

A benchmark of multiple sequence alignment programs upon structural RNAs

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Abstract

To date, few attempts have been made to benchmark alignment algorithms upon nucleic acid sequences. Frequently, sophisticated PAM or BLOSUM like models are used to align proteins, yet equivalents are not considered for nucleic acids; Instead, rather ad hoc models are generally favoured. Here we systematically test the performance of existing alignment algorithms on structural RNAs. The goals of this work are: (1) To determine conditions where it is appropriate to apply common sequence alignment methods to the structural RNA alignment problem. This indicates where and when researchers should consider augmenting the alignment process with auxiliary information, such as secondary structure. (2) To determine which sequence alignment algorithms perform well under the broadest range of conditions. We find that sequence alignment alone, using the current algorithms, is generally inappropriate below 50-60% sequence-identity, secondly we note that the probabilistic method ProAlign and the aging Clustal algorithms generally out-perform other sequence-based algorithms, under the broadest range of applications.

Introduction

Motivation

The use of multiple sequence alignments is an essential step for many RNA sequence analysis methods (e.g. RNA structure analysis [1–5], RNA homology search [6, 7], ncRNA detection [8, 9] and RNA based phylogenetic inference [10, 11]). Structural alignment of RNA is however an open problem. An algorithm for simultaneous structural RNA sequence alignment, structure prediction and phylogenetic reconstruction has been proposed [12], yet current implementations are limited in terms of functionality and sequence size [13–18]. A second structural RNA alignment approach employs (predicted) structures and aligns these directly [19–21], again these are generally limited in terms of sequence size, but primarily suffer from the inaccuracy of single sequence structure prediction [22–24] (although, pruning low probability base-pairs yields modest improvement [25, 26]). In addition, algorithms which attempt to include RNA structural information for sequence alignment fail when structure is not conserved (e.g. methylation-guide snoRNAs, Air-RNA). Many of these methods are also impractical when sequence length is large or when single sequence structure prediction [27, 28] performs poorly [24].

The performance of current sequence alignment methods has been thoroughly analysed in terms of protein alignment accuracy [29–32]. Benchmarking has also been performed for simulated non-coding DNA [33]. However, the results of these studies do not explore methods for the specific problem of aligning structural RNAs. In this work we extend these studies and test the performance of current alignment algorithms upon structural RNA data-sets.

The aims of this work are two-fold. Firstly, to identify the “twilight zone” of RNA sequence alignment - the homology range below which sequence alignment alone is unlikely to produce reliable results and researchers should seriously consider augmenting the alignment process with auxiliary information such as secondary structure. Secondly, to identify algorithms capable of reliably aligning structural RNA sequences under a range of sequence identi-

ties

Alignment algorithms

The simplest form of an alignment is the pairwise sequence alignment. This can be performed by aligning sequences globally [34] or locally [35], both employ dynamic programming, thus resulting in a quadratic time complexity. Different scoring schemes may be used, which already produce varying results. The alignment of multiple sequences is far more complex, as the mathematically optimal solution imposes exponential complexity. Therefore heuristics are used, which do not guarantee an optimal solution, but perform multiple sequence alignment in reasonable time.

One common approach is called progressive alignment [36], which builds a multiple alignment from pairwise alignments. The idea is that an alignment of sequences, which have more recently diverged, is more likely to be reliable. Thus high scoring pairwise alignment are aligned first and next closely related sequences (or alignments of sequences) are added progressively. The order of this progressive alignment is, in most cases, defined by a guide tree, which is created beforehand from a distance matrix, produced by aligning all $\frac{n(n-1)}{2}$ possible pairs of sequences first. The basic drawback of this method is the fact that gaps introduced in an early step cannot be removed during the later addition of sequences, e.g. errors made in an early step propagate during the alignment process (“Once a gap always a gap”). So called iterative methods prevent this by realigning sequences or sequence groups in the multiple alignment, thus, theoretically optimising the alignment until successive iterations fail to improve the alignment or reach a predefined limit (convergence).

