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TASSER_low-zsc: An approach to improve structure prediction using low z-score ranked templates

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Abstract

In a variety of threading methods, often poorly ranked (low z-score) templates have good alignments. Here, a new method, *TASSER_low-zsc* that identifies these low z-score ranked templates to improve protein structure prediction accuracy is described. The approach consists of clustering of threading templates by affinity propagation on the basis of structural similarity (*thread_cluster*) followed by TASSER modeling, with final models selected using a TASSER_QA variant. To establish generality of the approach, templates provided by two threading methods, SP³ and SPARKS², are examined. The SP³ and SPARKS² benchmark datasets consist of 351 and 357 medium/hard proteins (those with moderate to poor quality templates and/or alignments) of length ≤ 250 residues respectively. For SP³ medium and hard targets, using *thread_cluster*, the TM-scores of the best template improve by ~4% and ~9% over the original set (without low z-score templates) respectively; after TASSER modeling/refinement and ranking, the best model improves by ~7% and ~9% over the best model generated with the original template set. Moreover, *TASSER_low-zsc* generates 22% (43%) more foldable medium (hard) targets. Similar improvements are observed with low ranked templates from SPARKS². The template clustering approach could be applied to other modeling methods that utilize multiple templates to improve structure prediction.

Keywords

Structure prediction; threading; TASSER; tertiary structure

Introduction

Despite significant progress, the prediction of protein structure remains an unsolved problem in computational structural biology^{1–3}. Historically, structure prediction methods have been divided into three general categories: comparative modeling (CM)^{1,4–6}, threading^{7–11} and free modeling^{12–15}. The basic objective of CM and threading approaches is to identify a set of structurally related template proteins (with known tertiary structure) to the target sequence^{5,9}. CM methods identify template proteins with a clear evolutionary relationship to the target using sequence-based methods^{5,16}, whereas threading, by including protein structural information, strives to identify template proteins having a similar fold as the target irrespective of their evolutionary relationship^{3,8,9}. Due to the convergence of threading and CM methods, both are referred to as Template-based modeling (TBM)¹⁷. In TBM, once the related template is identified, the target sequence is aligned to the template structure either indirectly by performing a sequence alignment and then transferring this alignment to the associated position in the structure or by directly incorporating structural information into

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the alignment procedure^{5,9–11}. A full-length model is then generated by building the chain in the unaligned regions of the template. This full-length structure is then refined, with the goal of improving model quality relative to the initial TBM provided alignment. In contrast, in template free modeling (FM), one does not use any global template structural information as input^{12,13}. Thus, the possibility of assembling a novel fold exists.

In recent years, TBM has emerged as the most robust approach to protein structure prediction^{3,17}. Advances in better template identification and improved alignment accuracy resulted from use of profile-profile alignments^{18–20}, inclusion of structural properties such as solvent accessibility²¹ and structural profiles^{8,10,11,22–24}, hidden Markov models^{25,26}, machine learning approaches^{27,28} and the employment of meta-servers^{29–31}. Model refinement can be achieved by using multiple templates to generate better alignments^{32,33}, iterative refinement^{34,35} as well as physics and evolution-based potentials^{12,35,36}. The ultimate success of TBM requires that similar structures to those adopted by the target be found in the Protein Data Bank (PDB)³⁷. Recent studies have demonstrated that the current PDB library is most likely complete in that it can provide templates for all compact, single domain proteins from which low-to-moderate resolution structures can be built^{38,39}. However, the key issue is to select such templates and to generate high quality alignments.

Among the more successful structure prediction approaches is TASSER (Threading/ASSEMBly/Refinement)³² and its variants chunk-TASSER and pro-sp3-TASSER⁴⁰, where in large scale benchmarking, it was shown that reasonable models could be built for both template-based and template-free modeling for weakly/non-homologous proteins^{32,40–43}. Recent improvements resulted from incorporation of improved contact predictions as in TASSER_2.0⁴⁴, iterative TASSER^{iter}³⁴, and the use of templates identified by multiple threading algorithms such as in METATASSER⁴³ or pro-sp3-TASSER⁴⁰.

In threading, usually some type of knowledge-based scoring function is used to rank the particular sequence-structure alignment^{8,9,45}. Furthermore, the score significance of a target aligned to a given template is evaluated in terms its z-score or through use of a neural network to rank the templates^{46,47}. In that regards, the z-score of the sequence mounted in a given structure is defined by

$$Z=(E - \langle E \rangle)/\sqrt{(\langle E^2 \rangle - \langle E \rangle^2)}$$

the quantity in $\langle \rangle$ denotes the average of the best alignment given by dynamic programming over the template library, and E is the score or energy⁹. Usually, z-score based ranking identifies best-fit templates for the target sequence^{8,45,48}. However, we have observed that templates with good alignments to the target native structure (TM-score ≥ 0.40)⁴⁹, sometimes have quite poor ranks based on their z-scores. Identification of such templates/alignments is important in those situations when the top ranked template/alignments are of poor quality.

