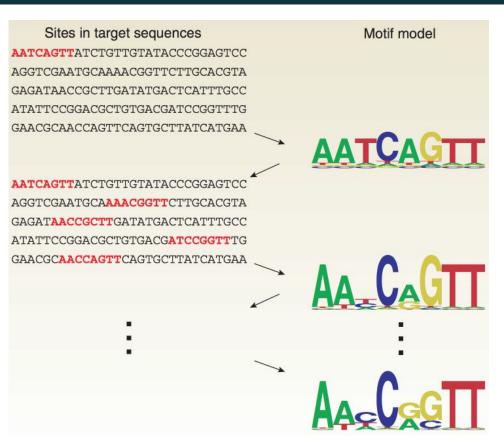
Week 06: Regulatory genomics

- DNA-binding sites/motifs
 - ChIP-seq
 - Position-weight matrices
 - Motif-finding
 - Expectation-Maximization
 - Gibbs Sampling

- 1. Define the probabilistic model and the likelihood function $P(X \mid \theta)$.
- 2. Identify the hidden variables (Z).
 - a. Here, they are the locations of the motifs in each sequence.
- Write the E step.
 - Compute the expected values of the hidden variables given current parameter values.
- 4. Write the **M step**.
 - Determine new parameters given the expected values of the hidden variables.
- 5. Repeat until convergence.

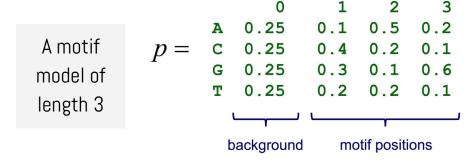


The **likelihood** of a model is the probability that the observed data could have been generated by the model under consideration.

- Easier to optimize the logarithm of this probability (hence '**log likelihood**') with respect to the parameters of the model:

$$\log L(\text{model} \mid \text{data}) = \log \Pr(\text{data} \mid \text{model})$$
$$= \sum_{i} \log \Pr(\text{data}_{i} \mid \text{model})$$

- MEME: Multiple EM for Motif Elicitation
- A motif is:
 - assumed to have a fixed width, W
 - represented by a matrix of probabilities: p_{c,k} (probability of character c in column k).
- The "background" (i.e. sequence outside the motif) is given by p_{c,0} (probability of base c in the background).
- Data is a collection of sequences, denoted X.
- Motif starting positions are represented by a matrix indicator variables (0/1) Z_i

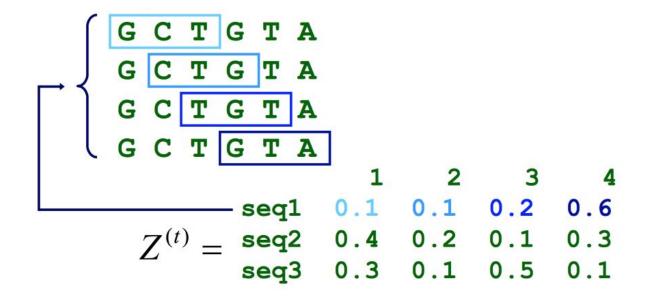


Given sequences L = 6. Possible starting positions m = L - W + 1

- 1. Define the probabilistic model and the likelihood function $P(X \mid \theta)$.
- 2. Identify the hidden variables (Z).
 - a. Here, they are the locations of the motifs in each sequence.
- 3. Write the **E step**.
 - a. Compute the expected values of the hidden variables given current parameter values.
- 4. Write the **M step**.
 - a. Determine new parameters given the expected values of the hidden variables.
- 5. Repeat until convergence.

```
given: width parameter \mathbf{W}, set of sequences
  t=0
  set initial values for p^{(0)}
  do
     ++t
     re-estimate Z^{(t)} from p^{(t-1)} (E-step)
     re-estimate p<sup>(t)</sup> from Z<sup>(t)</sup> (M-step)
  until change in p^{(t)} < \varepsilon
return: p<sup>(t)</sup>, Z<sup>(t)</sup>
```

