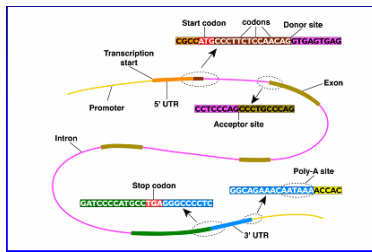


Computational gene finding

Gene finding in eukaryotic DNA



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Ab initio methods

- Use information embedded in the genomic sequence *exclusively* to predict the gene structure.
- Find structure G representing gene boundaries + internal gene structure which maximizes the probability $P(G|\text{genomic sequence})$.
- Hidden Markov models are the predominant generative method for modeling the problem.

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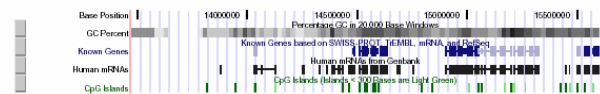
Ab-initio methods

- Advantages**
 - Intuitive, natural modeling
 - Prediction of 'novel' genes, *i.e.*, with no a priori known cDNA or protein evidence
- Caveats**
 - Not effective in detecting alternatively spliced forms, interleaved or overlapping genes
 - Difficulties with gene boundary identification
 - Potentially large number of false positives with over-fitting

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A simple example: CpG Islands



CpG nucleotides in the genome are frequently methylated. (Write CpG not to confuse with CG base pair)

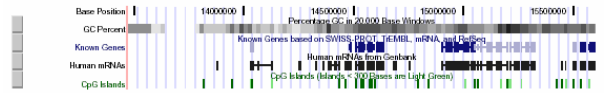
$C \rightarrow \text{methyl-C} \rightarrow T$

Methylation often suppressed around genes, promoters \rightarrow CpG islands

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Example: CpG Islands



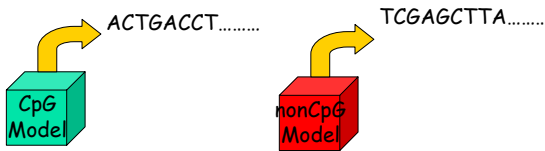
In CpG islands, CG is more frequent than in the rest of the genome

Two problems

- Given a short DNA sequence, does it come from a CpG island or not?
 - Is this part of a CpG island or not?
- How to find the CpG islands in a long sequence?



Generative models



Models generate sequences of strings in the A,T,C,G alphabet. Model parameters are tuned to reflect characteristics of CpG and non CpG islands.

Markov processes: a quick intro

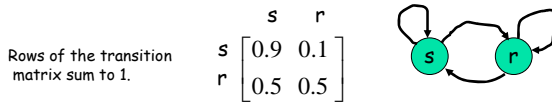
- We are interested in predicting weather, which can be either be sunny (s) or rainy (r).
- The weather on a given day depends only on the weather on the previous day.

$$P(w_t | w_{t-1}, \dots, w_1) = P(w_t | w_{t-1})$$

This is the Markov property.

Markov process example

- We have knowledge of the transition probabilities between sunny and rainy days.



- We know the initial probabilities of s and r.

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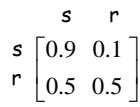
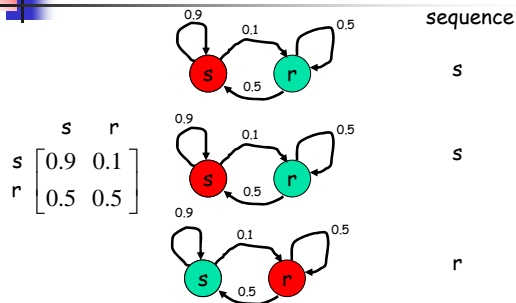
Generating weather sequences

- Let the probabilities of weather on the first day be $[0.5 \ 0.5]$. Let's say we start with a sunny day.
- Now we consult our transition matrix and find that $P(w|s) = [0.9 \ 0.1]$. It is more likely that the next day will be sunny too.
- We repeat this process, flipping coins biased by the probability $P(w_t|w_{t-1})$ to get a sequence representing weather for a consecutive set of days.

