

Part V

11 Construction of scoring matrix

11.1 Scoring schemes for protein sequence alignment

Applying an appropriate scoring scheme is critical to create biologically accurate alignments and phylogenetic trees.

Different types of scoring schemes for proteins

- Use of identity
- Use of the genetic code
- Use of a classification of amino acids
- Scoring matrix

Use of identity

The score is calculated by counting identical amino acids. It is equivalent with a simple scoring scheme with match: 1, mismatch: 0, and gap penalty: 0.

Example of “use of identity”

Calculate the SP score by counting identical amino acids.

Seq1 F-NV

Seq2 FPN-

Seq3 FC-V

$$S(\bar{s}^1, \bar{s}^2) = 2$$

$$S(\bar{s}^1, \bar{s}^3) = 2$$

$$S(\bar{s}^2, \bar{s}^3) = 1$$

$$S(\mathcal{A}) = S(\bar{s}^1, \bar{s}^2) + S(\bar{s}^1, \bar{s}^3) + S(\bar{s}^2, \bar{s}^3) = 2 + 2 + 1 = 5$$

Score: 5

Use of the genetic code

The score is based on the distance between two amino acids at the codon level.

Example of “use of the genetic code”

Seq1 FFFF

Seq2 FCNG

Phe (UUU, UUC) & Phe (UUU, UUC): 3
Phe (UUU, UUC) & Cys (UGU, UGC): 2
Phe (UUU, UUC) & Asn (AAU, AAC): 1
Phe (UUU, UUC) & Glu (GAA, GAG): 0

Score: 6

Use of a classification of amino acids

The score is based on the physio-chemical properties. For example, AACH (amino acid class hierarchy) can be used as a scoring scheme.

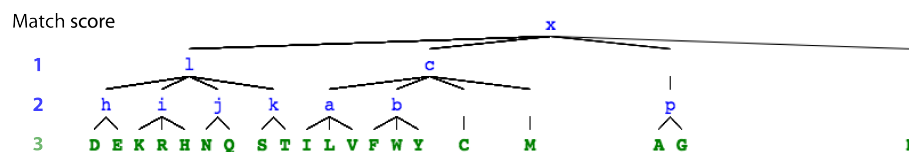


Figure 11.1: Example of amino acid class hierarchy (AACH)

Example of “Use of a classification of amino acids”

Calculate the score by using AACH.

Seq1 DDDP

Seq2 DEKD

D & D: 3, D & E: 2, D & K: 1, P & D: 0

Score: 6

Scoring matrix

- DNA/RNA: 4×4
- Protein: 20×20

PAM and BLOSUM

BLAST parameters

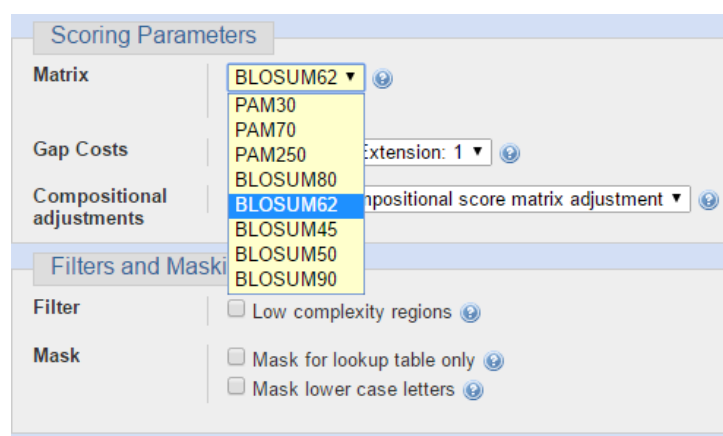


Figure 11.2: BLAST score parameters (source:)

Correspondence between PAM and BLOSUM

PAM 120	PAM 160	PAM 250
BLOSUM 80	BLOSUM 62	BLOSUM 45

Types of substitutions

There are several types of substitutions between two sequences from the common ancestor.

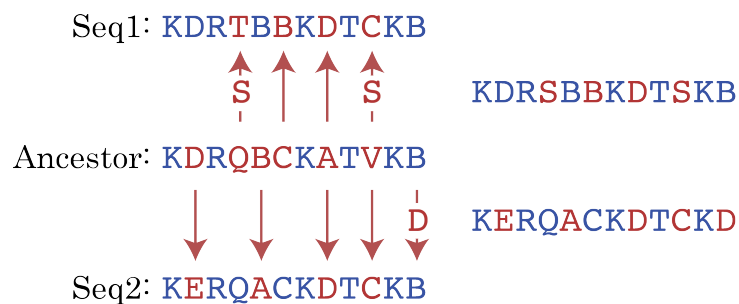


Figure 11.3: Different types of substitutions

Exercise 11.1

Calculate the score of the alignment by using different scoring schemes.

Seq1 K-RI
Seq2 KDCC

- Use the identity.
- Use the genetic code.

K	Lys	AAA, AAG
D	Asp	GAU, GAC
R	Arg	CGU, CGC, CGA
I	Ile	AUU, AUC, AUA
C	Cys	UGU, UGC

- Use AACH.

11.2 PAM accepted mutations

PAM is a popular scoring scheme for protein sequence alignments. It is based on substitution matrices created from experiment data.

Accepted point mutations

- Independent of positions and neighbor residues
- Independent from previous mutations at the same position
- Biological clock is assumed (the rate of mutations is constant)

PAM (point accepted mutation)

One PAM means one accepted point mutation per 100 residues.

Resources of constructing a PAM score

- 34 super-families
- 71 groups of homologous sequences (85% identity)

Preparations for constructing a PAM score

Counting the number of mutations is the first step to make a PAM score. Several sub-steps are involved.

- Create a phylogenetic tree
- Estimate ancestor sequences
- Count all occurrences of mutations

Frequencies of estimated mutations

Frequencies of estimated mutations are counted in internal nodes of the reconstructed tree.

f_{ab} : The number of mutations from a to b or from b to a

f_a : The total number of mutations in which a takes part

f : Twice the total number of mutations

Example of frequency calculation

Calculate f_{CA} , f_C , and f from the phylogenetic tree and the table below.

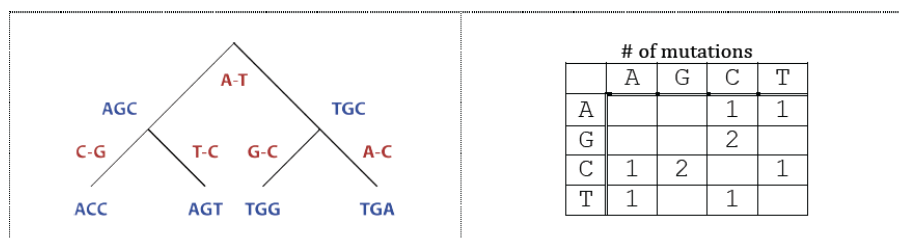


Figure 11.4: Phylogenetic tree and a table of the number of mutations

$$f_{CA} = 1$$

$$f_C = 1 + 2 + 1 = 4$$

$$f = 10$$

Background frequencies

The background probabilities are calculated from the data source.

p_a : The relative occurrence of a in the observed sequences

Example of background frequencies

Calculate p_G from the sequences below.

Seq1 ACC

Seq2 AGT

Seq3 TGG

Seq4 TGA

$$p_G = \frac{4}{12} \approx 0.333$$