In the present study, our goal is to develop a methodology that includes low ranked z-score but good quality templates to improve structure prediction using the TASSER methodology³². We can consider a large number of templates, having low z-score, as an input for TASSER. However, the main issue in considering more templates is that the consensus information from templates close to the native structure is usually diminished. To circumvent this issue, we have developed the ‘*thread_cluster*’ algorithm, which is used to filter and cluster structurally similar templates. Subsequently for each cluster, we perform TASSER simulations to generate an ensemble of models, which is ranked by our in-house

model ranking method described below. We refer to this protocol from template selection to model ranking as *TASSER_{low-zsc}*. We benchmarked this method on representative benchmark datasets using two threading methods, SP³¹⁰ and SPARKS²¹¹. In benchmarking, we show that this new approach of template clustering followed by TASSER modeling/refinement generates models which are better both in terms structure prediction accuracy and number of foldable targets compared to those generated using the original set of selected templates.

METHODS

We compiled a benchmark set of 691 proteins (whole chains or domains) of length ≤ 250 residues with pairwise sequence identity $\leq 40\%$ with the proteins in the PDB³⁷ template library. The template library consists of 17,888 proteins comprised of both whole chains and domains. Since for a multiple domain protein, both the whole chain and its domains are included, there is certain redundancy in the library. The benchmark proteins were released subsequent to the construction of template library. Thus, this benchmark is set up to mimic a Critical Assessment of Structure Prediction (CASP)-like⁵⁰ scenario. The list of proteins in the template library and benchmark dataset for SP³ and SPARKS² is available at: http://cssb.biology.gatech.edu/skolnick/files/TASSER_low-zsc. In addition to the above dataset, we have evaluated *TASSER_{low-zsc}* on 108 CASP8 targets⁵¹ of length ≤ 350 residues. Both SP³ and SPARKS² were used in the meta-threading procedure in CASP8⁵². Hence, we have used the threading output obtained during CASP8.

The protein structure prediction protocol described here consists of three main steps: a), filtering and clustering of templates via *thread_cluster* b) generation of an ensemble of models using TASSER and c) model ranking. *TASSER_{low-zsc}* approach is schematically described by the flowchart shown in Figure 1.

To evaluate the robustness of the *TASSER_{low-zsc}* methodology, we employ two threading programs, viz. SP³ and SPARKS². For each threading method, we classify targets into easy/medium/hard categories based on the z-score of the top template from the respective threading programs. In SP³, targets whose top template has a z-score ≥ 6.0 are classified as “easy”, those with a z-score ≤ 4.5 as “hard” and those having a $4.5 < \text{z-score} < 6.0$ as “medium”. In SPARKS², targets whose top template has a z-score ≥ 5.5 are classified as “easy”, those with z-score ≤ 4.0 as “hard”, and targets with $4.0 < \text{z-score} < 5.5$ as “medium”. In general, “easy” targets have templates with good threading alignments. “Medium” targets have good structure alignments to the native structure, but poor threading alignments, whereas, “Hard” targets generally have poor quality global structural alignments to the native state. Since for “easy” targets, usually the best template (that is closest to the native structure) is among the top ranked templates, here, we only consider medium/hard targets, as classified by their respective threading program, for the assessment of the method.

In the following, we discuss in detail each of the steps of *TASSER_{low-zsc}*:

1. Thread_cluster: filtering and clustering of templates

The top N templates as ranked by their z-score are clustered using the TM-score⁴⁹ structural similarity metric, which is calculated for the common aligned region between templates, where the length used for the TM-score calculation is the number of common aligned residues between two templates. For different threading methods, we obtain the maximum number of templates (N) to be used for clustering based on the enrichment factor of templates i.e. templates whose TM-score ≥ 0.40 to the native structure. For “medium” targets, from SP³ and SPARKS², we consider the top 60 and 75 z-score ranked templates

respectively, while for “hard” targets, we consider the top 70 and 80 z-score ranked templates, respectively. However, to reduce the number of templates considered, if there is significant structural similarity among the top 40 templates, (see below), we reduce N to 40.

We have used the Affinity Propagation (AP) method for clustering⁵³. Affinity propagation algorithm simultaneously considers all data points as potential “exemplars”. In the clustering step, real-valued messages are exchanged between data points until a good set of exemplars and the corresponding cluster emerges. There are two kinds of messages exchanged between data points “responsibility” and “availability”. In this, “responsibility” reflects the accumulated evidence of how well suited a data point is to act as an exemplar for another data point considering other possible exemplars, and “availability” is how appropriate it is for a data point to select a particular exemplar, taking into account the support from the other points for the exemplar. These real-values messages are updated during clustering. It has been shown that AP provides clusters with much lower error in comparison to other similar methods⁵³.