Another class of algorithms is called consistency based. Here a multiple alignment is constructed by extracting (maximum-scoring) pairwise alignments from a library such that these combined pairwise solutions are not contradictory or mutually exclusive.

Probabilistic methods are an increasingly popular way of generating solutions to biological problems. The basic premise is to produce a model that one believes best describes the system behaviour. Model parameters are sub-

sequently estimated from reliable data. In terms of sequence alignment a pairwise HMM based approach has been proposed [37], and now implemented and extended to multiple sequence alignment [38,39].

The structural RNA alignment approach of Sankoff [12], merges the recursions of Smith-Waterman [35] type sequence alignment and Nussinov (maximal base-pairing) [40] or Zuker-Stiegler (energy based) [41] RNA structure prediction [15]. The basic idea is to implicitly include base-pairing interactions into the alignment procedure such that homologous base-pairs are aligned correctly. Unfortunately, the algorithm is computationally expensive ($O(n^{3m})$ in time, and $O(n^{2m})$ in space, where n is the sequence length and m is the number of sequences). Current implementations, Dynalign [13,14], Foldalign [16], PMcomp [15] and Stemloc [17,18], are restricted implementations of the Sankoff-algorithm which impose practical limits on the size or shape of substructures. Additionally, sensible score routines such as thermodynamics [13,14], a combination of sequence and thermodynamic scores [16] and partition-function derived probability-matrices [15], can be used to score alignments.

Figure 1 shows a classification of all the programs used in this study into the above categories.

Materials and Methods

Accuracy measures

In order to evaluate alignment methods on structural RNAs we use two independent measures. The first is the traditional sum-of-pairs score (SPS) employed in many previous alignment benchmarks [29,32,33]. SPS is defined as the fraction out of all possible character pairs that are aligned in both the predicted and reference alignments. Perfectly predicted (concordant) alignments receive an SPS of one, absolutely imperfectly predicted (discordant) alignments receive an SPS of zero. Although, for nucleotide alignments an SPS of zero is rarely observed. Essentially, SPS provides a measure of the sensitivity of the prediction. The second measure, dubbed the “structure conservation index” (SCI) provides a measure of conserved secondary-structural

information contained within the alignment [9]. It is a derivative of the score calculated by the RNAalifold consensus folding algorithm [5, 42] which is based upon the sum of a thermodynamic and covariance term and, in contrast to the SPS, is independent from a reference alignment. The SCI is close to zero if RNAalifold identifies no common RNA structure in the alignment, whereas a set of perfectly conserved structures has a $SCI \approx 1$. A $SCI > 1$ indicates that there is a conserved RNA secondary structure which is, in addition, supported by compensatory and/or consistent mutations preserving the common structure. We want to note that the SCI scores alignment accuracy only in terms of secondary structure information. For example, if the helices of a secondary structure are accurately aligned, it does not affect the SCI whether the loop regions are well aligned in terms of sequence similarity. The SCI specifically points out the structural aspect of alignment accuracy and therefore appears to be a useful measure in addition to the SPS scores.

Alignment programs

We tested 11 sequence alignment programs and 4 structural alignment programs (see supplementary materials, Table 1). Where previous alignment benchmarks have only considered the default or “out of the box” behaviour we try testing a range of parameter combinations for each alignment method. Algorithm options are summarised in the supplementary materials (see Tables 2 & 3).

Test data-sets

We generated four diverse structural RNA data-sets of Group II introns, 5S, rRNA, tRNA and U5 spliceosomal RNA. The sequences and the reference alignments for calculating the SPS score were obtained from the Rfam v5.0 database. SRP from the SRPDB database was included in the original data-set but was later discarded due to poor comparability between predicted and structural alignments contained in this data-set (the results suggest that a fraction of the SRP reference sequences have been mis-aligned). Using the same procedure as described previously [42], we generated approximately 100 sub-alignments for each family. The alignments contained five sequences

each and encompassed a range of sequence identities. This large data-set (hereafter referred to as data-set 1) was divided into high ($\geq 75\%$ sequence-identity, 73 alignments), medium ($< 75\%$ and $\geq 55\%$ sequence-identity, 73 alignments) and low ($< 55\%$ sequence-identity, 242 alignments) sequence homology groups. An additional tRNA data-set was generated with just two sequences to each alignment (hereafter referred to as data-set 2). This was used to contrast pairwise structural alignment methods and sequence alignment methods.