- **E-step**: compute the expected values of Z given X and $p^{(t-1)}$
- Expected values: $Z^{(t)} = E[Z \mid X, p^{(t-1)}]$
- For example:



```
given: length parameter \mathbf{W}, set of sequences t=0 set initial values for p^{(0)} do  
++t 
re-estimate Z^{(t)} from p^{(t-1)} (E-step) 
re-estimate p^{(t)} from Z^{(t)} (M-step) 
until change in p^{(t)} < \mathbf{\epsilon} 
return: p^{(t)}, Z^{(t)}
```

$$P(Z_{i,j} = 1 | X_i, p^{(t-1)})$$

- **E-step**: compute the expected values of Z given X and $p^{(t-1)}$
- Expected values: $Z^{(t)} = E[Z \mid X, p^{(t-1)}]$
- Applying Bayes rule to: $P(Z_{i,j} = 1 \mid X_i, p^{(t-1)})$

$$Z_{i,j}^{(t)} = \frac{P(X_i \mid Z_{i,j} = 1, p^{(t-1)})P(Z_{i,j} = 1)}{\sum_{k=1}^{m} P(X_i \mid Z_{i,k} = 1, p^{(t-1)})P(Z_{i,k} = 1)}$$

$$Z_{i,j}^{(t)} \propto P(X_i | Z_{i,j} = 1, p^{(t-1)})$$

given: length parameter
$$\mathbf{W}$$
, set of sequences $t=0$ set initial values for $p^{(0)}$ do $++t$ re-estimate $Z^{(t)}$ from $p^{(t-1)}$ (E-step) re-estimate $p^{(t)}$ from $Z^{(t)}$ (M-step) until change in $p^{(t)} < \mathbf{\epsilon}$ return: $p^{(t)}$, $Z^{(t)}$

Assuming that it is equally likely that the motif will start in any position

$$P(Z_{i,j} = 1) = \frac{1}{m}$$

Probability of a Sequence Given a Motif Starting Position

$$P(X_i \mid Z_{i,j} = 1, p) = \prod_{k=1}^{j-1} p_{c_k,0} \prod_{k=j}^{j+W-1} p_{c_k,k-j+1} \prod_{k=j+W}^{L} p_{c_k,0}$$

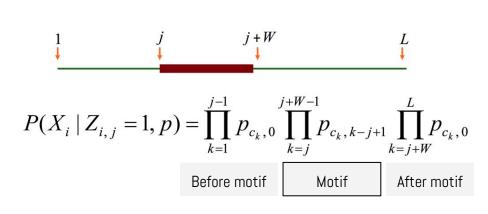
Before motif

Motif

After motif

- X; is the i th sequence
- Z_{i,i} is 1 if motif starts at position j in sequence i
- $\mathbf{c}_{\mathbf{k}}$ is the base at position k in sequence i

Probability of a Sequence Given a Motif Starting Position



0.25 0.2 0.2

 $X_i = G C T G T A G$

- X, is the ith sequence
- \mathbf{Z}_{ii} is 1 if motif starts at position j in sequence i
- $\mathbf{c}_{\mathbf{k}}$ is the base at position k in sequence i

$$P(X_i | Z_{i,3} = 1, p) =$$

$$p_{G,0} \times p_{C,0} \times p_{T,1} \times p_{G,2} \times p_{T,3} \times p_{A,0} \times p_{G,0} =$$

$$0.25 \times 0.25 \times 0.2 \times 0.1 \times 0.1 \times 0.25 \times 0.25$$

 $P(X_i | Z_{i,1} = 1, p^{(t-1)})$?