The AP program requires the similarity between the data points and a preference parameter (P_{pref}) of a point, which is the *a priori* suitability of a point to serve as exemplar. Using AP, one can obtain the desired number of clusters by changing the value of P_{pref} . A high value of P_{pref} will cause AP to find many clusters, while a low value of P_{pref} will lead to a small number of clusters.

In practice, the following algorithm is employed for filtering and clustering the templates:

1. For each of the top 40 templates, calculate the number of templates with pairwise TM-score ≥ 0.45 . If the median of this distribution is ≥ 10 , then set $N=40$; otherwise, use the top N templates as defined previously.
2. For each of the N templates, calculate the number of templates with a TM-score ≥ 0.45 . If the median of this distribution is ≥ 10 , set $P_{pref}=0.45$; otherwise, set $P_{pref}=0.35$. Next, remove any template whose best TM-score value to any other another template is less than P_{pref} . This removes templates with insignificant similarity within the list of N templates.
3. Calculate all-against-all TM-scores (for their common threading aligned region) for the filtered set of N templates. Next, perform AP clustering (with P_{pref} set in the previous step) on the basis of their TM-score similarity among templates. After clustering, all one member clusters are removed. In addition, from each cluster, remove those templates whose best structurally similar template, as given by TM-score, within the cluster has a TM-score $\leq P_{pref}$. Then, for each cluster, calculate the average of all-against-all TM-scores. For the next step, if the average TM-score is ≥ 0.65 , only one template with the highest z-score is considered; otherwise, all the members of the cluster are considered. This constitutes the filtered N_{mod} templates. Furthermore, add the top 10 z-score ranked templates to the N_{mod} templates and remove any redundant templates from the modified list of templates.
4. For the N_{mod} templates, calculate their all-against-all TM-scores. Perform AP clustering with P_{pref} set to the median obtained from the distribution of pairwise similarity values. In this step, the desired maximum number of clusters $N_{clus} \leq 5$. In practice, N_{clus} varies depending on the number of templates (N_{mod}). If $N_{mod} < 30$, then set $N_{clus} \leq 3$; otherwise, if $30 < N_{mod} < 40$, set $N_{clus} \leq 4$. The required number of clusters is achieved by iterating the AP clustering method with concomitant decrease in the value of P_{pref} . This step assigns every template in N_{mod} to one of the clusters. Usually, structurally similar templates are clustered together at the end of the AP clustering process.

2. TASSER simulations

As shown in Figure 1, we employed TASSER to generate the ensemble of models. Since TASSER has been extensively described in the literature^{32,41}, here we just present a brief overview of its essential components. The TASSER force-field consist of knowledge-based statistical potentials describing short-range backbone correlations, pairwise interactions, hydrogen-bonding, secondary structure propensities and a set of predicted side-chain contact and distance restraints derived from the initial structures. Generally, the structures that provide the restraints and the starting conformations are the same. The structures generated by TASSER are then clustered using SPICKER⁵⁴, which provides the list of models ranked by cluster density. Here, we limit the run time of TASSER to ~24 hrs to avoid long simulation times especially for longer proteins.

To generate a diverse set of models, the following sets of templates are considered:

1. For each of the N_{clus} clusters, templates within the cluster are used as an input for TASSER along with their consensus side-chain contacts (present in $\frac{3}{4}$ of the templates).
2. All the templates (N_{mod}) are used as input for TASSER.

The ensemble of models generated in the above procedure is ranked with the E_{frag} component of TASSER-QA⁵⁵. Next, for each of the top 15 ranked models, the three best threading templates (NR_{temp}) having TM-score ≥ 0.45 to the TASSER model are identified. If no templates could be identified with this TM-score cut-off, this value is reduced until there is at least a total of four templates identified by the top 15 ranked models. Usually for most targets, the TM-score cut-off value does not drop below 0.25. Using this set of models and templates, the following set of input structures are used in short TASSER simulations (~12 hrs)

1. Only the templates (NR_{temp}) are used as input structures for the simulation.
2. The templates (NR_{temp}) and those models (NR_{model}), which could identify templates with TM-score ≥ 0.45 (as described before), are used as input structures.

The main objective of performing this step of TASSER is to enrich the set of good (closer to native structures) models provided by the templates identified in the previous step. The final ensemble consists of models from the present and previous simulations.

