Sequence analysis

Using de novo protein structure predictions to measure the quality of very large multiple sequence alignments

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Abstract

Motivation: Multiple sequence alignments (MSAs) with large numbers of sequences are now commonplace. However, current multiple alignment benchmarks are ill-suited for testing these types of alignments, as test cases either contain a very small number of sequences or are based purely on simulation rather than empirical data.

Results: We take advantage of recent developments in protein structure prediction methods to create a benchmark (ContTest) for protein MSAs containing many thousands of sequences in each test case and which is based on empirical biological data. We rank popular MSA methods using this benchmark and verify a recent result showing that chained guide trees increase the accuracy of progressive alignment packages on datasets with thousands of proteins.

Availability and implementation: Benchmark data and scripts are available for download at http://www.bioinf.ucd.ie/download/ContTest.tar.gz.

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Supplementary information: Supplementary data are available at Bioinformatics online.

1 Introduction

Making a multiple sequence alignment (MSA) of nucleotide or amino acid sequences is a crucial step needed in a wide variety of bioinformatics studies. To overcome the extreme computational demands of exact algorithms, MSA software uses heuristics to find near-optimal alignments in a reasonable time. Different software packages, however, use different heuristics and objective functions and explicitly focus on different aspects of the MSA process such as accurate placement of gaps for phylogenetic reconstruction of small numbers of sequences (Lo¨ytynoja and Goldman, 2005) or sheer speed for making very large alignments e.g. MAFFT ParTTree (Katoh and Toh, 2007). Accurate benchmarks are therefore important to allow users of MSA software to choose the most appropriate tools and to guide developers in improving algorithms and heuristics (Iantorno et al., 2014).

As computational resources and sequence databases grow, and methods for creating MSAs are improved, the practical upper limit for the size of MSAs increases. Currently, alignments of tens of thousands of sequences can be made routinely on a desktop computer. It is therefore important that benchmarks can reflect alignment problems on this scale. Current protein MSA benchmarks can be divided into structure-, phylogeny-, simulation- and consistency-based benchmarks (Iantorno et al., 2014). Structure- and phylogeny-based benchmarks, in which scores are based on structural superpositions and accurate inference of phylogenetic trees, respectively, are strongly grounded in empirical biological data, but they focus on alignments of small numbers of sequences and are difficult to scale to larger datasets. Meanwhile, simulation- and consistency-based benchmarks are based on simulations of protein evolution and simple agreement between different MSA methods, respectively, and can involve alignments of arbitrary size. It is unclear, however, how well simulated sequences model actual biological sequences, while consistency measures only how similar the results of one heuristic method are to the results of other heuristic methods.

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MSAs containing large numbers of sequences are generally created by software using the progressive alignment method. Progressive alignment approximates the optimal alignment of multiple sequences by calculating a series of pairwise alignments according to a bifurcating tree known as a guide tree (Higgins et al., 1992). Recently, completely imbalanced—or ‘chained’—guide trees have been shown to increase the accuracy of some progressive MSA methods (Boyce et al., 2014) when aligning large numbers of sequences. This result is controversial (Boyce et al., 2015; Tan et al., 2015) and a benchmark is needed to accurately test the effect of chained guide trees on alignment accuracy.

Here, we describe a benchmark, ContTest, that is able to realistically test the accuracy of MSAs of very large numbers of amino acid sequences, making use of recent developments in the prediction of protein structures from large MSAs (Marks et al., 2011). We take the MSA that is to be tested and use it to predict a contact map for a protein in the alignment that also has a known three-dimensional (3D) structure. This predicted contact map is then compared with the known contact map for the same protein, and the alignment is scored based on their agreement (Fig. 1). The benchmark makes use of all the sequences in the alignment and does not reward over-alignment. It also avoids superposition of predicted and known structures as this step itself requires the use of heuristic algorithms. This process gives a very robust scoring system based on the assumption that, from a protein structure perspective, better alignment methods will result in more accurate contact maps.

We used the ContTest benchmark to compare the accuracy of some widely used MSA packages. We also used it to compare some different ways of using these packages, e.g. by varying the numbers of iterations and in particular using chained guide trees with the progressive alignment methods.

1.1 De novo contact prediction

Recently, several computationally efficient methods have been developed to predict protein structural information solely from an alignment of the sequence of interest with a large number of homologous sequences (Jones et al., 2012; Marks et al., 2011, 2012; Taylor et al., 2013). Residues that are in contact in the folded protein can exhibit a pattern of coevolution, where substitutions at one position may be compensated for by complementary mutations at nearby positions. EVfold-mfDCA (Marks et al., 2011) and PSICOV (Jones et al., 2012) detect correlations between the patterns of substitutions in pairs of columns of an MSA and output a list of pairs of residues predicted to be in contact.