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Generating sequences (Take 2)



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Prediction

- Suppose day is rainy. We will represent this as a vector of probabilities over the two values.
- $$\pi(1) = [0 \ 1];$$
- How do we predict weather on day 2 given $\pi(1)$ and the transition probabilities P ?
 - From P , we can see that the probability of day 2 being sunny is .5, and for being rainy is 0.5

$$\pi(1) * P = [0.5 \ 0.5];$$

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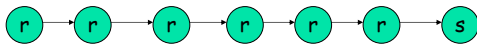
31

Probability of a sequence

- What is the probability of observing the sequence "rrrrrrs"?

$$P(X = rrrrrrs) = \pi(r)P(r|r)P(r|r)P(r|r)P(r|r)P(r|r)P(r|r)P(s|r)$$

$$= \pi(r) \prod_{t=2..7} P(x_t | x_{t-1}) = (0.5)^7$$



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Which weather pattern is more likely?

- Given a transition model

$$\begin{array}{c} s \quad r \\ s \begin{bmatrix} 0.9 & 0.1 \\ 0.5 & 0.5 \end{bmatrix} \\ r \end{array}$$

- And an initial state distribution: [0.5 0.5]
- And two sequences: rrrrrrs and ssssssr
Which is more likely, given the model?

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Comparing likelihoods

$$P(X = rrrrrrs | Model) = \pi(r)[P(r|r)]^5 P(s|r) = (0.5)^7$$

$$P(X = ssssssr | Model) = \pi(s)[P(s|s)]^5 P(r|s) = 0.5 * (0.9)^5 * 0.1$$

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Markov models (summary)

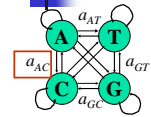
- States: $S = \{s_1, \dots, s_N\}$, N states
- Transition probability:
 - $a_{ij} = P(X_{t+1}=s_j | X_t=s_i)$, i, j in $[1..N]$
- Initial state probability
 - $\pi_i = P(X_1=s_i)$, i in $[1..N]$

Model generates sequences of states from S , and we can compute how likely a sequence is given the model.

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Markov models for CpG islands



A state for each of the four letters A, C, G, and T in the DNA alphabet

$$a_{st}^+ = \frac{c_{st}^+}{\sum_{t'} c_{st'}^+}$$

From a set of known CpG islands, and non CpG islands, estimate the transition probabilities

+	A	C	G	T
A	.180	.274	.426	.120
C	.171	.368	.274	.188
G	.161	.339	.375	.125
T	.079	.355	.384	.182

-	A	C	G	T
A	.300	.205	.285	.210
C	.322	.298	.078	.302
G	.248	.246	.298	.208
T	.177	.239	.292	.292

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Using the model

- To use the model for classification of a given sequence, calculate the log-odds ratio.
- Is the sequence more likely to come from a CpG island or a non-CpG region?

$$P(x | CpG) > P(x | nonCpG)$$

$$\frac{P(x | CpG)}{P(x | nonCpG)} > 1$$

$$\log \frac{P(x | CpG)}{P(x | nonCpG)} > 0$$

Log-odds ratio

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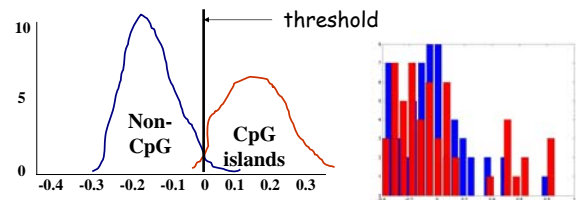
The log-odds ratio

$$S(x) = \log \frac{P(x | CpG)}{P(x | nonCpG)} = \sum_{i=1}^L \log \frac{a_{x_{i-1}x_i}^+}{a_{x_{i-1}x_i}^-}$$

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Histogram of log-odds scores



Given a short sequence x , does it come from CpG island (Yes-No question)?