3. Ranking of models

Model ranking is an important unsolved problem in the field of protein modeling⁵⁶. In our ranking procedure, we use a combination of model clustering and ranking with the E_{frag} component of TASSER-QA⁵⁵. The basic idea is to cluster structurally similar models and rank the representative model obtained from each cluster. In practice, our model ranking algorithm is as follows:

1. For all models, first calculate the all-against-all similarity score as measured by the TM-score. Next, cluster the models by AP method with the preference parameter set as the median of the similarity score distribution. Generate the modified list of models by only considering models from clusters containing at least three members. Next, for the modified list of models, iterate AP clustering by tuning the P_{pref} parameter to result in a maximum of five clusters.
2. For each cluster, rank the models within the cluster by the average highest TM-score to all other models in the cluster. Next, superimpose all the models to the top ranked model and generate the centroid model by averaging the coordinate positions within a root mean-square deviation, RMSD, of less than 6.5 Å after

superimposition. This centroid model is considered as the final model. PULCHRA⁵⁷ is then used to fix the artifacts in the centroid model due to averaging. If PULCHRA fails, the model closest to the average structure is considered as the final model.

3. Rank the final set of models using the E_{frag} component of TASSER-QA.

TASSER with original set of templates (TASSER_org)—In order to evaluate the improvement in structure prediction with *TASSER_low-zsc*, we generated models with TASSER using the numbers of templates benchmarked in previous studies⁴³. For both SP³ and SPARKS², we used the top 10 z-score ranked templates as an input for TASSER. These are the number of templates that would be used in a standard TASSER prediction scenario.

Results and discussion

TASSER_low-zsc is evaluated on a representative benchmark dataset and on CASP8 targets using two different threading methods, SP³ and SPARKS². In the present work, we have used the TM-score⁴⁹ to assess the quality of the structure template and the predicted full-length model. The benchmarking results of the method are followed with evaluation on CASP8 targets.

Targets are classified into easy/medium/hard sets based on the top template z-score (see Methods). For SP³, the number of targets classified into the easy, medium and hard set is 340, 148 and 203, respectively. For SPARKS², the number of targets in the easy, medium and hard set is 334, 155 and 202, respectively. As mentioned before, we consider only medium/hard targets for assessment. Thus, SP³ and SPARKS² are evaluated on 351 and 357 targets respectively. In what follows, we first assess the performance of the *thread_cluster* algorithm and then assess the accuracy of the structure prediction protocol, *TASSER_low-zsc*.

The main objective of the *thread_cluster* algorithm is to filter low z-score ranked templates to identify a structurally consistent set. When we consider low z-score ranked templates, the number of templates increases and the fraction of “good” templates (TM-score to native ≥ 0.40) could either be enriched or depleted. Hence, clustering structurally similar templates, as measured by their TM-score, attempts to retain these “good” templates. For “good” templates, the assumption here is that more than one such template is identified by the given threading procedure. First, we assess ability of *thread_cluster* to retain the best available template among all top N templates after clustering. For both SP³ and SPARKS², Table I summarizes the comparison among the best available templates from the original template set, the top N set, and templates after filtering and clustering. On average, the TM-score to the native of the best template among the top N templates for SP³ medium and hard targets is better than the original template set by ~8% and ~11% respectively. After applying *thread_cluster*, on average the TM-score to native structure of the best template among available templates is better by ~4% and ~9% for medium and hard targets respectively. Similarly, for SPARKS², after using *thread_cluster*, on average the TM-score to the native structure of the best template among available templates is better by ~8% and ~10% for medium and hard targets respectively. This suggests that for most targets *thread_cluster* can retain the best available template among the top N templates.

Next, we compare the performance of *thread_cluster* in terms of the number of targets with TM-score ≥ 0.4 (indicative of significant structural similarity to the native structure). As is evident from Table I, using SP³ (SPARKS²) for medium targets, the number of foldable targets in the top N template set increases by 16% (30%) in comparison to the original template set. For hard targets, the relative number of foldable proteins increases by 55%

(96%) respectively. Subsequent to the *thread_cluster* procedure, for medium targets, using SP³ (SPARKS²) the improvement in terms of number of foldable targets is still 13% (20%) in comparison to the original template set, while for hard targets the number increases by 49% (65%). The diminution relative to the best possible performance reflects the fact that for some targets, *thread_cluster* could not recover the best template. For most of these targets, there are a few “good” templates in the list of N templates used for clustering. Moreover, for some targets, the “good” template does not have significant structural similarity to any other template. Hence, these templates are filtered out in the clustering process of *thread_cluster*.

Subsequent to *thread_cluster*, the ensemble of full length models is generated and ranked as described in Methods. We present the benchmarking result of structure prediction using templates from SP³ and SPARKS² threading.