Caveat: tools improve

We note here that our data reflect the state of the art in early 2005. Most of the tools tested are relatively recent, and frequently still under development. Hence, not all the below observations will remain reproducible. In fact, we hope this study helps to obtain better results in the future.

Results

Data-set 1: applicability of pure sequence alignment to ncRNA sequences

All 11 sequence alignment algorithms were tested upon data-set 1. The results and relative algorithm ranks for each homology group are summarised in Table 1. We experimented with a variety of algorithm parameters. The results using default and any parameter combinations that increased algorithm performance are summarised in Figure 2. In order to measure relative algorithm performances a ranking for each algorithm (and parameter setting) was calculated within each of the three homology ranges. The rank is based upon the product of mean SCI and mean SPS scores.

For the high similarity data-sets (sequence-identity $\geq 75\%$), there is little difference in accuracy across most of the algorithms considered here (see Table 1). Align-m and Handel rank well on this data-set, yet the relative performance of both these methods dropped rapidly with decreasing sequence homology. Interestingly, for Align-m, this is the opposite to what has been observed for protein-based results [43].

PCMA ranked well on both the high and medium similarity data-sets, however the relative performance of this method dropped on the low similarity data-set.

ClustalW, MUSCLE, PCMA, POA (with both global and progressive modes - hereafter referred to as POA (gp)), ProAlign, Prrn and T-Coffee (when ClustalW is used to generate a library of pairwise alignments) perform comparatively well across all homology ranges, with little significant variation between these methods. Only ClustalW, ProAlign and POA (gp) consistently ranked in the top 10 across all the data-sets. There is some redundancy in ranking algorithms over all the combinations of algorithm parameters we use here, obviously some combinations produce similar (or even identical) results. However, this has had little impact on our conclusions.

The results suggest that $\sim 60\%$ sequence-identity is a crude threshold, whereby the structural content of predicted sequence alignments diverges from reference structural alignments (see Figures 2&3).

Data-set 2: comparison of structural and sequence methods

Now we contrast the relative performance of the comparatively good sequence based methods identified in the previous section with structural alignment methods using a smaller tRNA data-set. The structure based methods are generally computationally more intensive than the sequence based methods - hence the small size (in terms of the number of sequences and the sequence length) of this data-set.

We use data-set 2 to compare the relative performances of structure-based methods (e.g. Dynalign, Foldalign, PMcomp and Stemloc) to the “better” sequence based methods identified in the previous section (e.g. ClustalW, MUSCLE, PCMA, POA (gp), ProAlign and Prrn). We observe a dramatic divergence in relative performances below $\sim 60\%$ sequence-identity between the structure and sequence based methods (see Figure 3).

The structural methods Dynalign, Foldalign and PMcomp show high conservation of structural information (SCI) across all homology ranges. How-

ever, the SPS of Dynalign and PMcomp are significantly lower than that of Foldalign. The difference is most marked in the high to medium homology range. This is possibly because the current versions of Dynalign and PMcomp optimise scores solely based on secondary structure information and hence are likely to produce rather different alignments to those used in the test data-set, where sequence information is also included. Stemloc performs comparatively well in terms of SCI for sequence-identities above 40%. In terms of SPS however, Stemloc behaves much like the purely sequence-based methods. Given the apparent sophistication of this method and large computational resources required to run the algorithm [17,18] these results are rather disappointing. We hypothesise perhaps these are due to an overemphasis upon the sequence based component of the algorithm.

All of the above data-sets are freely available from:

<http://www.binf.ku.dk/users/pgardner/bralibase/>. As novel and updated algorithms become available updated results will also be made available from the webpage.

