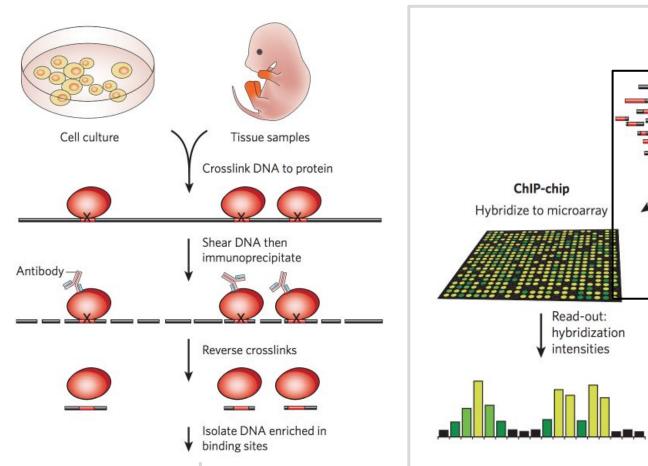
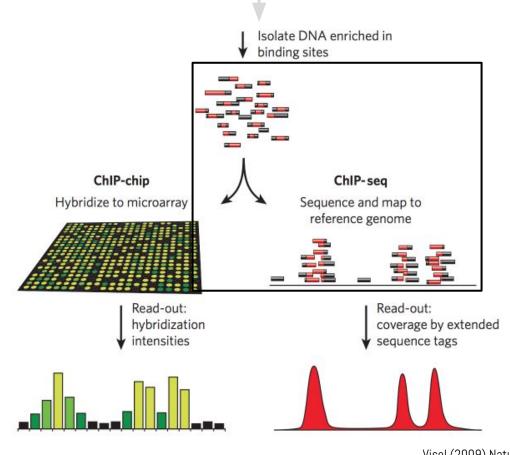
Regulatory genomics

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

Mapping of regulatory elements using ChIP-chip and ChIP-seq





Mapping of regulatory elements using ChIP-chip and ChIP-seq

Sequences are not aligned, we don't know motif positions.

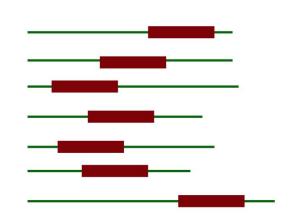
We also don't know what the motif looks like.

The motif model learning task:

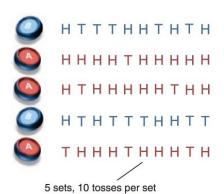
- Given: a set of sequences that are thought to contain occurrences of an unknown motif of interest
- Do:
 - o infer a model (PWM) of the motif, and
 - predict the locations of the motif occurrences in the given sequences.

Expectation-Maximization: Iteratively refine positions / motif profile

Gibbs sampling: Iteratively sample positions / motif profile



a Maximum likelihood



Coin A	Coin B
	5 H, 5 T
9 H, 1 T	
8 H, 2 T	
	4 H, 6 T
7 H, 3 T	
24 H, 6 T	9 H, 11 T

$$\hat{\theta}_A = \frac{24}{24+6} = 0.80$$

$$\hat{\theta}_B = \frac{9}{9+11} = 0.45$$

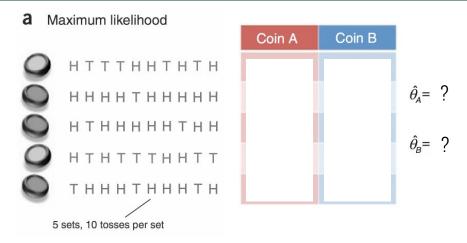
 $x = (x_1, x_2, ..., x_5) | x_i \in \{0,1,...,10\}$ is the no. of heads observed during the ith set of tosses.

 $z = (z_1, z_2, ..., z_5) \mid z_i \in \{A,B\}$ is the identity of the coin used during the ith set of tosses.

A coin-flipping experiment

- $\theta_A & \theta_B$ are the biases of two coins A & B.
- **Goal**: Estimate $\theta = (\theta_A, \theta_B)$ by repeating the following procedure five times:
 - Randomly choose one of the two coins (with equal probability)
 - Perform ten independent coin tosses with the selected coin.

