Accelerating Computation of DNA Multiple Sequence Alignment in Distributed Environment

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Abstract
Multiple sequence alignment (MSA) is a technique for finding similarity in many sequences. This technique is very important to support many Bioinformatics task such as identifying Single Nucleotide Polymorphism (SNP), creating phylogenetic tree, and metagenome fragments binning. The simplest algorithm in MSA is Star Algorithm. This algorithm consists of aligning all possible pairs of sequences, finding a sequence Star chosen from sequence that has maximum alignment score, and aligning all sequences referred to the sequence Star. Each of pairwise alignments is conducted using dynamic programming technique. The complexity of DNA multiple sequence alignment using dynamic programming technique is very high. The computation time is increased exponentially due to the increasing of the number and the length of DNA sequences. This research aims to accelerate computation of Star Multiple Sequence Alignment using Message Passing Interfaces (MPI). The performance of the proposed method was evaluated by calculating speedup. Experiment was conducted using 64 sequences of 800 bp Glycine-max-chromosome-9-BBI fragments yielded by randomly cut from reference sequence of Glycine-max-chromosome-9-BBI taken from NCBI (National Center for Biotechnology Information). The results showed that the proposed technique could obtain speed up three times using five computers when aligning 64 sequences of Glycine-max-chromosome-9-BBI fragments. Moreover, the increasing of the number of computers would significantly increase speed up of the proposed method.

Keywords: DNA multiple sequence alignment, distributed computing, star algorithm, message passing interface

1. Introduction
Multiple Sequence Alignment (MSA) is a technique for aligning multiple biological sequences [1] to find similarities and differences in the sequence [2]. MSA is used in phylogenetic analysis to assessing the origin of species. MSA is also used to search a single nucleotide polymorphism (SNP) for molecular-based plant breeding genetics [3]. The growth of biological sequence databases makes the sequence alignment computation increase significantly [1].

The complexity of sequence alignment is \(O(\text{LN})\), where \(L\) is the length of each sequence and \(N\) is the number of sequences [4]. The longer and the more sequences to be aligned the longer it’s computational time. The optimal time of sequence alignment is increased exponentially with increasing of the number and length of sequences [5].

Methods or tools are needed to obtain optimal computational process. Previous study to optimize the computational sequence alignment used a dynamic programming algorithm with complexity of \(O(mn)\) by Zhou and Chen [6]. Other studies used a linear space algorithm for pairwise sequence alignment with the complexity of \(O(n)\) [7, 8]. Others used parallel computing with MPI [9] and GPU-CUDA [1, 3]. Parallel computing used to propose new classification algorithm [12] and simulate properties of polymer chain [13] in other case of computational problem.

In this research, parallel computing is utilized on clusters of computers known as Distributed Memory. This research tries to improve a previous work conducted by Sunarto which used star algorithms in parallel environment using MPI [9]. The previous research had limitation in scalability. In this research, we proposed a scalable distributed computing of DNA
Multiple Sequence Alignment. The new approach could handle more number of sequences and was also easily extended by increasing the number of resources used to compute multiple sequence alignment.

2. Research Method
2.1. Research Data
In this research, we used the sequence of Glycine-max-chromosome-9-BBI obtained from NCBI (National Center for Biotechnology Information) [10]. Detailed research data is shown in Table 1.

<table>
<thead>
<tr>
<th>Gene Name</th>
<th>Length (bp)</th>
<th>Number</th>
</tr>
</thead>
<tbody>
<tr>
<td>Glycine-max-chromosome-9-BBI</td>
<td>811-850</td>
<td>64</td>
</tr>
</tbody>
</table>

2.2. Multiple Sequence Alignment (MSA) with Star Algorithm
Algorithms for MSA are actually developed based on probabilistic or heuristic approaches such as the Star method and the Progressive method [3]. This research used the Star method by improving and extending the previous research conducted by Sunarto AA et al. In the Star alignment algorithm, there are 3 stages, including (i) calculating pairwise alignment scores from all possible sequence pairs, (ii) selecting a Star sequence which has the best alignment score, and (iii) aligning the Star sequence with other sequences [3, 9]. The illustration of multiple sequence alignment using the Star algorithm can be described as follows:

