational method called confinement for computing the difference free energy between two conformational ensembles. We show that the difference free energy can be calculated on 100 residue sized proteins, even for large conformational changes. We have demonstrated that it can discriminate the folding preferences of a series of chameleon proteins. We show that the confinement method can discriminate between the native structure and structure predictions and can identify the best prediction reliably. Perhaps most importantly, we have shown that it can be used to give residue-level insights into what are the dominant structural factors in a protein that are responsible for the difference free energies. The confinement should be useful for protein design, structure prediction, and understanding the mechanism of conformational change.

A key advantage is that this method does not require any reaction coordinate, or sampling a pathway from conformation *A* to *B*.

1 Method

The confinement method has been described in details by Tyka et al. in reference¹⁹ and Cecchini et al.²⁰. The basic approach of the confinement approach is the same in both these papers. A thermodynamic cycle is used to compute the free energy between conformations *A* and *B*. However, there are some small technical differences between the two approaches. Here we briefly describe the procedure that we used.

1. In the first step, minimizations of *A* and *B* conformations are per-

formed. These minimized conformations (A^* and B^*) are the reference conformation of that state.

2. The free energy of confining the ensemble (A or B) to a tighter ensemble (A^* or B^*) is calculated. This is done by gradually confining each atom to its position in the reference conformation (A^* or B^*) using a series of progressively stronger position restraints. This is done by running 21 molecular dynamics simulation (each 20 ns long) for each leg of the thermodynamic cycle, where the harmonic restraint force constant was scaled from 0.00005 Kcal/Mol (mostly free) to 81.92 kcal/mol (tightly restrained). The free energy for this step is estimated using a thermodynamic integration approach developed by Tyka et. al.¹⁹. The confinement free energies calculated in this step are denoted as $\Delta G_{A,A^*}$ and $\Delta G_{B,B^*}$ in Figure 1.
3. The thermodynamic cycle is closed by calculating the free energy between the final restrained states A^* and B^* using normal mode analysis^{24;25} or quasiharmonic analysis^{26;27}. The free energy calculated in this way is shown as $\Delta G_{A^*,B^*}$ in Figure 1.
4. The full free energy, $\Delta G_{A,B}$ between the two state A and B is calculated as $\Delta G_{A,B} = \Delta G_{A,A^*} - \Delta G_{B,B^*} + \Delta G_{A^*,B^*}$.

One advantage of confinement method is that none of the simulations during the restraining step depends on each other. Therefore, with available computer resources it can be fast to compute. The computational cost for a single 20 ns simulation (along the confinement step) of 56-residue protein is only 4 hours in a single gpu computer [all the different confinements in one GPU?](#). All calculations were performed with the Amber 11 suit of programs^{43;44} in combination with the ff99SB forcefield⁴⁷ and the generalized born implicit solvent⁴⁸.

Interestingly, we extend the method for calculation of per residue free energy in an approximate way. For this purpose, the confinement energy, $\Delta G_{A,A^*}$ and $\Delta G_{B,B^*}$ of each residue is calculated in the usual numerical way as described by Tyka et al. in reference¹⁹. The internal energy of each residue is calculated using amber's "decomp" module using the final restrained trajectory. We call this method approximate as we ignore the entropic contribution from the normal mode or quasiharmonic analysis.

However, this contribution to the total free energy is much smaller, which allow us to study the mechanistic details of conformational preference of each residue.

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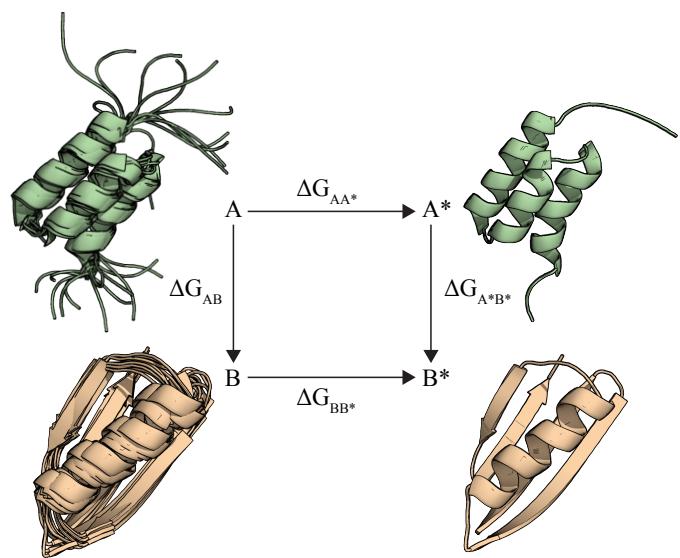
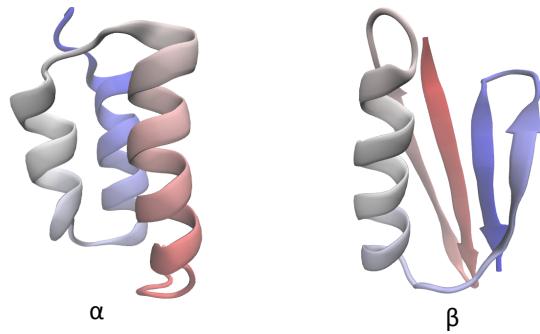