Benchmarking results using templates from SP³

The main objective of including low z-score ranking templates in TASSER modeling is to increase the accuracy of protein structure prediction. For assessment, we compared the models generated from two procedures, *i.e.* *TASSER_org*, which has models generated with TASSER using the original template set (see Methods) and *TASSER_low-zsc*, which has the models generated with TASSER using the input templates obtained after *thread_cluster* followed by ranking of the model ensemble (see Methods). Moreover, *TASSER_org* uses the standard TASSER structure prediction scenario^{32,43}. The comparison of performance between *TASSER_org* and *TASSER_low-zsc* methods is summarized in Table IIA. For the first model, *TASSER_low-zsc* shows on average a 5% and 7% TM-score improvement over *TASSER_org* for medium and hard targets respectively. Similarly, on average the best (among the top five) model TM-score improvement from *TASSER_low-zsc* over *TASSER_org* is ~7% and ~9% for medium and hard targets respectively. Moreover, for the medium (hard) set, *TASSER_low-zsc* provides 22% (43%) more foldable targets than *TASSER_org*, when the best of five models are considered.

For the detailed analysis of the improved performance of *TASSER_low-zsc*, we used the best (of top five) model (from both procedures), TM-score to the native structure, to classify the medium and hard targets into following four sets: a), targets having best model TM-score < 0.40 from both procedures; b), targets with a best model TM-score < 0.40 from *TASSER_org*, and TM-score ≥ 0.40 from *TASSER_low-zsc* method; c), proteins with best model TM-score ≥ 0.40 from *TASSER_org* and TM-score < 0.40 from *TASSER_low-zsc* method and d) proteins with best model TM-score ≥ 0.40 from both procedures. The average TM-score of best models from both methods for the targets classified into these four sets is summarized in Table IIB. The targets with a best model TM-score ≥ 0.40 to the native structure only from *TASSER_low-zsc* method (set b in the above classification) show an average TM-score improvement of 37% (40%) for medium (hard) targets over models from *TASSER_org*. Interestingly, even for targets whose best model TM-score to the native structure from both methods ≥ 0.40 (set d in above classification), *TASSER_low-zsc* on average shows an improvement of 4% in TM-score for both medium and hard targets (Table IIB). For very few targets, *TASSER_low-zsc* results in a model with lower accuracy in terms of its TM-score (Table IIB) to the native structure.

In Figures 2A and 2B, for the medium and hard targets respectively, we show the comparison of the best model from *TASSER_org* and *TASSER_low-zsc*. Overall, the best model from *TASSER_low-zsc* has a higher TM-score than *TASSER_org*. However for certain targets, the performance of *TASSER_low-zsc* is worse. These cases are partly due to the failure of the method to select and rank models. For example, target 2zb5_A_d1 has a *TASSER_org* generated best model with a TM-score of 0.457. In contrast, *TASSER_low-zsc*

results in a best model with a TM-score of 0.343. However, the best possible model among the ensemble of models from *TASSER_low-zsc* has a TM-score of 0.454. Next, we compare the distribution of the number of targets having models greater than or equal to a given TM-score threshold value. This comparison is shown in Figure 3 where the number of targets is plotted as a function of TM-score and demonstrates that *TASSER_low-zsc* is better for almost all TM-score threshold cut-off values.

In the above sections, we analyzed and discussed the results of *TASSER_low-zsc* models after selection and ranking from an ensemble of models generated after various TASSER simulations. In addition, from this ensemble, we can find the best model and determine the maximum possible improvement for *TASSER_low-zsc* over *TASSER_org*. For medium and hard targets, the best possible model from *TASSER_low-zsc* has an average TM-score to native of 0.442 and 0.425, respectively. The number of foldable proteins is 81 (100) for medium (hard) targets. Hence, given a perfect algorithm to select and rank models, we could achieve a TM-score improvement of ~16% and ~20% for medium and hard targets respectively over models from *TASSER_org* method. However, in *TASSER_low-zsc*, we are able to achieve a ~7–9% TM-score improvement on average with respect to *TASSER_org*. This demands that we further improve the model selection and ranking.

The improvement in structure prediction accuracy from *TASSER_low-zsc* with respect to *TASSER_org* is due to a combination of different factors: the identification of more “good” templates; the fact that structurally similar templates are clustered together enriches the consensus information with concomitant improvement in contact prediction accuracy, and the ability of TASSER to improve over the initial template structure. Finally, improved model selection and ranking also contributes to the success of the method, which requires having a set of similar models close to the native structure. Due to the convoluted effects of these factors, it is difficult to ascertain their relative contributions.

Below, we discuss examples where a combination of these factors contributes to the improvement in the model from *TASSER_low-zsc* as compared to *TASSER_org*. As shown in Figure 4, target 2qvv_A has a *TASSER_org* best model with a TM-score of 0.383. In the top 10 templates (the set used in *TASSER_org*), there is only one good template which has a TM-score of 0.572 to native structure, which is ranked 7th. There is another template with a TM-score of 0.434 to native structure; however, it is 41st ranked in z-score based ranking. Interestingly, *thread_cluster* could cluster these two good templates into a single cluster along with other templates having a TM-score ≥ 0.35 to the native structure. This probably contributed to the successful prediction from *TASSER_low-zsc*, which has the best of top five models with a TM-score of 0.647 to the native structure.