Maximum likelihood estimation: statistical model that has the highest probability of generating the observed data $-\theta$ that maximizes $logP(x,z;\theta)$.

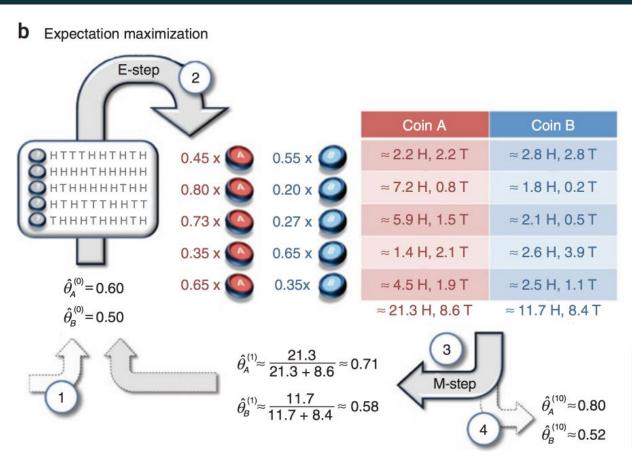


 $\mathbf{x} = (\mathbf{x_1}, \mathbf{x_2}, ..., \mathbf{x_5}) \mid \mathbf{x_i} \in \{0,1,...,10\}$ is the no. of heads observed during the \mathbf{i}^{th} set of tosses.

 $z = (z_1, z_2, ..., z_5) | z_i \in \{A,B\}$ is the identity of the coin used during the ith set of tosses. [Hidden variables / Latent factors]

A coin-flipping experiment

- $\theta_A \& \theta_B$ are the biases of two coins A & B.
- **Goal**: Estimate $\theta = (\theta_A, \theta_B)$ by repeating the following procedure five times:
 - Randomly choose one of the two coins (with equal probability; but you don't know which coin was chosen)
 - Perform ten independent coin tosses with the selected coin.



E-step:

- Estimate $P(x_i, z_i | \theta^{(t)})$ and the expected values of the hidden variables.

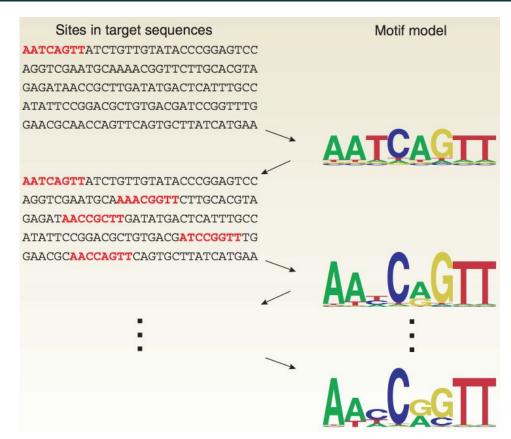
M-step:

Estimate new parameters θ^(t+1)
 given current estimates of hidden
 variables & parameters.

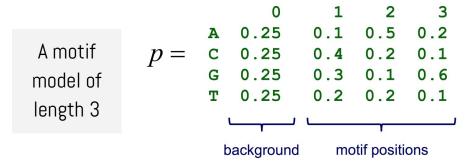
Repeat until convergence.

 $P(x_i, z_i | \theta^{(t)})$: Likelihood function, from here on also going to be written as $P(X, Z | \theta)$.

- 1. Define the probabilistic model and the likelihood function $P(X \mid \theta)$.
 - a. Here, it is the **position weight matrix**.
- 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.