Figure 1: Graphical representation of the thermodynamic cycle involving confinement method.

12345678901234567890123456789012345678901234567890123456
 2jws: TTYKLILNLKQAKEEAIKEVDA**G**I**A**E**K**Y**I**KL**I**ANAKTVEGVWTL**L**K**D**E**I**L**T**FT**V**T**E** GA88
 2jwu: TTYKLILNLKQAKEEAIKELVDA**A**T**A**E**K**Y**F**K**L****Y**ANAKTVEGVW**T****Y**K**D**E**T****K**T**F**T**V**T**E** GB88
 2kdl: TTYKLILNLKQAKEEAIKE**L**VDAGTA**E**K**I**KL**I**ANAKTVEGVW**T****L**K**D**E**I**K**T**F**T**V**T****E** GA95
 2kdm: TTYKLILNLKQAKEEAIKE**A**VDAGTA**E**K**F**K**L**IANAKTVEGVW**T****Y**K**D**E**I**K**T**F**T**V**T****E** GB95



Sequence	Experimental Fold		Calculated Fold		Calculated $\Delta G_{\beta} - \Delta G_{\alpha}$ (Kcal/Mol)
	α	β	α	β	
GA88	✓		✓		+3.94±0.51
GB88		✓		✓	-4.36±0.46
GA95	✓		✓		+3.48±0.47
GB95		✓		✓	-5.01±0.49

Figure 2: The confinement method correctly predicts the structural preferences of four chameleon sequences. The top part of the figure represent four sequences used in this study along with the protein data bank identifier. The experimentally observed fold, computationally predicted fold, and difference free energy between the two folds is reported for each sequence.