Another interesting example is target 2ras_A_d2 shown in Figure 4. For this protein, the best template has a TM-score of 0.539 (ranked 10th) among the top 10 templates. However, the remaining 9 templates have a TM-score to native < 0.20 . The best model from *TASSER_org* has a TM-score of 0.228, even though the best template is included in *TASSER_org* modeling. In the low z-score ranked templates, there are several templates with TM-score > 0.40 ; these are ranked at positions > 12 . The best of these has a TM-score to native of 0.605 and is ranked 27th. Interestingly, *thread_cluster* clusters all these templates having TM-score > 0.40 to native into one cluster. Furthermore, the selection and ranking procedure resulted in the best model (among top five models) having a TM-score of 0.596 (Figure 4).

This suggests that *TASSER_low-zsc* can improve the structure prediction accuracy over the standard *TASSER_org* models. Apart from providing better models for targets, which have

templates with TM-score < 0.40 in the top 10 templates, *TASSER_low-zsc* also provides better models for other targets whose TM-score > 0.4 .

Benchmarking results using templates from SPARKS²

In the previous section, we have demonstrated that using low z-score templates from SP³, *TASSER_low-zsc* improves structure prediction, as assessed by TM-score, with respect to the model generated with the original template set (*TASSER_org*). We next assess its applicability to templates generated from a different threading program, SPARKS². In Table I, we have already shown that for SPARKS² *thread_cluster* could recover the best template for most medium/hard targets. Here, we compare the prediction success of *TASSER_low-zsc* and *TASSER_org*. The structure prediction procedure is the same as that used for SP³ (see Methods). The performance comparison between *TASSER_low-zsc* and *TASSER_org* is presented in Table III. We note that the average TM-score to native of the best possible model among the ensemble of models is 0.428 and 0.365 for medium and hard targets respectively. Hence, given the ability to choose the best model from the ensemble, we could achieve an improvement of ~16% (~21%) for medium (hard) targets. In practice, for the first model, *TASSER_low-zsc* shows an average TM-score improvement of ~5% (~7%) over *TASSER_org* for medium (hard) targets. For the best of five models, a similar TM-score improvement of ~6% and ~7% is observed for medium and hard targets, respectively. Furthermore, the number of foldable targets using *TASSER_low-zsc* increases by 11% for medium proteins. For hard proteins, *TASSER_low-zsc* has almost twice the number of foldable targets as *TASSER_org*.

Next, for a detailed analysis of the improvement observed with *TASSER_low-zsc*, we classified the medium/hard targets into four sets as defined above for the evaluation of SP³ (see “Benchmarking results using templates from SP³”). The results are presented in Table S1 (see in Supporting Information). The improvement in targets with best (of top five) model TM-score to native structure ≥ 0.40 only from *TASSER_low-zsc* is ~42% (~43%) for medium (hard) proteins over *TASSER_org*. Similarly, targets with best model TM-score ≥ 0.40 to the native structure from both methods shows on average a TM-score improvement of ~7% (~2%). However, there are a few targets having a TM-score ≥ 0.40 from *TASSER_org*, for which *TASSER_low-zsc* gives poor quality models (TM-score < 0.40). This is partly due to issues in model ranking. Thus, the behavior of *TASSER_low-zsc* when SPARKS² provides the set of threaded templates is very similar to that of SP³.

In Figures 5A and 5B, we show the comparison of the best model from *TASSER_org* and *TASSER_low-zsc* for medium and hard targets respectively. On average, *TASSER_low-zsc* performs better than *TASSER_org* as was the case when templates from SP³ are used. Similarly as shown in Figure S1 (see in Supporting Information), for TM-scores ≥ 0.3 , the improvement of *TASSER_low-zsc* is seen over all values of the TM score. For some targets, the predicted model from *TASSER_low-zsc* has a lower TM-score in comparison to the *TASSER_org* model. As discussed before, this is partly due to issues in model ranking.

Prediction results for CASP 8 targets

We have used *TASSER_low-zsc* with the templates derived from SP³ and SPARKS² for the assessment of CASP8 targets.

The targets are classified into easy/medium/hard categories based on z-score of the first template. For SP³ the number of targets classified into the easy, medium and hard set is 87, 13 and 8, respectively. Similarly, in the case of SPARKS², the number of easy medium and hard targets is 91, 3 and 14 respectively. Since little if any improvement from *TASSER_low-zsc* is expected for easy targets, we are limited to a very small subset of CASP8 proteins on

which to perform the analysis, and the results are not likely to be statistically significant. Nevertheless, it is of interest to see how *TASSER_low-zsc* would have performed in CASP8. In the analysis, we have used 21 and 17 medium/hard targets from SP³ and SPARKS² respectively (see Table S2 in Supporting Information). In the following section, we present the assessment of structure prediction employing templates selected by SP³ and then from SPARKS².