- MEME: Multiple EM for Motif Elicitation
- A motif is:
 - \circ assumed to have a fixed width, ${f W}$
 - \circ 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) \mathbf{Z}_{ii}

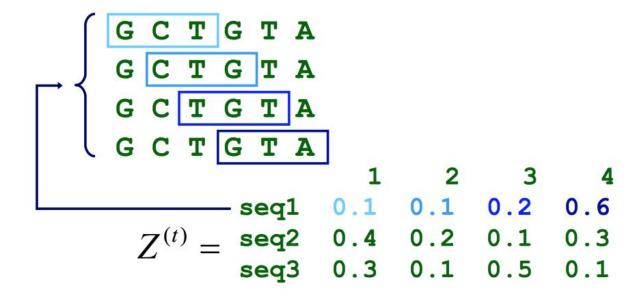


Given sequences L = 6. Possible starting positions m = L - W + 1

- 1. Define the probabilistic model and the likelihood function $P(X|\theta)$.
 - a. Here, it is the position weight matrix.
- 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 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)}
```

- **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: width parameter 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)}
```

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

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

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

given: width parameter W, set of sequences t=0set initial values for $p^{(0)}$ do
++tre-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: $p^{(t)}$, $Z^{(t)}$

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

the motif will start in any position

Assuming that it is equally likely that

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

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_i is the i^{th} sequence
- \mathbf{Z}_{ij} is 1 if motif starts at position j in sequence i
- \mathbf{c}_k is the base at position k in sequence i

Probability of a Sequence Given a Motif Starting Position

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

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

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

0.25 0.2 0.2

- X_i is the i^{th} sequence
- \mathbf{Z}_{ii} is 1 if motif starts at position j in sequence i
- \mathbf{c}_k is the base at position k in sequence i

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

= 0.25 \times 0.25 \times 0.25 \times 0.1 \times 0.1 \times 0.25 \times 0.25

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0.1

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

$$X_i = \mathbf{G} \ \mathbf{C} \ \mathbf{T} \ \mathbf{G} \ \mathbf{T} \ \mathbf{A} \ \mathbf{G}$$

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

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

given: width parameter 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: $p^{(t)}$, $Z^{(t)}$

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

...

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

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- **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: width parameter 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_{ck} represents the prob. of base c in position k.
- k=0 represents the background.

$$p_{c,k}^{(t)} = \frac{n_{c,k} + d_{c,k}}{\sum\limits_{b \in \{A,C,G,T\}}} \qquad n_{c,k} = \begin{cases} \sum\limits_{i} \sum\limits_{\{j \mid X_{i,j+k-1} = c\}} Z_{i,j}^{(t)} & k > 0 \\ \\ n_{c} - \sum\limits_{j=1}^{W} n_{c,j} & k = 0 \end{cases}$$
 total # c's in the dataset

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

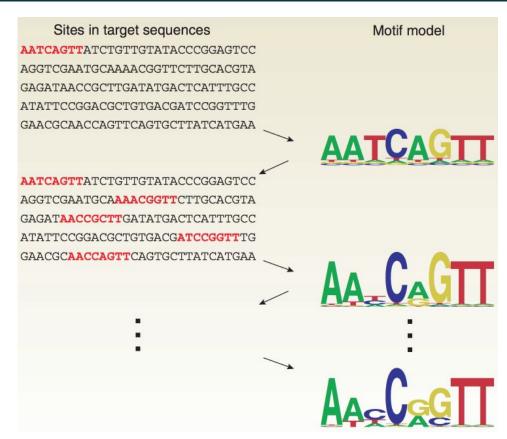
ACAGCA

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

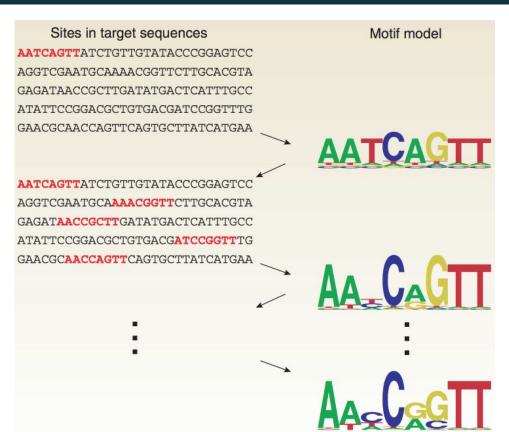
$$p^{(t)}_{C,2} = \frac{Z^{(t)}_{1,1} + Z^{(t)}_{1,4} + Z^{(t)}_{2,3} + Z^{(t)}_{3,1} + 1}{Z^{(t)}_{1,1} + Z^{(t)}_{1,2} \dots + Z^{(t)}_{3,3} + Z^{(t)}_{3,4} + 4}$$