Decision rule: if $S(x) > 0$ then CpG else non-CpG

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How to locate CpG islands?

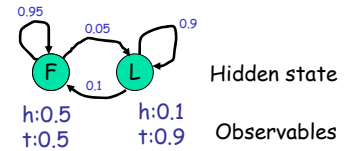
- Given a DNA sequence, find the CpG islands in it, if any.
- Approach: Calculate the log-odds score for a window of w nucleotides around every base in the sequence. Predict as CpG islands, those with a positive log-odds score.
- Problem: What should the size of the window w be? Predictions are sensitive to choice of w .

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The occasionally dishonest casino

- A casino uses a fair coin most of the time, but occasionally they switch to a loaded coin. You can't see which coin they are using, just the results of the flips (heads and tails) are visible.



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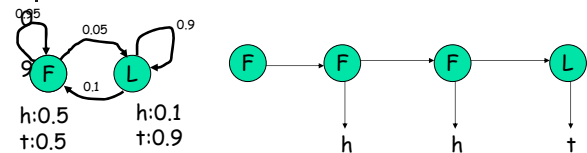
Generating coin flips

- Start in one of the states, F or L (i.e., pick a fair or loaded coin to start with) (initial probabilities).
- Move to the next state (F or L), based on the transition probabilities. Generate an h or t based on the emission probabilities of that state.
- Repeat above step.

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Generating flips (take 2)



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Hidden Markov Models

- $S = \{s_1, \dots, s_N\}$, N states
 - $O = \{o_1, \dots, o_M\}$, M observation symbols
 - $a_{ij} = P(S_{t+1}=s_j | S_t=s_i)$, i, j in $[1..N]$; **transition probabilities**
 - $b_i(k) = P(E_t=o_k | S_t=s_i)$, k in $[1..M]$, i in $[1..N]$; **emission probabilities**
 - $\pi_i = P(S_1=s_i)$, i in $[1..N]$; **initial state probabilities**
- $\lambda = (A, B, \pi)$ specifies the HMM model

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Dishonest casino as an HMM

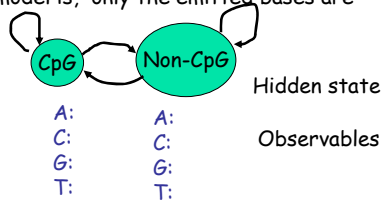
- $N = 2$, $S = \{F, L\}$
- $M = 2$, $O = \{h, t\}$
- $A = \begin{matrix} & F & L \\ F & \begin{bmatrix} 0.95 & 0.05 \end{bmatrix} \\ L & \begin{bmatrix} 0.10 & 0.90 \end{bmatrix} \end{matrix}$
- $B = \begin{matrix} & h & t \\ F & \begin{bmatrix} 0.5 & 0.5 \end{bmatrix} \\ L & \begin{bmatrix} 0.1 & 0.9 \end{bmatrix} \end{matrix}$
- $\pi = [1 \ 0]$

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A generative model for CpG islands

- There are two hidden states: CpG and non-CpG. Each state is characterized by emission probabilities of the 4 bases. You can't see which state the model is, only the emitted bases are visible.



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Filtering or the forward computation

- Given an HMM model (A, B, π) , and an observation sequence $o_1 \dots o_t$, can we find the most likely hidden state at time t , S_t ?
 - $P(S_t | o_1 \dots o_t)$: filtering

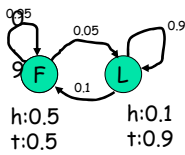
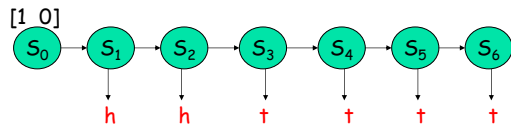
Observation sequence: h h t t t t

What is the hidden state here (F or L)?

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Filtering (contd.)