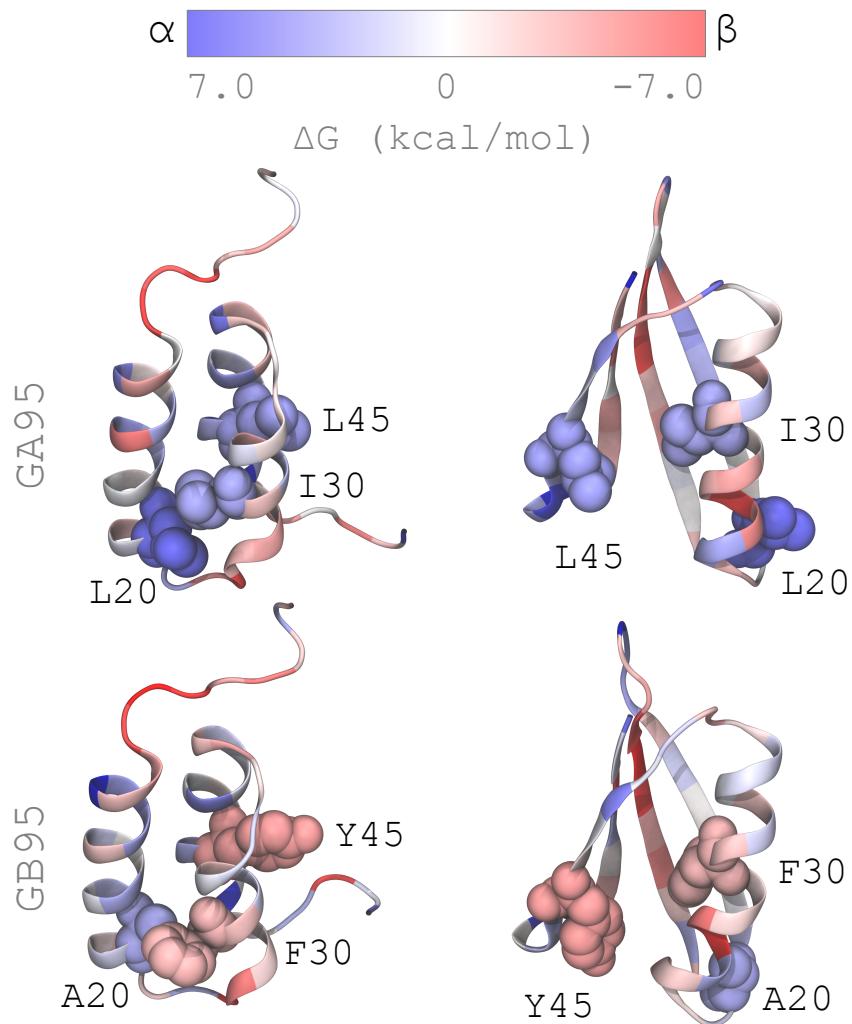


Figure 3: Per-residue difference free energies for the α and β conformations of the GA95 and GB95 sequences. Residues colored in blue favor the α structure, residues in red favor the β structure, and white residues have no preference.

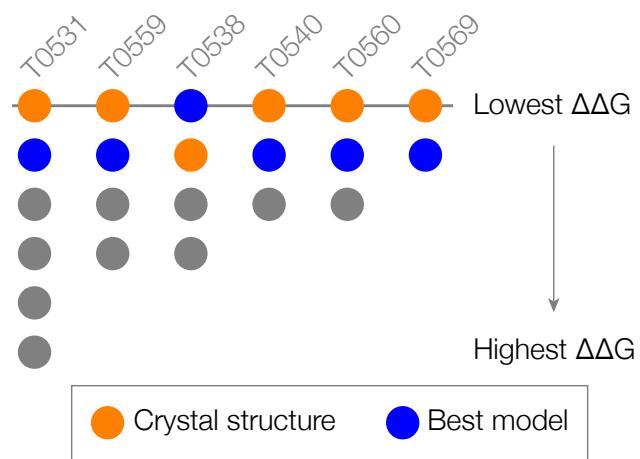


Figure 4: The confinement method is usually able to identify the native structure and the best model (the model with the highest GDT-TS) from a set of decoys.

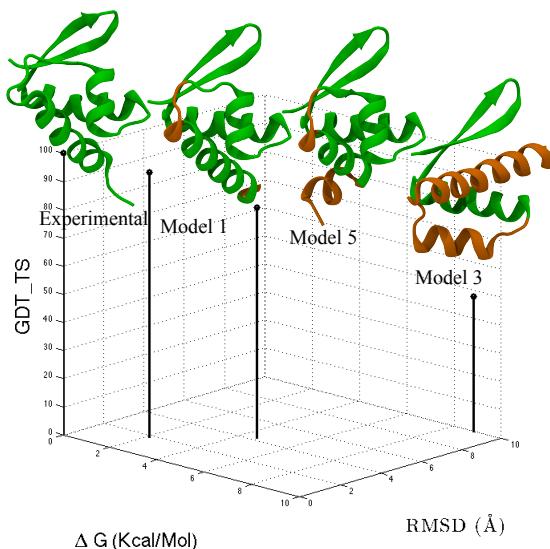


Figure 5: The confinement method can correctly rank the native structure and three predictions produced by a single method for target T0559. The backbone region of the predicted structures which differ substantially compare to the experimental structure upon superposition are colored brown.

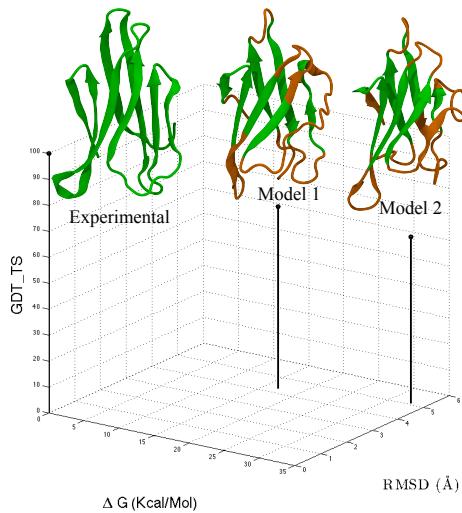


Figure 6: The confinement method can correctly rank the native structure and two models submitted by different prediction groups.

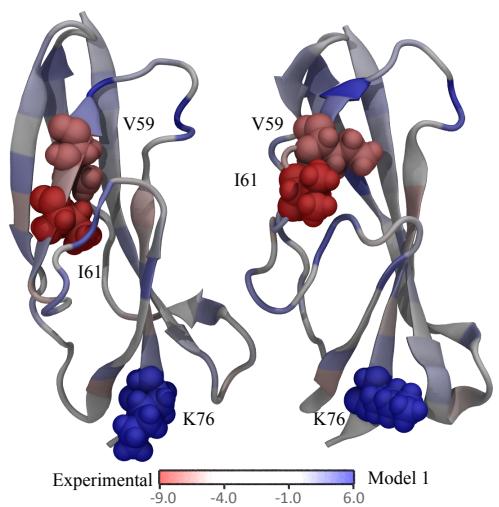


Figure 7: Per-residue difference free energy between the experimental NMR structure and the best prediction for CASP target T0569. The amino acid residues that are colored in deep red and deep blue stabilizes the NMR structure and the prediction, respectively; the residues with light blue color does not have a strong preference.