Discussion

An aim of this work is to determine the boundaries between when pure sequence alignment methods perform well and when augmentation of the alignment with structure is necessary. We wish to highlight that benchmarks based purely upon structural protein alignments do not adequately test all the uses of sequence alignment. Also, we are pleased to note that our two independent measures of alignment fitness (SCI and SPS) produce similar results.

In some cases we found altering algorithm parameters produced a dramatic improvement over the defaults (e.g. T-Coffee performance improves using Clustal to generate a library of pairwise alignments and POA performance improves dramatically using a combination of the global and progressive modes).

We find the conclusions of previous studies based upon structural protein alignments do not necessarily hold for the alignment of structural ncRNA. For example, DIALIGN, identified as a method which performed well for low

homology protein alignment did not generally improve (relative to the alternative methods) on low homology data-sets [32]. Another surprising discovery was that T-Coffee, touted as an excellent method for high homology data-sets, did not perform well (again, relative to the alternative methods) on the ncRNA data-sets [32]. Another surprise was that the supposedly out-dated, yet still widely used method ClustalW, performed consistently well across all homology data-sets. This is possibly a consequence of the fact that more recent algorithms are heavily optimised for protein alignment. The relatively new methods ProAlign, POA (gp) and MUSCLE also performed consistently well. ProAlign in particular produced (comparatively) reliable alignments and ranked in the top 5 across all homology ranges. This is possibly due to the fact that ProAlign is one of the few algorithms to use a scoring scheme derived from reliable nucleic acid sequence alignments. The performance of POA (gp) is also remarkable, not only because it employs a very fast method (said to accurately align 5,000 EST sequences in 4 hours on a Pentium II [44]) but also because it performed consistently well over all test-sets.

Another conclusion of this work is that the “twilight-zone” of ncRNA alignment - the homology-range where little-to-no structural information of predicted alignments (using the current state of the art algorithms) for structurally homologous sequences is retained - is in the 50-60% sequence-identity range. This is dramatically higher than that of protein sequences which is 10-20% [29]. Much of this difference is, of course, due to the different alphabet sizes and the generally limited models and score-matrices for nucleotide alignment.

It is interesting to note that three of the structural methods (Dynalign, Foldalign and PMcomp) for a short homology range (40-60% sequence-identity) have higher SCI scores than the reference alignment and that in these same regions there is a dip in the performance when Dynalign, Foldalign and PMcomp performance is measured using SPS. This suggests that the reference alignments themselves may be improved upon in this homology range.

Based upon these results the Foldalign score routines seem to have optimised the delicate balance between the sequence and structure based scores.

This implementation of Sankoff’s algorithm employs a light-weight energy model [13, 41, 45, 46] in concert with substitution matrices similar to those of RIBOSUM [47] and BLOSUM [48] which seem to produce excellent predictions. However, the computational complexity of this algorithm is still an issue, global alignment is restricted to sequences of approximately 200 nucleotides or less, in practice. Further optimization may increase this bound, however.

The profile based approach of Hofacker et al. (`pmcomp --fast` [15, 49]), holds promise for producing fast and reasonably accurate alignments in satisfactory time across all homology ranges. It by no means produces “optimal” alignments in terms of sequence or structure, but is a reasonable compromise between the sequence and structure based methods in terms of improved accuracy for the former and dramatically reduced computational requirements for the latter. This method is in the process of being re-implemented in C with affine gap costs and an adjustable sequence-weighting parameter. This is available as “RNApaln” with the Vienna package version 1.5 or greater (pers. commun. Ivo Hofacker).

Summary

The results and main conclusions of our study can be summarised as follows:

- The two independent measures of global alignment accuracy SPS and SCI are generally in agreement. These measures only differ significantly on methods, such as Dynalign and PMcomp, that perform only structural alignment. The SCI is, therefore, a useful score for assessing the accuracy of structural RNA alignments.
- The relative performance of multiple sequence alignment programs on RNA alignments can differ remarkably from the performance observed on protein alignments.
- The multiple sequence alignment algorithms ClustalW, MUSCLE, PCMA, POA (gp), ProAlign and Prn perform well on high-to-medium homology data-sets.
- ClustalW, ProAlign and POA (gp) consistently ranked in the top 10 across

all homology ranges.