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0.1

- **E-step**: compute the expected values of Z given X and $p^{(t-1)}$
- Expected values: $Z^{(t)} = E[Z \mid X, p^{(t-1)}]$

$$X_i = G C T G T A G$$

```
0.25 0.2 0.2
```

++† re-estimate $Z^{(t)}$ from $p^{(t-1)}$ (E-step) 0.1 re-estimate $p^{(t)}$ from $Z^{(t)}$ (M-step) until change in $p^{(t)} < \varepsilon$ return: p^(t), Z^(t)

$$Z_{i,j}^{(t)} \propto P(X_i \mid Z_{i,j} = 1, p^{(t-1)})$$

$$Z^{(t)}_{i,1} \propto P(X_i | Z_{i,1} = 1, p^{(t-1)}) = 0.3 \times 0.2 \times 0.1 \times 0.25 \times 0.25 \times 0.25 \times 0.25$$
$$Z^{(t)}_{i,2} \propto P(X_i | Z_{i,2} = 1, p^{(t-1)}) = 0.25 \times 0.4 \times 0.2 \times 0.6 \times 0.25 \times 0.25 \times 0.25$$

Normalize so that
$$\sum_{j=1}^m Z^{(t)}{}_{i,j} = 1$$

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given: length parameter W, set of sequences

set initial values for $p^{(0)}$

t=0

do

- **E-step**: compute the expected values of Z given X and $p^{(t-1)}$
- Expected values: $Z^{(t)} = E[Z \mid X, p^{(t-1)}]$

$$p = \begin{bmatrix} 0 & 1 & 2 & 3 \\ A & 0.25 & 0.1 & 0.5 & 0.2 \\ C & 0.25 & 0.4 & 0.2 & 0.1 \\ G & 0.25 & 0.3 & 0.1 & 0.6 \\ T & 0.25 & 0.2 & 0.2 & 0.1 \end{bmatrix}$$

ACAGCA

$$Z^{(t)}_{1,1} = 0.1, \ Z^{(t)}_{1,2} = 0.7, \ Z^{(t)}_{1,3} = 0.1, \ Z^{(t)}_{1,4} = 0.1$$

AGGCAG

$$Z^{(t)}_{2,1} = 0.4, \ Z^{(t)}_{2,2} = 0.1, \ Z^{(t)}_{2,3} = 0.1, \ Z^{(t)}_{2,4} = 0.4$$

TCAGTC

```
Z^{(t)}_{3,1} = 0.2, \ Z^{(t)}_{3,2} = 0.6, \ Z^{(t)}_{3,3} = 0.1, \ Z^{(t)}_{3,4} = 0.1
```

```
given: length parameter \mathbf{W}, set of sequences t=0
set initial values for p^{(0)}
do  
++t
re-estimate Z^{(t)} from p^{(t-1)} (E-step)
re-estimate p^{(t)} from Z^{(t)} (M-step)
until change in p^{(t)} < \varepsilon
return: p^{(t)}, Z^{(t)}
```

- **M-step**: Estimate $p^{(t)}$ given X and $Z^{(t)}$.
- $p_{c,k}$ represents the prob. of base c in position k.
- k=0 represents the background.

$$p_{c,k}^{(t)} = \frac{n_{c,k} + d_{c,k}}{\sum\limits_{b \in \{A,C,G,T\}}} \\ n_{c,k} = \begin{cases} \sum\limits_{i} \sum\limits_{\{j \mid X_{i,j+k-1} = c\}} Z_{i,j}^{(t)} & k > 0 \\ \\ n_{c,k} = \begin{cases} \sum\limits_{i} \sum\limits_{\{j \mid X_{i,j+k-1} = c\}} Z_{i,j}^{(t)} & k > 0 \\ \\ n_{c} - \sum\limits_{j=1}^{W} n_{c,j} & k = 0 \end{cases}$$
 sum over positions where c appears

- M-step: Estimate $p^{(t)}$ given X and $Z^{(t)}$.
- $p_{c,k}$ represents the prob. of base c in position k.
- k=0 represents the background.