We investigated if we could use the accuracy of contact predictions to infer the accuracy of the input MSAs. For a target protein with a resolved 3D structure in the Protein Data Bank (PDB) (Berman et al., 2000) and a large number of homologous sequences in the Pfam database (Finn et al., 2014), we compare contact predictions made using PSICOV or EVfold-mfDCA from an alignment of the sequences to a list of known contacts derived from the protein’s 3D coordinates. Accuracy is evaluated as the precision of the top 1/5 long range contacts predicted, where L is the length of the target protein. Long range is defined to mean any contact between residues separated by at least 23 other residues.

2 Materials and methods

2.1 Software versions and parameters
We used the following software to make MSAs:

- Clustal Omega 1.2.0 (Sievers et al., 2011). For one, two or three iterations we use the parameters:
  clustalo -i ... --guidetree-out=... -o ...

- ‘Chained’ guide trees are generated internally using the --pileup option:
  clustalo --pileup -i ... -o ...

- Clustal W2.1 (Larkin et al., 2007). We reuse the guide trees created by Clustal Omega:
  clustalw2 -quiet -infile=... -outfile=... -usetree=... -outorder=input

- HMMER 1.8.5 (hmmt) (Eddy, 1998)
  hmmt -o ... hmm_out in

- Kalign 2.04 (Lassmann et al., 2009)
  kalign -q -i ... -o ...

- Kalign 1.04 (Lassmann and Sonnhammer, 2005)
  kalign -f -q -i ... -o ...

- MAFFT v7.029b (Katoh and Standley, 2013) Default (FFT-NS-2):
  mafft --anysymbol in > out

  To make alignments using ‘chained’ guide trees, we create an external guide tree file in MAFFT tree format, which is read using the --treein option:
  mafft --anysymbol --treein ... in > out

- NW-NS-PartTree-1:
  mafft --retree 1 --maxiterate 0 --nofft --parttree --anysymbol in > out

- PSICOV version 2.1 (Jones et al., 2012) and FreeContact version 1.0.21 (Kajan et al., 2014) to predict residue-residue contacts from MSAs. We used PSICOV with default parameters and FreeContact with the --parprof evfold option.
We used CMView 1.1.1 (Vehlow et al., 2011) to extract contact maps from PDB files.

2.2 Test case selection and preparation

We based our test cases on the 150 proteins used to test contact prediction software by Jones et al. (2012). Each test case consists of the full set of sequences from a single Pfam family and a list of experimentally determined residue–residue contacts for a protein in that family. We retrieved 3D co-ordinates for each target protein from the PDB and extracted the list of long range residue–residue contacts using CMView (Vehlow et al., 2011), using an 8 Ångström Cα–Cα distance threshold (Cα for glycine) and a minimum sequence separation of 24 positions. We identified Pfam families for each target protein and downloaded sequences from Pfam in FASTA format, with all gaps represented by ‘-’ characters and all characters uppercase. Sequence orders were randomly shuffled before making alignments. Where the sequence of a protein differs between the PDB and Pfam, we perform a pairwise alignment between the PDB sequence and the Pfam sequence to correctly map columns of the alignment to beta carbons in the known structure.

Test cases were excluded where PSICOV was unable to make predictions from any of the alignments tested due to the alignments containing too few sequences. Test cases were also excluded where there were too many sequences in the Pfam family to be aligned by one or more of the alignment packages tested. For instance, Clustal W2 is unable to align more than 40,000 sequences, and Kalign 2 is unable to align more than 60,000 sequences.) One test case had too few true positive contacts to be useful for the benchmark. Fourteen test cases were excluded using these criteria, leaving a total of 136 test cases for the benchmark. Details of the test cases are given in Supplementary Table S1.

2.3 Scoring predicted contacts

The PSICOV and Evfold-mDCA algorithms both assign scores to each predicted contact. We assess the quality of long range contact predictions; for amino acid residues at positions i and j in the protein chain, a contact is defined as long range if |j − i| > 23. For a test case with a target protein of length L, we calculate the precision of the top scoring L/5 long range contacts output by PSICOV and EVfold-mDCA, which we refer to as the PSICOV precision and the EVfold-mDCA precision for that test case, respectively.