- The “twilight-zone” of ncRNA alignment is in the 50-60% sequence-identity range.
- Below this limit, algorithms incorporating structural information (Dynalign, Foldalign, PMcomp, Stemloc) outperform pure sequence based methods. However, these algorithms are computationally demanding which severely limits their use in practise.

Future directions

One rather interesting result of this study is that the structure profile alignment method (`pmcomp --fast`) produces reasonable structural alignments across all homology ranges in a dramatically short time. This method in combination with, as yet undeveloped, iterative alignment refinement strategies seems poised to become a method of choice for RNA researchers in the near future. This also has interesting implications for the notoriously difficult problem of ncRNA homology search. A combination of a database of locally stable regions [50] (analogous to the index creation of the BLAST procedure) and the profile alignment method is likely to produce a superior homology detection tool. This application and the extension of this method to multiple alignment is an area of active research.

Other potentially fruitful research areas to explore are: (1) The implementation of light-weight Sankoff-like algorithms, which produce reasonable alignments in a short time-frame and use score routines combining energy and sequence scores similar to those of Foldalign. (2) In analogy to the improvement of structure prediction accuracy by including stacking parameters (nearest-neighbour model), perhaps the alignment of RNA sequences over a di-nucleotide alphabet could produce improvements in sequence-based alignment. (3) The current score matrices for nucleotide alignment are generally *ad hoc*, it is likely that significant improvements could be gained by using RIBOSUM-like matrices [47] for scoring alignment. (4) “Intelligent” alignment algorithms which employ sequence information when this is reasonable, or structure alignment when this is better.

Note added to proof

Since embarking on this project the alignment algorithms Align-m, Handel and MAFFT have been updated. Preliminary analysis of the updated algorithms suggests that improvements to Align-m and Handel have resulted in modest performance increases across the high, medium and low similarity groups of data-set 1. The improvements to MAFFT however have resulted in major performance increases across all similarity groups of data-set 1. In fact, across all similarity groups MAFFT (ver. 5) now ranks second only to ProAlign. However, gap-parameters for this algorithm have been estimated directly from data-set 1, this bias could be alleviated by determining optimal gap-parameters for all methods prior to benchmarking. Preliminary work in this direction shows algorithm performances on RNAs can in some cases be enhanced by optimising gap-parameters (personal communication K. Katoh).