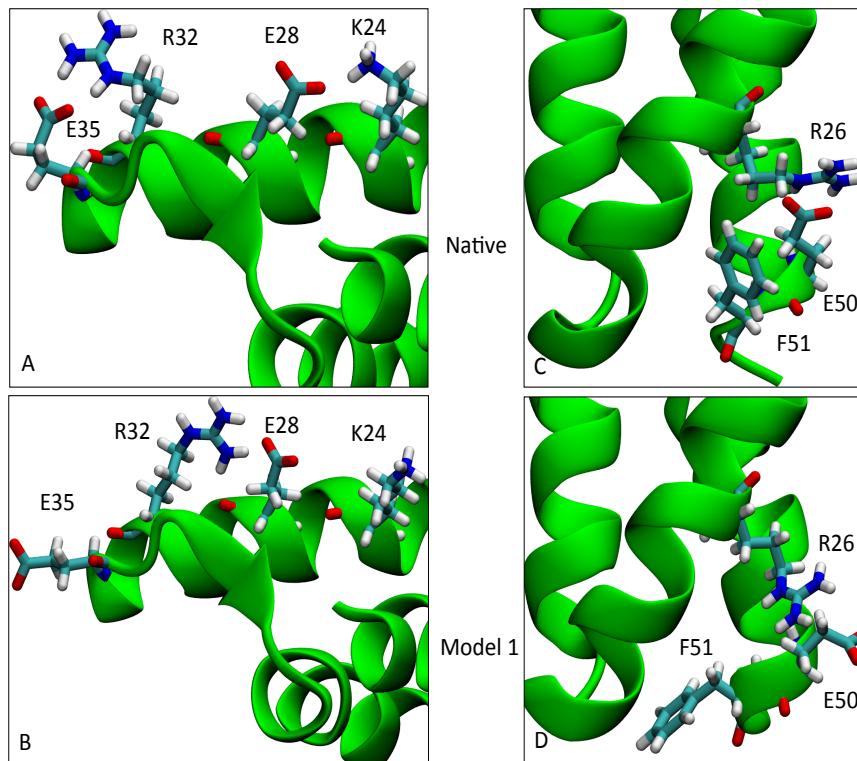


Figure 8: The per residue free energy calculation found some differences of the side chain orientation in computer generated model 1 and experimental structure of CASP target T0538. (A) Two salt bridges between residues E35-R32 and E28-K24 exist in the native structure. These are absent in the computer generated model structure. Instead (B) a new salt bridge is formed between residues R32-E28 in model. (C) Another salt bridge between residues R26-E50 in the experimental structure. (D) In comparison to the native structure, orientation of F51 differs in the model.

Table 1: The confinement method assigns a more favorable free energy to the experimentally determined structure than to computer-generated predictions. For each target, we examined as many as five predictions submitted by CASP participants. We report the free energy difference between the most favorable decoy and the experimentally determined structure. Positive $\Delta\Delta G (= G_{best\ decoy} - G_{native})$ values indicate that the experimental structure is predicted to be more favorable than any of the decoys.

CASP Target	PDB Identifier	$\Delta\Delta G$ (kcal/mol)	GDT-TS
T0531	2KJX	11.15 ± 0.70	44
T0538	2L09	-3.00 ± 0.47	96
T0540	3MX7	16.94 ± 0.49	70
T0559	2L01	2.10 ± 0.24	94
T0560	2L02	18.00 ± 0.49	94
T0569	2KYW	20.01 ± 0.69	78

Table 2: The six different CASP targets that are used in this study. The corresponding PDB Identifier and the organism type are also listed.

CASP Target	PDB Identifier	Organism type
T0531	2KJX	extracellular domain of the jumping translocation breakpoint protein
T0538	2L09	protein asr4154 from Nostoc sp. PCC7120
T0540	3MX7	human Fas apoptotic inhibitory molecule
T0559	2L01	protein BVU3908 from Bacteroides vulgatus
T0560	2L02	protein BT2368 from Bacteroides thetaiotaomicron
T0569	2KYW	domain of adhesion exoprotein from Pediococcus pentosaceus