COS 551: Introduction to Computational Molecular Biology Lecture 4: September 26, 2000 Lecturer: Mona Singh Scribe: Aneel Delawalla

## Multiple Sequence Alignments I

**Definition:** Given *k* strings,  $S_1$ ,  $S_2$ ,  $\ldots$ ,  $S_k$ , a multiple sequence alignment (MSA) is obtained by inserting gaps in the strings to make them all the same length.

E.g., the following is a MSA of 4 sequences MQPILLLV, MLRLL, MKILLL, and MPPVLILV.



No column may be all gaps.

Multiple sequence alignments are used for many reasons, including:

- (1) to detect regions of variability or conservation in DNA or protein families,
- (2) to provide stronger evidence than pairwise similarity for structural and functional inferences,
- (3) to serve as the first step in phylogenetic reconstruction, in RNA secondary structure prediction, and in building profiles (probabilistic models) for protein families or DNA signals.

For pairwise alignments, we scored each column by looking at matches, mismatches, and gaps in the two sequences (in practice, protein sequences are scored using substitution matrices). However, now we have multiple characters in each column, and it is not obvious what the best way to score a column is. There are many possibilities. The sum-of-pairs (SP) is a common scoring scheme. Here, each column in an alignment is scored by summing the scores of all pairs of symbols in that column. The score of the entire alignment is then summed over all column scores. We will assume that a match  $= 1$ , a mismatch  $= -1$ , and a gap  $= -2$ . For example, the sum-of-pairs score of the  $4<sup>th</sup>$  column of the MSA given earlier is:

 $SP(I, -I, V)$  $=$  score(I,-) + score(I, I) + score(I,V) + score(-,I) + score (-,V) + score(I,V)  $= -2 + 1 + -1 + -2 + -2 + -1 = -7$ 

Although there is never an entire column of gaps, if we look at any 2 sequences in the alignment, there may be columns where both have gaps. The value of  $score(-,-)$  is set equal to 0; this eliminates double counting (or extra penalization) for the use of gaps.

Let MSA-SP denote the optimum (i.e., highest scoring) MSA under the SP scoring system.

Note that it is not at all clear that sum-of-pairs is the best scoring system to use. The SP scoring system tends to overweight contributions of differences from many very similar sequences.





The sum-of-pairs score for this column is  $3 - 3 = 0$ .

However, evolutionarily speaking, say the relationship between the sequences is:



Then a single  $C \longrightarrow A$  mutation can explain the data, and thus SP tends to overcount mutations.

**Note:** An optimum MSA-SP may not give the optimal pairwise alignments. For example, if a match = 1, a mismatch = -1, and a gap = -2, then the optimum MSA-SP for the sequences AT, A, T, AT and AT is as follows, with the induced alignment between A and T also shown.



However, the optimum pairwise arrangement for the two sequences A and T is  $^{-1}$ , which, if it were part of the MSA would yield:

A T

$$
\begin{array}{ccc}\nA & T \\
A & - \\
T & - \\
A & T \\
A & T \\
\hline\n2 & -9 \\
= -7\n\end{array}
$$

We now outline the algorithm for finding an optimum MSA under the SP measure. It is straighforward to see how to adapt it for other similiarity measures. However, this algorithm is usually not used in practice because it is computationally too time consuming. This approach uses dynamic programming.

Assume for simplicity that our *k* sequences  $S_1$ ,  $S_2$ ,  $\ldots$ ,  $S_k$  all have length *n*. If we were aligning 2 such sequences, we would need an array of size  $[n+1][n+1]$  to account for all possible solutions. Similarly, for *k* sequences, we need an array of size  $[n + 1]^k$ , or a *k*-dimensional array.

As with the pairwise alignments, let's look at the last column in the alignment. For each sequence, there are 2 possibilities: either there is a letter or a gap in the last character position. Taken for all *k* sequences, there are then  $2<sup>k</sup> - 1$  possibilities (we subtract 1 because the case of every sequence having a gap in the last character position is not allowed).