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Figure legends

Figure 1

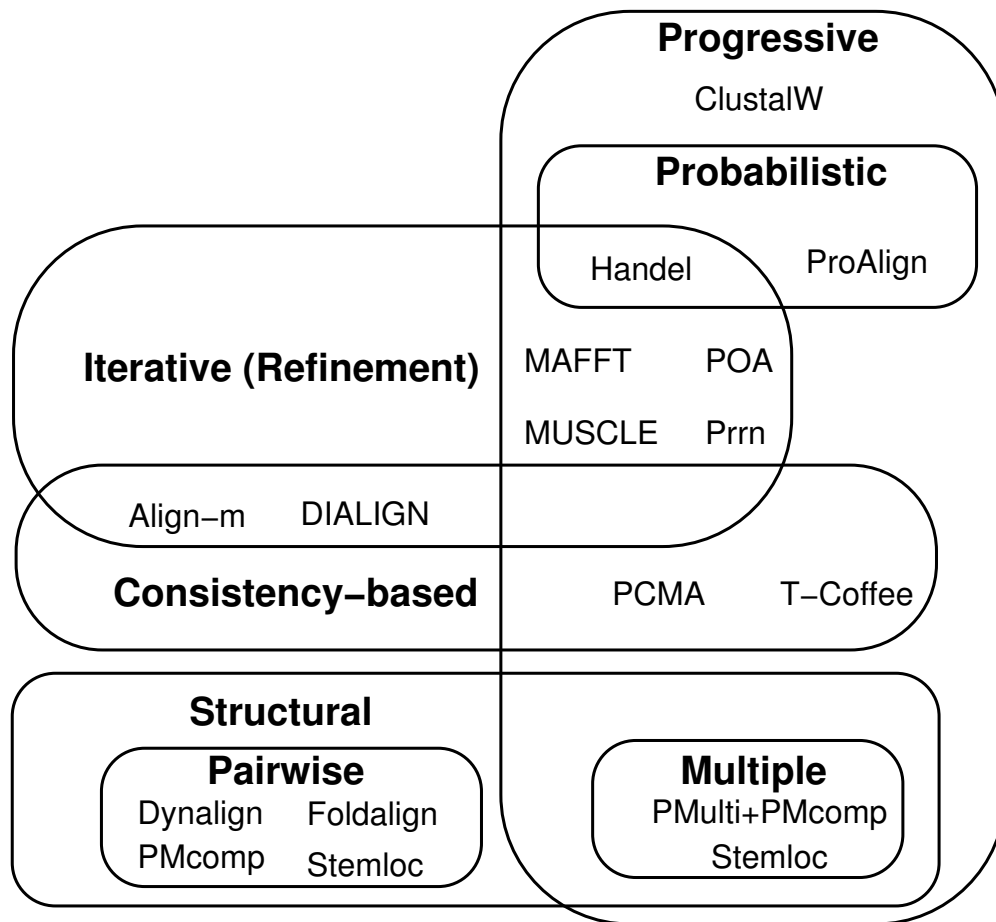


Fig. 1. An overview of alignment programs used in this work. Programs were classified into the categories described in detail in the “Alignment algorithms” section.

Figure 2

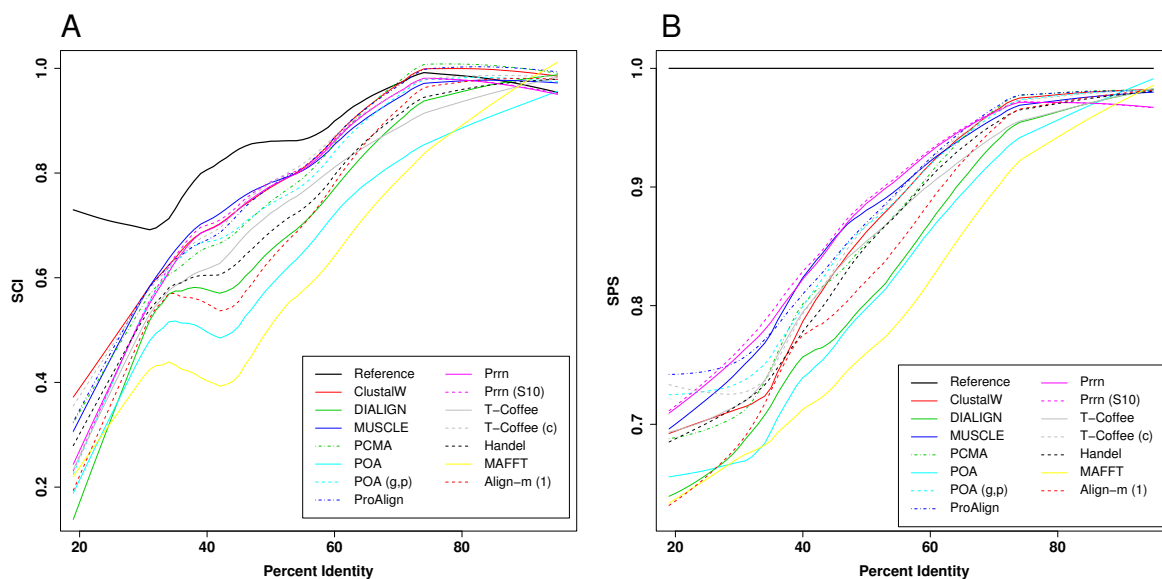


Fig. 2. Figure **A** displays the structure conservation index (SCI), figure **B** displays sum-of-pairs score (SPS). Both measures of structural RNA alignment correctness are plotted as functions of the mean-pairwise-sequence-identity (calculated using the reference alignments). The curves are fit to data-set 1 (see text for details) using lowess (Local Weighted Regression) smoothing. At most, two curves are plotted for each alignment package - one corresponding to the default parameters, the other corresponds to best parameter combination we could identify.