Because of the dependence of the output MSA on input sequence order for most methods tested, where possible we made three alignments for each test case with each alignment method, randomizing the order of the input sequences before each alignment. We calculated the average PSICOV precision and average EVfold-mDCA precision over the three replicates. Then, we calculated the geometric mean of the average PSICOV and average EVfold-mDCA precisions to arrive at a single number representing the precision of contact predictions for that test case. We call this the ContTest score for that test case. We rank alignment methods by the mean ContTest score over all test cases. Pfam stores only a single alignment for each family, while Clustal Omega runs extremely slowly when using chained guide trees. Therefore, results for Pfam and Clustal Omega (chained guide tree) are based on a single replicate only.

While this scoring method involves only a subset of columns in the alignment, we found that using receiver-operating characteristic curves to score predicted contacts produced almost the same rankings of alignments (Supplementary Text and Tables S2 and S3).

2.4 Statistical significance of benchmark scores

We calculate the statistical significance of the difference in two benchmark scores by using the Wilcoxon signed rank test where the ContTest scores for each of the 136 test cases are paired between the two conditions.

2.5 Making ‘bad’ alignments

To validate our benchmark, we use two methods to artificially create alignments which we expect to be worse than some reference alignment. First, given a reasonable starting alignment of a set of sequences, we expect that shifting a random subset of the sequences out of alignment with the others will result in an alignment that is poorer than the original alignment. We shift sequences out of alignment by adding a gap character to the start of the aligned sequence and truncating the last character. For each test case, we take the Pfam alignment and shift 0.5%, 1%, 2%, 3%, 4% or 5% of the sequences out of alignment, creating a series of alignments which we expect to be of decreasing quality and which should therefore result in decreasing benchmark scores.

Second, we perform a similar analysis by mutating random residues, which we expect should decrease contact prediction accuracy. Starting with the Pfam alignments, we mutate between 0 and 20% of residues in each alignment to a different residue. We expect that increasing numbers of mutations should again result in a decreasing ContTest score.

2.6 Adding HOMSTRAD reference sequences to ContTest alignments

We found mappings from 80 of the ContTest test cases to HOMSTRAD (Mizuguchi et al., 1998) alignments using the PFAM.db package for R (Carlson et al., n.d.). For each of these 80 test cases, we added the dealigned HOMSTRAD sequences to the unaligned Pfam sequences and randomly shuffled the sequence order before making alignments and proceeding with the benchmark process as usual. We then calculated both the average ContTest score and the average sum-of-pairs score (SPS) for the HOMSTRAD sequences, thus arriving at two measures of quality for the same alignments.

2.7 Measures of guide tree imbalance

The Sackin score of a rooted, binary tree is defined as the sum of the depths of its leaves (Sackin, 1972). The Sackin score for a tree with N leaves, that is as balanced as possible is N × log2(N); the Sackin score for a perfectly imbalanced (chained) tree with N leaves is (N + 2) × (N − 1)/2. Concrete values for trees of up to eight leaves can be found in Sievers et al. (2014).

The expected value of the Sackin score depends on the underlying evolutionary model. In the Equal Rates Markov or Yule (Yule et al., 1925) model, trees are built up, beginning with a node with just two leaves, by repeatedly selecting a leaf at random and replacing it with a node with two leaves until the required number of leaves is reached. The expected Sackin score for a tree with N leaves is calculated under the Equal Rates Markov model as:

\[ EY[S(N)] = 2N \sum_{i=1}^{N} 1/i \]

In contrast, under the uniform or Proportional to Distinguishable Arrangements model, all trees with the same number of leaves are assumed to be equally likely. This is not strictly a model of evolution but merely of tree growth. The expected Sackin
score under the uniform or Proportional to Distinguishable Arrangements model $EU[S(N)]$ can be calculated as a hypergeometric function $3F2$ (Mir et al., 2013), which for large values of $N$ is asymptotic to $\sqrt{2}\sqrt{N}/2$.

3 Results

We developed ContTest, a benchmark for large protein MSAs based on the accuracy of de novo contact map prediction. ContTest contains 136 test cases with between 1467 and 43,910 sequences (median 7098) and an average length of between 31.9 and 370 residues (median 113.6) (see Supplementary Table S1 and Fig. S1). We used the benchmark to score alignments made by Clustal Omega (Sievers et al., 2011), Kalign 2 (Lassmann et al., 2009), MAFFT (Katoh and Standley, 2013), MUSCLE (Edgar, 2004), Clustal W2 (Larkin et al., 2007), Kalign 1 (Lassmann and Sonnhammer, 2005) and hmmt (Eddy, 1998). Additionally, the Pfam database contains MSAs of all sequences in each protein family, and we used the benchmark to score the full alignments from Pfam 27. Clustal Omega, Kalign 2, MAFFT and hmmt were all run with default parameters. To make alignments in a reasonable time, Kalign 1 was run with the fast heuristic `-f` option, MUSCLE was limited to two iterations and Clustal W2 reused the guide tree generated by Clustal Omega. Benchmark scores for these alignment methods are listed in Table 1.