For example, supposed we have DNA sequence data as follows:
Seq 1 = ATGG
Seq 2 = ATGC
Seq 3 = ATCC
Seq 4 = AGCG

The first stage is calculating pairwise similarity scores from all possible sequence pairs. This stage has $O(n^2)$ complexity where $n$ is number of sequences. This stage begins by calculating the number of sequence pairs using Equation (1):

$$(n-1) + (n-2) + \ldots + 1 \text{ or } n(n-1)/2$$

Where $n$ is number of sequences.

The next step is creating identity matrix containing the sequences, symbolized by $K_n$. Figure 1 shows the identity matrix.

<table>
<thead>
<tr>
<th>Seq 1</th>
<th>Seq 2</th>
<th>Seq 3</th>
<th>Seq 4</th>
</tr>
</thead>
<tbody>
<tr>
<td>Seq 1</td>
<td>K1</td>
<td>K2</td>
<td>K3</td>
</tr>
<tr>
<td>Seq 2</td>
<td>K1</td>
<td>K4</td>
<td>K5</td>
</tr>
<tr>
<td>Seq 3</td>
<td>K2</td>
<td>K4</td>
<td>K6</td>
</tr>
<tr>
<td>Seq 4</td>
<td>K3</td>
<td>K5</td>
<td>K6</td>
</tr>
</tbody>
</table>

Figure 1. Pairwise sequence identity matrix

The results of the determination of DNA pairwise sequence are:
K1 = Seq 1 (ATGG) and Seq 2 (ATGC)
K2 = Seq 1 (ATGG) and Seq 3 (ATCC)
K3 = Seq 1 (ATGG) and Seq 4 (AGCG)
K4 = Seq 2 (ATGC) and Seq 3 (ATCC)
K5 = Seq 2 (ATGC) and Seq 4 (AGCG)
K6 = Seq 3 (ATCC) and Seq 4 (AGCG)
Each pair of sequences above is aligned using Needleman Wunsch algorithm. This algorithm uses dynamic programming approach to find global similarity of sequences. Needleman Wunsch algorithm consists of matrix initialization, filling the value of each cell matrix and tracing back. Similarity score of pairwise sequence is calculated using Equation (1) below:

$$A_{mn} = \max \begin{cases} 
A_{m-1,n-1} + S_{m,n} \\
A_{m,n-1} + W \\
A_{m-1,n} + W \\
0 
\end{cases}$$

Where:
- $m$ is the row and $n$ is the column.
- $A$ is a matrix of values for each cell is expressed as $(A_{m,n})$
- $S$ is the score of each cell is expressed as $(S_{m,n})$, match score = 5 and unmatch score = -3
- $W$ expressed as gap alignment with value = -4

The calculation results of similarity scores for all pairs of sequences are shown in Figure 2.

![Figure 2. Sequence alignment similarity scores](image)

The second stage of star algorithm is selecting a Star sequence. A Star sequence was selected by comparing the sequence alignment similarity scores of all sequences. A sequence with the best similarity score was chosen as a Star sequence. Next, the Star sequence was updated by aligning it to the other sequences. The complexity of this stage is $O(n)$, where $n$ is the number of sequences. Illustration of this stage is shown in Figure 3.

![Figure 3. Selecting a Star sequence](image)
The third stage of star algorithm is realignment. This stage is realigning the Star sequence to others sequences by using dynamic programming technique. Complexity is $O(n)$, where $n$ is the amount of sequences. Illustration for realignment is shown in Figure 4.