Figure 3

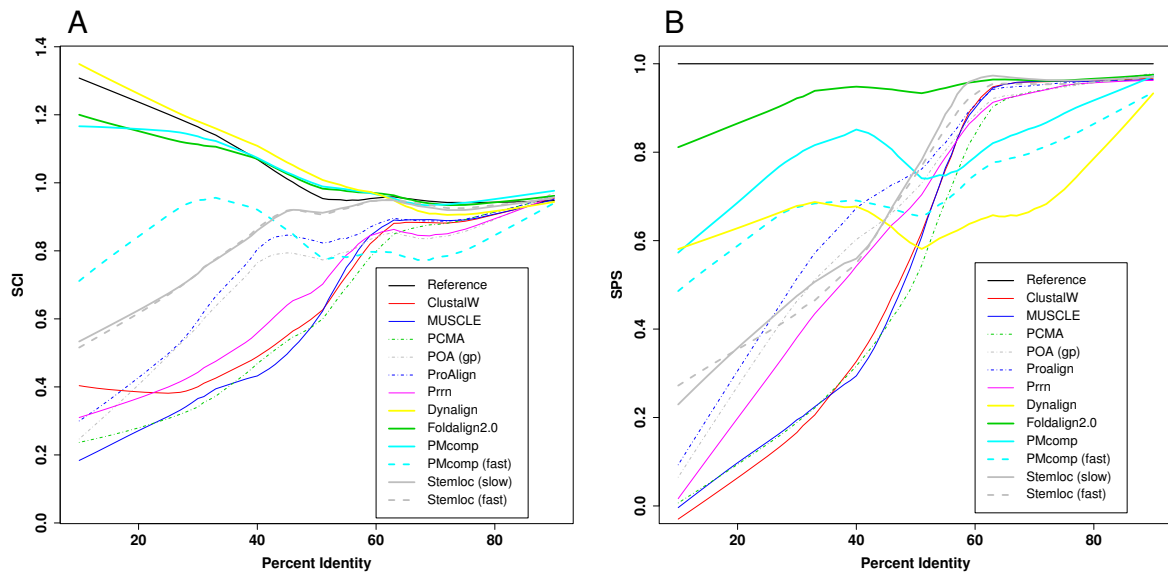


Fig. 3. SCI (figure **A**) and SPS (figure **B**) as functions of the sequence-identity for data-set 2 (see text for details). Five structural algorithms are shown; The Sankoff-based methods Dynalign, Foldalign, PMcomp and Stemloc and the base-pair-probability-profile alignment method implemented in PMcomp (fast). These are contrasted with hand-curated-structural alignments and six of the better sequence-based alignment algorithms (ClustalW, MUSCLE, PCMA, POA (gp), ProAlign, Prn).