At the top of the rankings are the hmmt, Kalign 1 and Pfam alignments. These results are striking and unexpected. Since the method by which the Pfam alignments are created is not directly applicable to general MSA problems, the quality of unexpected alignments has not previously been assessed in MSA benchmark studies. Similarly, neither Kalign 1 nor hmmt have, to our knowledge, been considered in any recent benchmarking studies, as Kalign 1 is superseded by Kalign 2 and hmmt is not included in recent versions of HMMER.

The Pfam alignments are created in two steps. First, a manually curated ‘seed’ alignment is created from a small set of sequences

Table 1. Benchmark scores for alignments from Pfam and seven MSA packages

| Alignment method | Mean PSICOV precision | Mean EVfold-mdDCA precision | ContTest score |
|------------------|-----------------------|-----------------------------|----------------|
| hmmt             | 0.526                 | 0.590                       | 0.551          |
| Kalign1 (fast)   | 0.527                 | 0.581                       | 0.550          |
| Pfam             | 0.535                 | 0.577                       | 0.545          |
| Kalign 2         | 0.507                 | 0.533                       | 0.513          |
| Clustal Omega    | 0.497                 | 0.390                       | 0.428          |
| MAFFT (default)  | 0.433                 | 0.379                       | 0.396          |
| Clustal W2       | 0.413                 | 0.373                       | 0.381          |
| MUSCLE (2 iterations) | 0.445           | 0.329                       | 0.371          |

Mean PSICOV precision and mean EVfold-mdDCA precision are the mean precisions of the top 1/L long range contacts predicted by PSICOV and EVfold-mdDCA, respectively. ContTest Score is the geometric mean of the PSICOV and EVfold-mdDCA precisions. Statistical significances are indicated for the differences in consecutive pairs of ContTest scores. * $P < 0.007$; ** $P < 0.0014$; *** $P < 0.00014$ (0.05, 0.001 and 0.001 with Bonferroni correction for seven tests, respectively); NS, not significant.

3.1 Varying alignment parameters

To verify that the ContTest score correctly ranks alignment methods by accuracy, we created alignments using different parameter sets of the same MSA packages. For a given alignment package, we expect that certain parameter sets produce more accurate alignments than others, on average. These relationships are reflected in the benchmark scores (Table 2).

Both Clustal Omega and MUSCLE are capable of iteratively refining MSAs and using more iterations results in an increase in benchmark score for these methods (Table 2). In addition, Clustal Omega can use an external HMM to guide alignments. Using the Pfam HMMs for each family during alignment also increases the benchmark score for Clustal Omega. The NW-NS-PartTree-1 parameter set of MAFFT is recommended for making fast alignments of large numbers of sequences and is expected to have a lower average

![Fig. 1. Flowchart of benchmark process for one test case. Sequences from Pfam are aligned with the method of interest and the resulting MSA is used to predict residue–residue contacts for one of the proteins in the alignment. The 3D coordinates of this target protein are used to calculate the true residue–residue contacts. The two lists of contacts are compared to calculate a score for the alignment](image-url)
3.2 Adding errors to alignments decreases the benchmark score

As a further test of benchmark accuracy, we studied how the addition of deliberate errors to a set of alignments affects the benchmark score. First, we selected random subsets of sequences in each Pfam alignment to ‘misalign’ by adding a gap to the start of the aligned sequence, while truncating the final character (see Section 2.5). We observed that greater the percentage of sequences in each alignment that were misaligned, the greater the drop in contract prediction precision from the starting alignments. Figure 3 shows alignment that were misaligned, the greater the drop in contract precision (Supplementary Fig. S6).

3.3 Adding HOMSTRAD sequences to alignments allows a second measurement for alignment quality

To compare the ContTest benchmark method with the more traditional structure-superposition benchmark method, we created a hybrid benchmark by adding HOMSTRAD reference sequences to the Pfam sequences in 80 test cases for which we identified a match between the Pfam family and a HOMSTRAD alignment. We then calculated both the ContTest score of each alignment and the SPS for the embedded HOMSTRAD sequences. Scores for the HOMSTRAD sequences are listed in Table 3. A plot of the ContTest score against the mean HOMSTRAD SPS is shown in Supplementary Figure S7. The SPS and the ContTest scores notably disagree on Clustal Omega, which is the most accurate method as ranked by SPS. MAFFT NW-NS-PartTree-1 and Kalign 2 also differ slightly in their rankings between the two benchmark scores. It is unclear why Clustal Omega is ranked so differently by the two accuracy measures. Since the SPS measures only the number of correctly aligned residue pairs in an alignment, it does not penalize ‘overalignment’, i.e. false-positive aligned residues in the MSA. It is possible that MSA methods which are tuned for sensitivity over specificity may be ranked higher when measured by SPS or total column score than when ranked by ContTest or other accuracy measures.

