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# **Bachelor's Thesis**

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## **My Title**

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I hereby declare that I have written this thesis independently without any help from others and without the use of documents or aids other than those stated. I have mentioned all used sources and cited them correctly according to established academic citation rules.

Göttingen, 09. May 2020



## Abstract

*Here comes the abstract...*





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## **Chapter 1**

# **Introduction**



## **Chapter 2**

### **Basics**

#### **2.1 Multiple sequence alignment**



## **Chapter 3**

### **Prior Work**





## **Chapter 4**

# **Algorithm**

In this chapter, the analysis of ...



## **Chapter 5**

# **Implementation**

In this chapter, the implementation of ...



## Chapter 6

# Evaluation

### 6.1 BALiBASE 3

The third version of the BALiBASE benchmark protein alignment database has been released in 2005 and is widely employed for the comparison of multiple alignment programs [1, 2]. It is constructed in a semi automatic process as shown in fig. 6.1 and suitable to evaluate global and local alignment programs. The database is split into 5 reference sets with different characteristics representing distinctive multiple alignment problems. It is divided into:

- reference set 1 subset V1, for which any two sequences share <20% identity and no internal insertions over 35 residues long
- reference set 1 subset V2, consisting of families with at least four equidistant sequences for which any two sequences share 20-40% identity and no large insertions
- reference set 2, for which all sequences share >40% identity and at least one 3D structure is known. Additionally an "Orphan" sequence with <20% identity is chosen per family
- for reference set 3, all sequences in the same sub-family have >40% identity, whereas sequences from

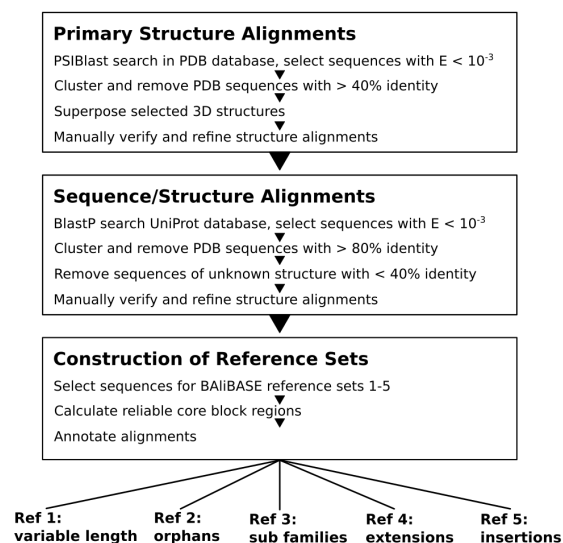


Figure 6.1: Flow chart showing the semi automatic process used to establish the reference sets TODO cite self

different subfamilies share <20% identity

- for reference sets 4 and 5, every sequence shares at least 20% with one other sequence, including sequences with large N/C-terminal extensions (ref 4) or internal insertions (ref 5)

### 6.1.1 Core blocks

Evaluating and comparing alignment programs is a difficult problem due to the uncertainty of supposedly "real" alignments of actual sequences. The BALiBASE database marks alignment columns which can be reliably aligned as so called "core blocks". These core blocks are calculated and manually verified, making up 19% of the full length sequences which are used in the evaluation of *spam-align* [1].

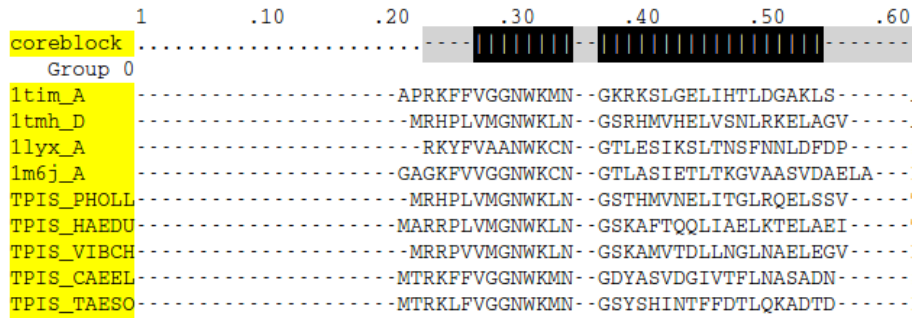


Figure 6.2: BALiBASE web interface. Black columns indicate core blocks. lbgi.fr

## 6.2 Sum-of-pairs and column score

Comparing the alignment output of different methods can be done by computing the sum-of-pairs and column scores.

Given a test alignment  $A_t$  and a reference alignment  $A_r$  with  $M$  sequences and  $N_t, N_r$  columns respectively, the sum-of-pairs and column score is defined according to Thompson et al. [3].

**Definition 6.2.1 (Sum-of-pairs score)**

The sum of pairs score is the ratio of correctly aligned individual residues. Formally it is defined as:

$$p_{ijk} = \begin{cases} 1 & \text{if residues } A_{t_{ij}} \text{ and } A_{t_{ik}} \text{ are aligned in } A_r \\ 0 & \text{otherwise} \end{cases}$$

$$S_i = \sum_{j=1}^M \sum_{k=i+1}^M p_{ijk}$$

$$SPS = \frac{\sum_{i=1}^{N_t} S_i}{\sum_{i=1}^{N_r} S_{r_i}}$$

with  $S_{r_i}$  being the number of correctly aligned residues in the reference.

**Definition 6.2.2 (Column score)**

The column score is the ratio of correctly aligned columns.

$$C_i = \begin{cases} 1 & \text{if all the residues in the } i\text{-th column are aligned correctly} \\ 0 & \text{otherwise} \end{cases}$$

$$CS = \frac{\sum_{i=1}^{N_t} C_i}{N_r}$$

Note that  $C_i = 1$  only if all the residues in the  $i$ -th column are aligned correctly and no residue belonging to this column is part of another one. For this reason, the numerator is smaller or equal to the denominator.

The definition of the column score is slightly different than that provided by the authors of BALiBASE [3] but resembles the actual implementation in the included BaliScore tool and its reimplementaion provided with this thesis.

These scores are only calculated for the core blocks of the BALiBASE alignments, meaning that for the following evaluation,  $A_t$  is an alignment over the full sequences, while  $A_r$  contains only the aligned residues inside the core blocks.

## 6.3 MAFFT

## 6.4 Results





## **Chapter 7**

# **Conclusion**

### **7.1 Further work**



# Bibliography

- [1] J. D. Thompson, P. Koehl, R. Ripp, and O. Poch, "Balibase 3.0: latest developments of the multiple sequence alignment benchmark," *Proteins: Structure, Function, and Bioinformatics*, vol. 61, no. 1, pp. 127–136, 2005.
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