Tables

Algorithm	High homology			Med. homology			Low homology		
	75% \leq Seq. Id.			75% $<$ Seq. Id. \leq 55%			Seq. Id. $<$ 55%		
	SCI	SPS	Rank	SCI	SPS	Rank	SCI	SPS	Rank
Structural	0.9789	1.0000	0	0.9297	1.0000	0	0.7846	1.0000	0
Align-m (1)	0.9827	0.9600	5	0.8453	0.8825	22	0.4957	0.6748	25
Align-m (2)	0.9827	0.9600	6	0.8453	0.8825	21	0.4957	0.6748	24
Align-m (3)	0.9778	0.9593	7	0.8040	0.8742	26	0.4691	0.6635	29
Align-m (4)	0.9778	0.9593	8	0.8040	0.8742	25	0.4691	0.6635	28
Align-m (5)	0.8995	0.9419	30	0.7597	0.8583	30	0.4777	0.6460	30
Clustal	0.9438	0.9741	9	0.9100	0.9194	2	0.6064	0.7423	8
Clustal (qt)	0.9315	0.9743	12	0.8996	0.9123	9	0.6076	0.7345	9
DIALIGN	0.9071	0.9577	27	0.8018	0.8712	27	0.4979	0.6659	26
DIALIGN (o)	0.9077	0.9601	26	0.8568	0.8860	20	0.5202	0.6721	23
DIALIGN (it)	0.8590	0.9491	34	0.7519	0.8556	31	0.4669	0.6546	32
DIALIGN (it,o)	0.8492	0.9486	35	0.7092	0.8456	33	0.4467	0.6467	33
Handel	0.9604	0.9560	11	0.8570	0.8954	19	0.5360	0.7283	19
MAFFT (fftns)	0.8321	0.9145	36	0.5864	0.8030	37	0.3538	0.6448	37
MAFFT (fftinsi)	0.8840	0.9427	32	0.6655	0.8378	36	0.3845	0.6634	36
MAFFT (nwns)	0.9297	0.9502	25	0.6712	0.8349	35	0.3941	0.6724	35
MAFFT (nwnsi)	0.9330	0.9526	22	0.7004	0.8451	34	0.4071	0.6812	34
MUSCLE	0.9222	0.9684	19	0.8988	0.9181	5	0.6065	0.7668	2
MUSCLE (nj)	0.9268	0.9707	16	0.8841	0.9110	17	0.5902	0.7503	12
MUSCLE (mi32)	0.9222	0.9683	21	0.8959	0.9167	8	0.6069	0.7666	1
MUSCLE (mi32,mt6)	0.9222	0.9683	20	0.8959	0.9167	7	0.6068	0.7664	3
MUSCLE (nj,mt6)	0.9268	0.9707	15	0.8841	0.9110	16	0.5902	0.7503	11
MUSCLE (nj,mi32)	0.9268	0.9708	14	0.8855	0.9112	14	0.5897	0.7501	14
MUSCLE (nj,mi32,mt6)	0.9268	0.9708	13	0.8855	0.9112	13	0.5898	0.7501	13
PCMA	1.0030	0.9635	3	0.9196	0.9059	3	0.5339	0.6890	20
PCMA (agi20)	1.0030	0.9635	2	0.9255	0.9068	1	0.5621	0.7058	16
PCMA (agi60)	1.0030	0.9635	1	0.8938	0.8941	18	0.5270	0.6827	21
POA	0.8478	0.9644	33	0.7666	0.8739	29	0.4656	0.6740	27
POA (g)	0.9253	0.9722	17	0.8836	0.9130	15	0.5581	0.7543	15
POA (p)	0.8668	0.9656	31	0.7814	0.8814	28	0.5079	0.6964	22
POA (gp)	0.9444	0.9726	10	0.8929	0.9188	10	0.5843	0.7726	6
ProAlign (bw400)	0.9978	0.9631	4	0.9163	0.9072	4	0.6045	0.7490	5
Prrn	0.9364	0.9458	24	0.9036	0.9064	11	0.5903	0.7549	10
Prrn (S10)	0.9371	0.9458	23	0.8997	0.9086	12	0.5964	0.7596	4
T-Coffee	0.8867	0.9656	29	0.8129	0.8989	24	0.5322	0.7337	18
T-Coffee (c)	0.9201	0.9733	18	0.8956	0.9194	6	0.5972	0.7543	7
T-Coffee (f)	0.8867	0.9656	28	0.8129	0.8989	23	0.5322	0.7337	17
T-Coffee (s)	0.7892	0.9536	37	0.7151	0.8637	32	0.4447	0.6934	31

Table 1

The table displays mean SCI and mean SPS scores computed for data-set 1 (see text for details). Data-set 1 has been divided into 3 homology groups: the high-homology group ($75\% \leq \text{seq. id.}$), the medium-homology group ($75\% < \text{seq. id.} \leq 55\%$) and the low-homology group ($\text{seq. id.} < 55\%$). Rankings are computed from the product of SCI and SPS. The top ten ranks are highlighted in bold. Abbreviations of the parameter switches used to produce these results are shown in parentheses - further details of these can be found in supplementary materials.