Thus, we are keeping track of a *k*-dimensional array sim, where  $\sin(\vec{i}) = \sin(i_1, i_2, ..., i_k)$ Then,

 $\sin(\vec{i}) = \max_{b,c} {\sin(\vec{i} - \vec{b})} + SP(\vec{c})$ , where  $\vec{b} \neq 0$ , over all binary vectors  $\vec{b}$  of length *k* 

$$
\vec{c}_j = \begin{cases} S_j[i_j], \text{ if } \vec{b}_j = 1 \\ \cdots \cdots, \text{ if } \vec{b}_j = 0 \end{cases}
$$

Vector  $\vec{b}$  enumerates over all  $2^k-1$  possibilities. That is, when  $\vec{b}$   $j = 1$ , we're considering the case where the  $i_j^{\text{th}}$  character of a sequence *j* is used. Vector  $\bar{c}$  keeps track of the actual characters used.

Therefore, we must:

- (1) fill out the entire array which has  $[n + 1]^k$  cells,
- (2) consider  $2^k 1$  possibilities for each cell in the array, and
- (3) compute the sum-of-pairs measure for each of these possibilities.

Thus, this algorithm would take roughly  $(n + 1)^k (2^k - 1)(k^2)$  time for the SP measure. This is impractical even for numbers as small as  $k = 10$ . Thus, most packages for multiple sequence alignment do not compute the optimum MSA. Instead, most packages use heuristics to compute multiple sequence alignments. The basic idea of these heuristics is to compute pairwise alignments and to merge alignments consistently.

**Note:** Given all pairwise alignments, it is usually impossible to arrange an MSA consistent with all of them.





Note, however, that given any *k*-1 alignments relating *k* sequences, there is always an MSA consistent with all *k*-1 pairwise alignments. That is, there is an MSA such that the induced pairwise alignments in the MSA include the given *k*-1 alignments.

With  $k = 5$ , there are 4 alignments relating all the sequences. For example, we could have pairwise alignments for the edges in the following graph.



We can use this fact to devise multiple sequence alignment heuristics.

**Star Alignment:** One such heuristic for MSA is to just add the sequences to an alignment, one by one, maintaining consistency and following the "once a gap, always a gap" rule. The twist in the star alignment heuristic is that you must (1) find the sequence that is the most similar to all the rest using pairwise alignment (see below) and (2) use it as the "center of a star" when aligning all other sequences.



Formally speaking, (1) is simply saying to find S*<sup>c</sup>* such that *i*≠*c*  $\sin(S_i, S_c)$  is maximized.

For example, say your sequences are:



Then, we first compute all pairwise alignments and scores. The following matrix gives the alignment scores for all pairs, as well as the sum of these scores for each sequence.

and



Now, let's build the alignment.



For consistency, once a gap is added, it is never removed.

The running time is dominated by computing the pairwise alignments. If *k* sequences are length *n*, then:

(1) we compute 2  $\frac{k(k-1)}{2}$  pairwise alignments (2) each alignment takes time  $n^2$ .

Thus, the running time for computing all pairwise alignments is  $O(k^2n^2)$ . If *l* is an upperbound on alignment lengths, then we can merge alignments in  $k^2 l$  time. Therefore, the total running time is  $O(kn^2 + k^2l)$ . Note that the Star Alignment does not optimize the SP criterion. Therefore, there is a trade-off between optimization and practicality. Actually, the star alignment method is not used much. In the next lecture, we'll discuss a similar method that is very widely used: the progressive alignment method of the ClustalW package. ClustalW uses the same basic method as the Star Alignment. However, it compares pairwise alignments and then adds individual sequences based on a given order of similarity.

## **Further Reading**

Setubal & Meidanis. *Introduction to Computational Molecular Biology*, PWS Publishing Company, 1997. Chapter 3.