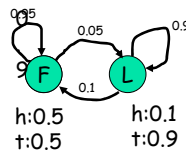
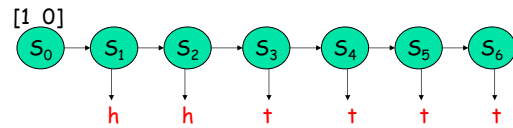


What is the distribution of S_1 ?
 Since, $s_0=F$, we can say that $P(S_1|S_0)=[0.95 \ 0.05]$, based on the transition probabilities alone.
 But is that all we know?

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More filtering



We have also observed h at time 1.
 How can we fold it in into the assessment of the distribution of S_1 ?

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Filtering (contd.)

$$P(S_1 | o_1) = \frac{P(o_1 | S_1)P(S_1)}{P(o_1)}$$

$$P(S_1 = F | o_1 = h) = \alpha P(h | F)0.95 = \alpha(0.5)(0.95)$$

$$P(S_1 = L | o_1 = h) = \alpha P(h | L)0.05 = \alpha(0.1)(0.05)$$

$$\alpha(0.5)(0.95) + \alpha(0.1)(0.05) = 1$$

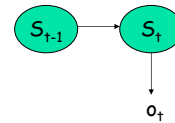
Therefore, $P(S_1)=[0.99 \ 0.01]$

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Filtering computation

F L
 [p 1-p]



Recursively computed

$$P(S_t | o_t, o_1 \dots o_{t-1}) = P(o_t | S_t) \sum_{s_{t-1}} P(S_t | s_{t-1}) P(s_{t-1} | o_1 \dots o_{t-1})$$

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Summary: filtering

Find $P(S_t | o_1, \dots, o_t) = cP(S_t, o_1, \dots, o_t)$.

Define $\alpha_t(i) = P(o_1, \dots, o_t, S_t = s_i)$.

Initialize: $\alpha_0(i) = \pi_i, 1 \leq i \leq n$

Recursion: $\alpha_{t+1}(j) = b_j(o_{t+1}) \sum_{i=1}^n \alpha_t(i) a_{ij}, 0 \leq j \leq n, 1 \leq t \leq T-1$

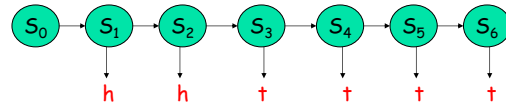
Termination: $\alpha_T(i), 1 \leq i \leq n$

Time complexity $O(n^2T)$

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Smoothing/posterior decoding



Question: can we re-estimate the distribution at S_k where $k < t$, using information about the observed sequence upto time t ?

That is, what is $P(S_k | o_1 \dots o_t)$?

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Backward computation

$$P(S_k | o_1, \dots, o_t) = c \overbrace{P(o_{k+1}, \dots, o_t | S_k)}^{\text{Backward computation}} \underbrace{P(S_k | o_1, \dots, o_k)}_{\text{Forward computation}}$$

Define $\beta_k(i) = P(o_{k+1}, \dots, o_t | S_k = s_i)$.

Initialize: $\beta_T(i) = 1, 1 \leq i \leq N$.

Recursion: $\beta_k(i) = c \sum_{j=1}^N a_{ij} b_j(o_{k+1}) \beta_{k+1}(j), 1 \leq i \leq N, T-1 \leq k \leq 1$

Time complexity: $O(n^2T)$

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Posterior decoding

$$P(S_k = i | o_1, \dots, o_t) = c \beta_k(i) \alpha_k(i)$$

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Full Decoding

- Given HMM model (A, B, π) , and an observation sequence $o_1 \dots o_T$, can we find the most likely hidden state sequence $s_1 \dots s_T$?
 - $\text{argmax}_{\{s_1 \dots s_T\}} P(s_1 \dots s_T | o_1 \dots o_T)$