$$Z^{(t)}_{1,1} = 0.1, \ Z^{(t)}_{1,2} = 0.7, \ Z^{(t)}_{1,3} = 0.1, \ Z^{(t)}_{1,4} = 0.1$$
A G G C A G
 $Z^{(t)}_{2,1} = 0.4, \ Z^{(t)}_{2,2} = 0.1, \ Z^{(t)}_{2,3} = 0.1, \ Z^{(t)}_{2,4} = 0.4$
T C A G T C
 $Z^{(t)}_{3,1} = 0.2, \ Z^{(t)}_{3,2} = 0.6, \ Z^{(t)}_{3,3} = 0.1, \ Z^{(t)}_{3,4} = 0.1$

ACAGCA

$$p^{(t)}_{A,1} = \frac{Z^{(t)}_{1,1} + Z^{(t)}_{1,3} + Z^{(t)}_{2,1} + Z^{(t)}_{3,3} + 1}{Z^{(t)}_{1,1} + Z^{(t)}_{1,2} \dots + Z^{(t)}_{3,3} + Z^{(t)}_{3,4} + 4}$$

$$p^{(t)}_{\mathrm{C},2} =$$

- M-step: Estimate $p^{(t)}$ given X and $Z^{(t)}$.
- $p_{c,k}$ represents the prob. of base c in position k.
- k=0 represents the background.

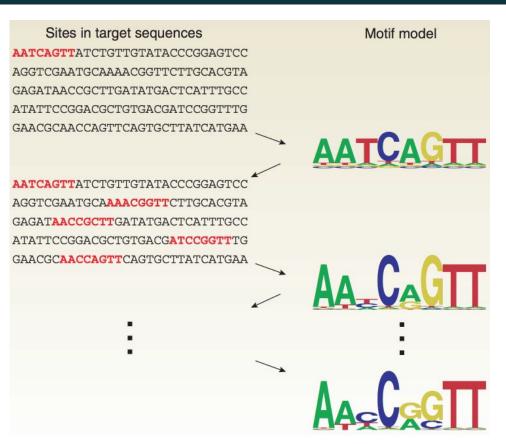
$$Z^{(t)}_{1,1} = 0.1, \ Z^{(t)}_{1,2} = 0.7, \ Z^{(t)}_{1,3} = 0.1, \ Z^{(t)}_{1,4} = 0.1$$
 A G G C A G
$$Z^{(t)}_{2,1} = 0.4, \ Z^{(t)}_{2,2} = 0.1, \ Z^{(t)}_{2,3} = 0.1, \ Z^{(t)}_{2,4} = 0.4$$
 T C A G T C
$$Z^{(t)}_{3,1} = 0.2, \ Z^{(t)}_{3,2} = 0.6, \ Z^{(t)}_{3,3} = 0.1, \ Z^{(t)}_{3,4} = 0.1$$

ACAGCA

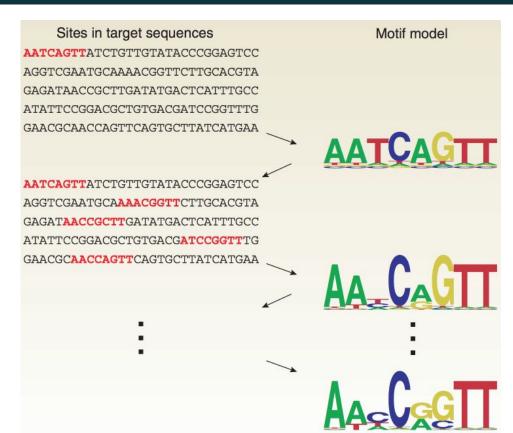
$$p^{(t)}_{A,1} = \frac{Z^{(t)}_{1,1} + Z^{(t)}_{1,3} + Z^{(t)}_{2,1} + Z^{(t)}_{3,3} + 1}{Z^{(t)}_{1,1} + Z^{(t)}_{1,2} \dots + Z^{(t)}_{3,3} + Z^{(t)}_{3,4} + 4}$$

$$p^{(t)}_{C,2} = \frac{Z^{(t)}_{1,1} + Z^{(t)}_{1,4} + Z^{(t)}_{2,3} + Z^{(t)}_{3,1} + 1}{Z^{(t)}_{1,1} + Z^{(t)}_{1,2} \dots + Z^{(t)}_{3,3} + Z^{(t)}_{3,4} + 4}$$