3.4 ContTest rankings differ from PREFAB rankings

PREFAB (Edgar, 2004) is a structure-superposition-based MSA benchmark containing 1682 test cases. Test cases each contain two sequences for which a reference alignment is available and up to 48 other homologs. We calculated the SPS on the PREFAB benchmark for all MSA methods for which we previously calculated ContTest scores (Supplementary Table S4). The rankings of alignment methods differ greatly between ContTest and PREFAB, with hmmt and ‘chained’ alignments scoring particularly poorly on PREFAB, while...
being among the highest scoring methods when ranked by ContTest. The difference in ranking is unsurprising given the large difference in numbers of sequences between PREFAB and ContTest. ‘Chained’ guide trees have previously been shown to be effective only for alignments of large numbers of sequences (Boyce et al., 2014). Meanwhile, the greater sequence diversity present in the larger ContTest test cases is likely to facilitate the HMM training at the core of hmmt’s alignment process. It should be noted that for alignments of tens of sequences, as in PREFAB, many slower, more accurate MSA methods are available which cannot make MSAs of the size required for the ContTest benchmark and thus were not included in our study.

4 Discussion

ContTest is the first protein MSA benchmark to realistically test alignments of large numbers of sequences and base scores on empirical biological data. Although structure-based benchmarks may contain alignments with many sequences, e.g. HomFam (Blackshields et al., 2010), scores are always based on reference alignment of a small subset of sequences contained in the MSA. This could allow large errors in non-reference sequences to go unpunished. In addition, the structure-based reference alignment must itself be created, and this alignment depends on heuristic structure comparison algorithms. Meanwhile, the phylogeny-based species-discordance benchmark of Dessimoz and Gil (2010)—in which phylogenetic trees are inferred from MSAs and compared with a known reference tree—is restricted to testing alignments of six sequences. Simulation-based benchmarks allow for arbitrary numbers of sequences in test alignments but depend on simplified models of evolution and may not realistically model gap placement in real proteins. Our benchmark method uses information from all sequences in the input alignments. Test cases must contain a minimum of about 1000 sequences, so there is sufficient information in the alignment to make contact predictions, but there is no practical upper limit on the number of sequences.

Our benchmark scores confirm the utility of completely chained guide trees, as described by Boyce et al. (2014). That result has been controversial (Tan et al., 2015) but, at least for alignments of many sequences from structurally conserved regions, the results appear clear and robust. Alignments made with fully or even largely chained guide trees consistently outscore alignments made with more traditional guide trees. Each of the alignment packages that accepts user-specified guide trees benefited from using chained guide trees, and the increase in score due to chained guide trees was greater than the increase due to extra iterations of Clustal Omega or MUSCLE or using a high quality external HMM with Clustal Omega. Our scores also demonstrate the quality of the alignments in the Pfam database, which may have previously been neglected, as well as suggesting that both Kalign 1 and hmmt be
revisited as useful methods for creating alignments of large numbers of sequences.

We have shown that the ContTest benchmark is in general agreement with structure-based benchmarks for large numbers of sequences. However, we observe that the methods that produce the most accurate alignments for large numbers of sequence do not necessarily perform well for small number of sequences, as measured by the PREFAB benchmark (Supplementary Table S4). ContTest is likely to be most sensitive to alignment errors in regions which are structurally conserved but without very high levels of sequence conservation. That is, regions of the protein in which amino acid substitutions are accepted but must be compensated for at co-evolving sites. A limitation of this benchmark is that all alignments inherently involve only sequences of a single protein domain, without any large insertions. It is likely that certain methods might perform poorly on more general alignment problems involving multiple domains. In particular, for hmmt alignments, it is necessary that a sensible HMM can be trained on the input sequences, which may not be possible in the general case. However, these are problems that may be addressed by future benchmarks, and alignments of single protein domains are common and important in many applications.

Finally, these results give some clear pointers toward the best methods for making MSAs intended for use with de novo structure prediction methods and show that choice of alignment method can have a large influence on the quality of contact predictions, even when using the same software package with different parameters.

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