Seq 2 (A T G C - )
Seq 1 (A T G G )
Seq 3 (A T C C )
Seq 4 (A G C G )

\[
\begin{align*}
\text{Seq 2} & \quad \text{Seq 1} & \quad \text{Seq 3} & \quad \text{Seq 4} \\
A & - & T & G \\
A & T & G & G \\
A & T & C & C \\
A & G & C & G \\
\end{align*}
\]

The results of the sequence alignment

2.3 Parallelization MSA with Message Passing Interfaces (MPI)

In this research, we used Foster’s Methodology for implementing parallel programming. This methodology consists of partitioning, communication, agglomeration and mapping stage [8]. Parallelization was conducted for computing pairwise similarity scores. The calculation began by dividing the sequence pairs into multiple processes (threads) symbolized by $K1, K2 \ldots Kn$. In the case when the amount of computers is equal to the amount of processes, each process will be allocated to each computer. If $Pk$ is a computer for $k=1,2,\ldots,n$ this allocation can be defined as $P1=K1, P2=K2 \ldots Pn=Kn$. Unfortunately the number of sequence pairs is often larger than the number of computers. Sequence pairs which has not been allocated yet must wait until the previous process has been completed. Allocation of processes (threads) when the number of sequence pairs are more than the number of computers or when $Kn > Pn$ can be described in Figure 5.

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Illustration of data distribution of DNA sequence with MPI can be seen in Figure 6. In this research, communication of MPI used point to point communication with a blocking send and receives operations. Parallelization scheme for MSA is to assign a computer as the data divider and other computers as data processors. A data divider called rank 0 distributed sequence pairs using MPI_Send() to other computer as data processors (Figure 7). Data processors which were symbolized by the rank 1...n received sequence pairs using MPI_Recv() (Figure 8). Next, each data processors conducted pairwise sequence alignment to compute pairwise similarity scores in parallel. The calculation results of each data processor were transmitted to data divider (rank 0) by using MPI_Send(). Data divider with rank 0 received the similarity scores using MPI_Recv() command and completing the MSA process by selecting a Star sequence and realigning all sequence to the Star sequence.
Figure 6. Illustration the distribution of sequence data with MPI

MPI_Init(&argc, &argv);
MPI_Comm_rank (MPI_COMM_WORLD, &rank);//Computer initialization
MPI_Comm_size (MPI_COMM_WORLD, &p_size);//Initialize the number of computers
If rank == 0 {
    c = ((js*js)-js)/2; // Calculate the many pairwise sequences
    p_bts = p_size-1; // Limit Number of Computers
    p=1; // Limit Number of computers for processing data (other than rank 0)
    for (i=1; i<=js-1; i++){
        for (j=i+1; j<=js; j++){
            us1 = i-1; // Initialization the order of sending data sequence
            us2 = j-1; // Initialization the order of sending data sequence
            MPI_Send (&c, 1, MPI_INT, p, 1, MPI_COMM_WORLD);
            MPI_Send (&SIZE, 1, MPI_INT, p, 3, MPI_COMM_WORLD);
            MPI_Send (&p_bts, 1, MPI_INT, p, 5, MPI_COMM_WORLD);
            MPI_Send (&us1, 1, MPI_INT, p, 6, MPI_COMM_WORLD);
            MPI_Send (&us2, 1, MPI_INT, p, 7, MPI_COMM_WORLD);
            MPI_Send (s[us1], SIZE+1, MPI_CHAR, p, 8, MPI_COMM_WORLD);
            MPI_Send (s[us2], SIZE+1, MPI_CHAR, p, 9, MPI_COMM_WORLD);
            p++;
            if (p>p_bts){p=1;} // Looping sequence data sending besides to rank 0
        }
    }
} // Tutup rank 0