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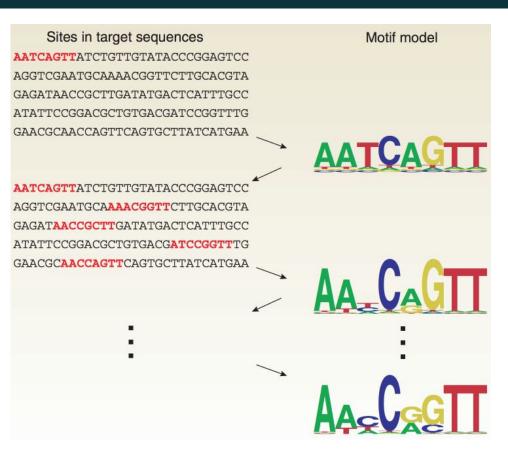


- 1. Assume zero or more motif occurrences per sequence.
- 2. Choosing the width of the motif.
- 3. Finding multiple motifs in a group of sequences.
- 4. Choosing good starting points for the parameters.
- 5. Using background knowledge to bias the parameters.



MEME:

- EM is susceptible to local maxima; so, try multiple starting points.
- Motif must be similar to some subsequence in data set
- For every distinct subsequence of length W in the training set
 - derive an initial p matrix from this subsequence
 - o run EM for 1 iteration
- Choose motif model (i.e. p matrix) with highest likelihood.
- Run EM to convergence.



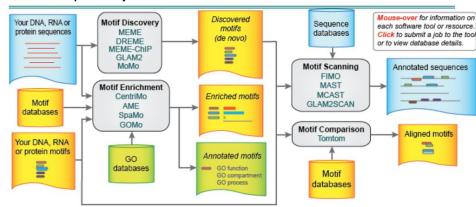
D'haeseleer (2016) Nat. Biotech. Gitter @ U. Wisconsin

MEME:

- Lawrence & Reilly (1990) "An expectation maximization (EM) algorithm for the identification and characterization of common sites in unaligned biopolymer sequences", Proteins.
- Bailey & Elkan (1994) "Fitting a mixture model by expectation maximization to discover motifs in biopolymers", Proceedings of the Second International Conference on Intelligent Systems for Molecular Biology.
- http://meme-suite.org/

The MEME Suite

Motif-based sequence analysis tools

























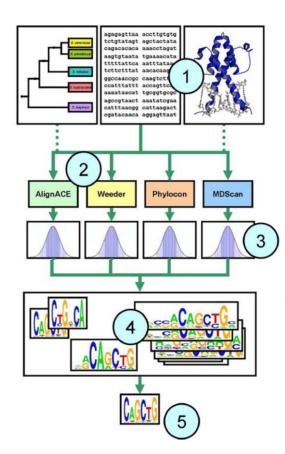








Practical strategies for finding motifs



- Assemble input data. Results may be improved by restricting the input to high-confidence sequences.

 Some algorithms achieve improved performance by using phylogenetic conservation information from orthologous sequences or information about protein DNA-binding domains.
- Choose several motif discovery programs for the analysis. For recommended programs see Figure 3.
- Test the statistical significance of the resulting motifs. Use control calculations to estimate the empirical distribution of scores produced by each program on random data.
- Clustering and post-processing the motifs. Motif discovery analyses often produce many similar motifs, which may be combined using clustering. Phylogenetic conservation information may be used to filter out statistically significant, but non-conserved motifs that are more likely to correspond to spurious sequence patterns.
- Interpretation of motifs. Algorithms exist for linking motifs to transcription factors and for combining motif discovery with expression data.