The *thread_cluster* procedure on templates from SP³ shows an average improvement of ~10% and ~5% for medium and hard targets respectively (Table S3 in Supporting Information). For both medium and hard categories, only one additional protein has a TM-score ≥ 0.40 after *thread_cluster*. We used the modeling procedure as described in the Methods section. In figure 6, for medium/hard targets, we show the comparison of the best model from *TASSER_org* and *TASSER_low-zsc*. For medium targets, the first (best) model using *TASSER_low-zsc* shows an average TM-score improvement of ~4% (~6%) over *TASSER_org* (Table S4 in Supporting Information). However, for hard targets, the average TM-score of the best model from *TASSER_low-zsc* becomes slightly worst in comparison to *TASSER_org* (Table S4 in Supporting Information). Further, detailed analysis showed that for most of hard targets, good templates (TM-score ≥ 0.40) are not in top N set of templates. Thus, their best model has a TM-score < 0.40 for both modeling procedures (Table S5A in Supporting Information). An interesting example is target T0480, classified in the medium category. The best model from *TASSER_org* has a TM-score to native of 0.390. The clustering method (*thread_cluster*) could cluster low-rank templates (at positions 15, 28, 31, 33 and 40) having TM-scores > 0.40 into one cluster. This resulted in a best model from *TASSER_low-zsc* with a TM-score to native of 0.456.

Using templates from SPARKS², *thread_cluster* is not able to recover the best possible template after clustering for one of the medium targets, T0514. For this target, there is only one template with TM-score > 0.40 and this could not cluster with any other templates. A similar issue exists for hard target T0399. Subsequent to modeling, the best models from *TASSER_low-zsc* on average do not show significant improvement with respect to *TASSER_org* (Table S4 in Supporting Information). We then classified the various medium/hard targets as to whether their TM-scores are above or below 0.4 for the *TASSER_org* and *TASSER_low-zsc* in Table S5B (see in Supporting Information). In case of medium targets, because of T0514, *TASSER_org* method uses a template with TM-score > 0.4 but *TASSER_low-zsc* does not identify this template; this reduced the TM-score from 0.434 to 0.282, but these results are anecdotal, as just one target is considered.

Figure 6 shows the comparison of the best model from both modeling procedures for medium/hard targets. For most hard targets, good templates (TM-score ≥ 0.40) were not present in top N set of templates. Hence, an improvement in structure prediction accuracy using *TASSER_low-zsc* similar to the benchmark studies could not be observed with CASP8 targets. An interesting case is T0482, which is classified as a hard target by SPARKS². The best model from *TASSER_org* has a TM-score to native of 0.383, after *TASSER_low-zsc* two templates at position 7 and 8 with TM-score > 0.38 clustered together with other similar templates. This then provides the best model from *TASSER_low-zsc* with a TM-score to native of 0.441. But, again, caution should be taken in interpreting the generality of these results as there are very few targets in the medium/hard regime.

These results suggest that *TASSER_low-zsc* could be extended to other threading procedures as well. In addition, *thread_cluster* could be combined with other modeling procedures, which use information from multiple templates as an input to improve structure prediction.

Conclusions

We have developed the *TASSER_{low-zsc}* approach to improve protein structure prediction using low z-score ranked, but good quality templates. Template clustering is performed using the *thread_cluster* algorithm, which attempts to retain only structurally similar templates as measured by their TM-score. *TASSER_{low-zsc}* was benchmarked for medium/hard targets from the SP³ and SPARKS² threading methods. The best model from *TASSER_{low-zsc}* shows an average TM-score improvement of ~6–9% with respect to the best model generated from *TASSER_{org}*. Furthermore, the number of foldable targets is significantly improved in *TASSER_{low-zsc}*.

A key unresolved issue is why are there good quality alignments in poorly ranked templates? Some unpublished work examining the set of structural alignments of the top ranked templates to native indicates that the majority have a good structural alignment to the native fold over at least a portion of their structure. If one requires that a set of three templates generate mutually consistent structural alignments, then ~14% of the residues have >90% probability of being part of the best structural alignment to native. In other words, there is a residual core of aligned residues that are commonly identified by threading (with the implication that the set of proteins are evolutionary related) and that this is the signal that is detected in the poorly ranked templates. In future work, we plan on exploring this issue in much greater detail.