Figure 7. Source code data divider

If rank <> 0 {
    MPI_Recv(&c, 1, MPI_INT, 0, 1, MPI_COMM_WORLD, &status);
    MPI_Recv(&SIZE, 1, MPI_INT, 0, 3, MPI_COMM_WORLD, &status);
    MPI_Recv(&p_bts, 1, MPI_INT, 0, 5, MPI_COMM_WORLD, &status);
    p=1;
    for (h=0; h<=c-1; h++){
        if (rank==p){
            MPI_Recv(&us1, 1, MPI_INT, 0, 6, MPI_COMM_WORLD, &status);
            MPI_Recv(&us2, 1, MPI_INT, 0, 7, MPI_COMM_WORLD, &status);
            MPI_Recv(s[us1], SIZE+1, MPI_CHAR, 0, 8, MPI_COMM_WORLD, &status);
            MPI_Recv(s[us2], SIZE+1, MPI_CHAR, 0, 9, MPI_COMM_WORLD, &status);
        }
    }
} //end rank <> 0

Figure 8. Source code data processors
3. Results and Analysis

This research used 5 computers with specification Intel Core CPU i3-3220@3.30GHz with 4 CPU cores, 8 GB of RAM memory. A whole genome data with FASTA format of *Glycine-max-chromosome-9-BBI* was divided randomly into some numbers of sequences such as 3, 4, 8, 12, 14, 16, 20, 24, 28, 32, 36, 40, 44, 48, 52, 56, 60 and 64 sequence. Testing evaluation was conducted in 3, 4 and 5 computers.

3.1. Results

The evaluation results was described in Figure 9-11. Figure 9 showed the performance of our proposed method in term of execution time.

![Figure 9. The execution time in various number of sequences](image)

The result of sequence alignment computation showed that the parallel computation time was faster than the sequential computation time. Figure 9 showed the difference in computation time of data FASTA Glycine-max-chromosome-9-BBI with the length of 811-850 basepair. Using 3, 4 and 8 sequences the computational time was only slightly different. However, with the increasing of the number of sequences the computational time was significantly different.

3.2. Analysis

The performance analysis or evaluation of this proposed method was conducted by calculating speedup [11]. Speedup can be obtained by using Equation (2).

\[
\text{Speedup} = \frac{T_s}{T_p} \tag{2}
\]

Description:
- \(T_s\) = Sequential execution time
- \(T_p\) = Parallel execution time

The results of the parallel computing with 3, 4 and 5 PC (Personal Computer) showed that increasing of the number of computers (PCs) affected the parallel execution time. Parallel computational speedup increased as the number of computers was increased. Plot of parallel computing speedup for 3, 4 and 5 PCs were shown in Figure 10.
Figure 10. The speedup of parallel computing in various number of sequences

The other metric for showing the parallel performance is efficiency. Increasing the number of processors decreased the value of efficiency, and conversely an increase in the size of data will increase the efficiency [14]. The constant efficiency value indicates that the performance of a parallel system is scalable to the size of the data. Scalable parallel systems means the system is able to maintain performance with increasing number of processors [14] to process the data in a certain size. Efficiency is calculated using Equation (4). The results of efficiency calculation in this study can be seen in Figure 11.

\[
\text{Efficiency} = \frac{S(n)}{p}
\]

(4)

Limitation efficiency value is \(1 / p < \text{efficiency} < 1\)

Description:
\(S(n) = \text{Speedup of parallel computing}\)
\(p = \text{Many processors}\)

Figure 11. Efficiency of parallel computing in various number of sequences

Figure 11 showed that the use of 3, 4 and 5 processor were scalable in processing data with the length of 811-850 basepair and the number of sequence of 32-64. This efficiency will increase when the size and the length of sequence increased with increasing of the number of processor.
4. Conclusion

Parallel computing using MPI is recommended for MSA. This study show that the speed up of using MPI in distributed environment for conducting MSA was increased by increasing the size and the number of DNA sequences aligned. The trend of efficiency was almost constant when using 56 of sequences or more. This tendency indicated that this method was scalable to the size of data.

Future research can be conducted by using the larger size of data sequence and increase the number of processors to improve the speedup and efficiency of parallel computing for MSA. In this research, pairwise sequence alignment was conducted in sequential computation. In future, we could conduct hybrid computing by conducting the pairwise sequence alignment in parallel using MPI and CUDA GPU to increase speedup significantly.

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References