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

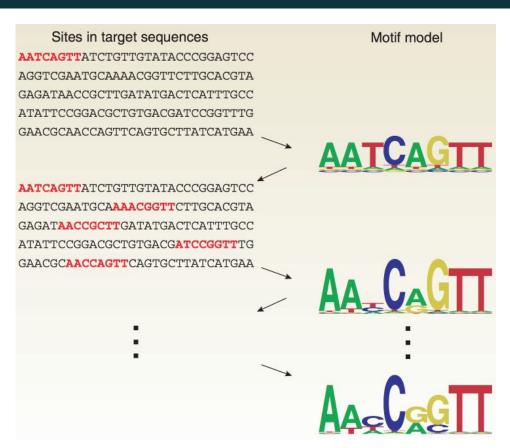


- 1. Assume zero or more motif occurrences per sequence.
- 2. Choosing the width of the motif.
- 3. Finding multiple motifs in a group of sequences.
- 4. Choosing good starting points for the parameters.
- 5. Using background knowledge to bias the parameters.



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.



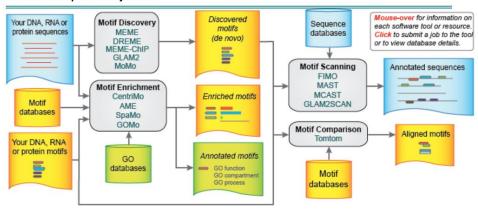
D'haeseleer (2016) Nat. Biotech. Gitter @ U. Wisconsin

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

































Motif finding using Gibbs sampling instead of EM

EM can get trapped in local minima

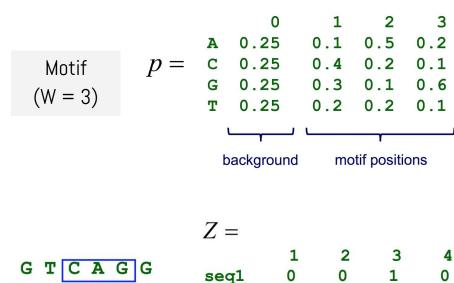
• One approach to alleviate this limitation: try different (perhaps random) initial parameters.

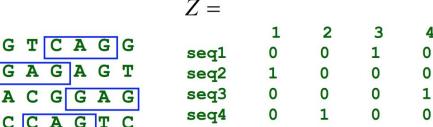
Gibbs sampling exploits randomized search to a much greater degree:

- Can be viewed as a stochastic analog of EM for this task.
- In theory, Gibbs sampling is less susceptible to local minima than EM.

Motif finding using Gibbs sampling

- A motif is:
 - \circ assumed to have a fixed width, ${f W}$
 - 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_{ii}.





Motif finding using Gibbs sampling

- 1. Choose initial **Z** containing the motif starting position in each sequence at random.
- 2. Loop through each sequence X_i :
 - Update p (position frequency matrix of background + motif) based on all sequences except X_i.
 - b. Based on the *updated* \mathbf{p} , calculate the location of best match in sequence $\mathbf{X}_{\mathbf{i}}$ and update the corresponding row in \mathbf{Z} .
- 3. Repeat until convergence.

$$p_{c,k}^{(t)} = \frac{n_{c,k} + d_{c,k}}{\sum_{b \in \{A,C,G,T\}} (n_{b,k} + d_{b,k})}$$

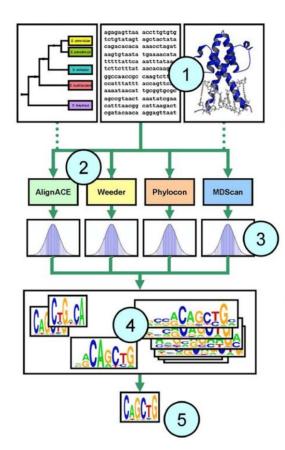
$$Z_{i,j}^{(t)} \propto P(X_i \mid Z_{i,j} = 1, p^{(t-1)})$$

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

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

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

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.