At this juncture, what is encouraging is that *TASSER_{low-zsc}* significantly increases the fraction of medium/hard targets that are foldable. For example, if the best of top five models are considered, for medium/hard targets from SP³, the percentage of foldable proteins increases from 39%/26% using *TASSER_{org}* to 48%/37%; this is precisely the regime of target difficulty where progress has been quite slow. Despite this success, additional extensions are required. *Thread_cluster* will be used with other protein structure prediction methods which exploit information from multiple templates as well as meta-approaches. Work along this direction is in progress.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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Figure 1.
Flowchart of the *TASSER_{low-zsc}* approach.

Figure 2. Using SP³ threading, A) For medium targets, scatter plot of the best model from *TASSER_org* versus the best model from *TASSER_low-zsc*. B) Same in A, but for hard targets.

Figure 3. Histogram comparison between *TASSER_org* and *TASSER_low-zsc* on the 351 target SP³ benchmark dataset. TM-scores are from the best of top five models.

Figure 4.

Examples showing an improved predicted model from *TASSER_low-zsc* in comparison to *TASSER_org* for two targets from the SP³ benchmark dataset. For each target, the superposition of the native structure (thin backbone) and best model (thick backbone) from *TASSER_org* (on the left) and from *TASSER_low-zsc* (on the right) are shown. Blue to red goes from the N- to the C-terminus.

Figure 5. Using SPARKS² threading, A) For medium targets, scatter plot of the best model from *TASSER_org* approach versus the best model from *TASSER_low-zsc*. B) Same as in A, but for hard targets.

Figure 6.

For CASP8 targets, scatter plot of the best model from *TASSER_org* versus the best model from *TASSER_low-zsc* using templates from SP³ or SPARKS².

Table I

Comparison of best template before and after *thread_cluster*

Threading	Target type	No. of targets	<TM _{org} >	<TM _{topn} >	<TM _{clus} >
SP ³	Medium	148	0.373 (61)	0.402 (71)	0.389 (69)
	Hard	203	0.349 (51)	0.388 (79)	0.380 (76)
SPARKS ²	Medium	155	0.358 (50)	0.404 (65)	0.385 (60)
	Hard	202	0.287 (26)	0.338 (51)	0.317 (43)

TM_{org}, TM_{topn} and TM_{clus} refer to the TM-score of the best template to the native structure among original template set (top 10 templates), among the top N templates and among templates identified after the filtering/clustering step in *thread_cluster* respectively. The number in the parenthesis is the number of proteins with TM-score ≥ 0.40 .

Table IIA

Summary of structure prediction results for SP³ medium/hard targets

Target type	No. of targets	First model		Best of top five models	
		<TMM _{org} >	<TMM _{clus} >	<TMM _{org} >	<TMM _{clus} >
Medium	148	0.345 (48)	0.363 (54)	0.380 (58)	0.407 (71)
Hard	203	0.316 (37)	0.340 (43)	0.354 (53)	0.385 (76)

TMM_{Org} and TMM_{clus} refer to TM-score of the model from TASSER_{org} and TASSER_{low-zsc}, respectively. The number in the parenthesis is the number of proteins with TM-score ≥ 0.40.

Summary of structure prediction results for SP³ targets classified into foldable/non-foldable categories either from one or both methods

Table IIB

Target classification	Target type	<TMB _{org} >	<TMB _{zsc} >	Target type	<TMB _{org} >	<TMB _{zsc} >
TMB _{org} < 0.40 and TMB _{zsc} < 0.40	Medium (75)	0.261	0.273	Hard (125)	0.284	0.297
TMB _{org} < 0.40 and TMB _{zsc} ≥ 0.40	Medium (15)	0.340	0.467	Hard (25)	0.345	0.484
TMB _{org} ≥ 0.40 and TMB _{zsc} < 0.40	Medium (2)	0.468	0.365	Hard (2)	0.419	0.387
TMB _{org} ≥ 0.40 and TMB _{zsc} ≥ 0.40	Medium (56)	0.547	0.570	Hard (51)	0.529	0.550

TMB_{org} and TMB_{zsc} refer to TM-score of the best model (of the top 5) to native structure from *TASSER_org* and *TASSER_low-zsc*, respectively. The number in parenthesis is the number of proteins classified into that particular category.

Table III

Summary of structure prediction results for SPARKS² medium/hard targets

Target type	No. of targets	First model		Best of top five models	
		<TMM _{org} >	<TMM _{clus} >	<TMM _{org} >	<TMM _{clus} >
Medium	155	0.335 (43)	0.351 (44)	0.369 (57)	0.395 (63)
Hard	202	0.267 (16)	0.283 (22)	0.302 (20)	0.324 (38)

TMM_{Org} and TMM_{clus} refer to the TM-score of the model from the *TASSER_low-zsc* and *TASSER_org*, respectively. The number in the parenthesis is the number of proteins with TM-score ≥ 0.40 .