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The Viterbi algorithm

$$\delta_t(i) = \max_{x_1, \dots, x_{t-1}} P(s_1, \dots, s_{t-1}, S_t = i, o_1, \dots, o_t)$$

$$\text{Initialize: } \delta_0(i) = \pi_i, 1 \leq i \leq n$$

$$\text{Recursion: } \delta_{t+1}(j) = \max_i \delta_t(i) a_{ij} b_j(o_{t+1}),$$

$$1 \leq t \leq T-1, 1 \leq j \leq n$$

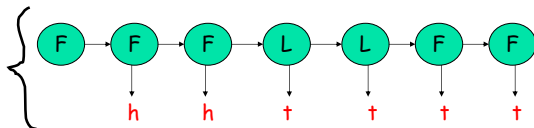
Computational complexity = $O(Tn^2)$

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Learning an HMM: case 1

- Given observation sequences, and the corresponding hidden state sequences, can we find the most likely model (A, B, π) which generated it?



Training data

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Parameter estimation

- Initial state distribution
 - Fraction of times state i is state 1 in training data
- Transition probabilities
 - a_{ij} = (number of transitions from i to j) / (number of transitions from i)
- Emission probabilities
 - $b_k(i)$ = (number of times k is emitted in state i) / (number of times state i occurs)

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Learning an HMM: case 2

- Given just the observation sequences, can we find the most likely model $\lambda = (A, B, \pi)$ which generated it?

$$\operatorname{argmax}_{\lambda} P(o_1 \dots o_t | \lambda)$$

Annotated training data is difficult to get; so we would like to derive model parameters from observable sequences.

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The EM algorithm

1. Guess a model λ
2. Use observation sequence to estimate transition probabilities, emission probabilities, and initial state probabilities.
3. Update model
4. Repeat 2 and 3 till no change in model

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Re-estimating parameters

- What is the probability of being in state i at time t and moving to state j , given the current model and the observation sequence O ?

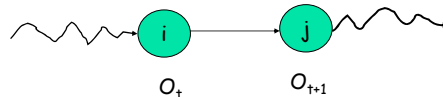
$$\xi_t(i, j) = P(S_t = i, S_{t+1} = j | O, \lambda)$$

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Using forward and backward computation

$$\xi_t(i, j) = \frac{\alpha_t(i) a_{ij} b_j(o_{t+1}) \beta_{t+1}(j)}{\sum_{i=1}^n \sum_{j=1}^n \alpha_t(i) a_{ij} b_j(o_{t+1}) \beta_{t+1}(j)}$$



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Re-estimating a_{ij}

- The transition probabilities a_{ij} can be re-estimated as follows

$$\hat{a}_{ij} = \frac{\sum_{t=1}^{T-1} \xi_t(i, j)}{\sum_{t=1}^{T-1} \sum_{j'=1}^n \xi_t(i, j')}$$

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Initial state probabilities

$$\gamma_t(i) = \sum_{j=1}^n \xi_t(i, j) \quad \text{Expected number of times in state } i$$

Initial state probabilities are simply $\gamma_1(i)$

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Emission probabilities

$\hat{b}_i(k) = \frac{\text{expected number of times in state } i \text{ and observe symbol } k}{\text{expected number of times in state } i}$

$$\hat{b}_i(k) = \frac{\sum_{t=1}^T \gamma_t(i)}{\sum_{t=1}^T \sum_{a_i=k} \gamma_t(i)}$$

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The EM algorithm

1. Guess a model $\lambda = (a, b, \pi)$
2. Use observation sequence to estimate

$$\xi_t(i, j) \text{ and } \gamma_t(i)$$

3. Use these estimates to recalculate

$$\lambda' = (a', b', \pi')$$

4. Repeat 2 and 3 till no change in model

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How to use the CpG island HMM

- Given a DNA region x , the **Viterbi** algorithm predicts locations of CpG islands on it.
- Given a nucleotide x_i , the **Viterbi** parse tells whether x_i is in a CpG island in the most likely sequence.
- **Posterior Decoding** can assign locally optimal predictions of CpG islands.
- A fully annotated training data set can be used to estimate the generating HMM.
- Even without annotations, we can use the EM procedure to derive model parameters.