CSE 527 Autumn 2006 Lectures 6-7 MLE, EM, Expression

Outline

- MLE: Maximum Likelihood Estimators
- EM: the Expectation Maximization Algorithm
- Bio: Gene expression and regulation

• Next: Motif description & discovery

MLE

Maximum Likelihood Estimators



Parameter Estimation

- Assuming sample $x_1, x_2, ..., x_n$ is from a parametric distribution $f(x|\theta)$, estimate θ .
- E.g.:

$$f(x) = \frac{1}{\sqrt{2\pi\sigma^2}} e^{-(x-\mu)^2/(2\sigma^2)}$$

$$\theta = (\mu, \sigma^2)$$

Maximum Likelihood Parameter Estimation

- One (of many) approaches to param. est.
- Likelihood of (indp) observations $x_1, x_2, ..., x_n$

$$L(x_1, x_2, \dots, x_n \mid \theta) = \prod_{i=1}^n f(x_i \mid \theta)$$

• As a function of θ , what θ maximizes the likelihood of the data actually observed

• Typical approach: $\frac{\partial}{\partial \theta} L(\vec{x} \mid \theta) = 0$ or $\frac{\partial}{\partial \theta} \log L(\vec{x} \mid \theta) = 0$

Example I

n coin flips, $x_1, x_2, ..., x_n$; n_0 tails, n_1 heads, $n_0 + n_1 = n$; $\theta = \text{probability of heads}$

$$L(x_1, x_2, \dots, x_n \mid \theta) = (1 - \theta)^{n_0} \theta^{n_1}$$

 $\log L(x_1, x_2, \dots, x_n \mid \theta) = n_0 \log(1 - \theta) + n_1 \log \theta$ $\frac{\partial}{\partial \theta} \log L(x_1, x_2, \dots, x_n \mid \theta) = \frac{-n_0}{1 - \theta} + \frac{n_1}{\theta}$

Setting to zero and solving:

$$\theta = \frac{n_1}{n}$$

(Also verify it's max, not min, & not better on boundary)

Ex. 2:
$$x_i \sim N(\mu, \sigma^2), \ \sigma^2 = 1, \ \mu$$
 unknown
 $L(x_1, x_2, \dots, x_n | \theta) = \prod_{1 \le i \le n} \frac{1}{\sqrt{2\pi}} e^{-(x_i - \theta)^2/2}$
 $\ln L(x_1, x_2, \dots, x_n | \theta) = \sum_{1 \le i \le n} -\frac{1}{2} \ln 2\pi - \frac{(x_i - \theta)^2}{2}$
 $\frac{d}{d\theta} \ln L(x_1, x_2, \dots, x_n | \theta) = \sum_{1 \le i \le n} (x_i - \theta)$
And verify it's max, $(\sum n) = 0$

And verify it's max, not min & not better on boundary

3

2

5

4

-3 -4 -5 -6 -7

$$= \left(\sum_{1 \le i \le n} x_i\right) - n\theta = 0$$

$$\hat{\theta} = \left(\sum_{1 \le i \le n} x_i\right) / n = \bar{x}$$

Ex 3:
$$x_i \sim N(\mu, \sigma^2), \ \mu, \sigma^2$$
 both unknown

$$\ln L(x_1, x_2, \dots, x_n | \theta_1, \theta_2) = \sum_{1 \leq i \leq n} -\frac{1}{2} \ln 2\pi \theta_2 - \frac{(x_i - \theta_1)^2}{2\theta_2}$$

$$\frac{\partial}{\partial \theta_1} \ln L(x_1, x_2, \dots, x_n | \theta_1, \theta_2) = \sum_{1 \leq i \leq n} \frac{(x_i - \theta_1)}{\theta_2} = 0$$

$$\hat{\theta_1} = \left(\sum_{1 \leq i \leq n} x_i\right) / n = \bar{x}$$

$$\ln L(x_1, x_2, \dots, x_n | \theta_1, \theta_2) = \sum_{1 \le i \le n} -\frac{1}{2} \ln 2\pi \theta_2 - \frac{(x_i - \theta_1)^2}{2\theta_2}$$
$$\frac{\partial}{\partial \theta_2} \ln L(x_1, x_2, \dots, x_n | \theta_1, \theta_2) = \sum_{1 \le i \le n} -\frac{1}{2} \frac{2\pi}{2\pi \theta_2} + \frac{(x_i - \theta_1)^2}{2\theta_2^2} = 0$$
$$\hat{\theta}_2 = \left(\sum_{1 \le i \le n} (x_i - \hat{\theta}_1)^2 \right) / n = \bar{s}^2$$

A consistent, but *biased* estimate of population variance. (An example of *overfitting*.) Unbiased estimate is:

$$\hat{\theta}_2 = \sum_{1 \le i \le n} \frac{(x_i - \hat{\theta}_1)^2}{n - 1}$$

Moral: MLE is a great idea, but not a magic bullet

EM

The Expectation-Maximization Algorithm













Likelihood

$$L(x_1, x_2, \dots, x_n | \overbrace{\mu_1, \mu_2, \sigma_1^2, \sigma_2^2, \tau_1, \tau_2}^{\theta})$$

$$= \prod_{i=1}^n \sum_{j=1}^2 \tau_j f(x_i | \mu_j, \sigma_j^2)$$

- Messy: no closed form solution known for finding θ maximizing L
- But what if we knew the $z_{ij} = \begin{cases} 1 & \text{if } x_i \text{ drawn from } f_j \\ 0 & \text{otherwise} \end{cases}$

EM as Egg vs Chicken

- IF z_{ij} known, could estimate parameters θ
- IF parameters θ known, could estimate z_{ii}
- But we know neither; (optimistically) iterate:
 - E: calculate expected z_{ii}, given parameters
 - M: calc "MLE" of parameters, given E(z_{ii})

Simple Idea: "Classification EM"

- If z_{ij} < .5, pretend it's 0; z_{ij} > .5, pretend it's 1
 i.e., *classify* points as component 0 or 1
- Now recale θ , assuming that partition
- then recalc \boldsymbol{z}_{ij} , assuming that $\boldsymbol{\theta}$
- then re-recale θ , assuming new z_{ii}
- etc., etc.

Full EM

 x_i 's are known; θ unknown. Goal is to find MLE θ of: $L(x_1,\ldots,x_n \mid \theta)$ (hidden data likelihood) Would be easy if z_{ij} 's were known, i.e., consider: $L(x_1, \ldots, x_n, z_{11}, z_{12}, \ldots, z_{n2} \mid \theta)$ (complete data likelihood) But z_{ij} 's aren't known. Instead, maximize *expected* likelihood of visible data $E(L(x_1,\ldots,x_n,z_{11},z_{12},\ldots,z_{n2} \mid \theta)),$ where expectation is over distribution of hidden data $(z_{ij}$'s)

The E-step

- Assume θ known & fixed
- A (B): the event that x_i was drawn from f_1 (f_2)

• A (B): the event that
$$\mathbf{x}_{i}$$
 was drawn from \mathbf{f}_{1} (\mathbf{f}_{2})
• D: the observed datum \mathbf{x}_{i}
• Expected value of \mathbf{z}_{i1} is P(A|D)
• E = 0 · P(0) + 1 · P(1)
• E = 0 · P(0) + 1 · P(1)
• P(A|D) = $\frac{P(D|A)P(A)}{P(D)}$
P(D) = $P(D|A)P(A) + P(D|B)P(B)$
= $f_{1}(x_{i}|\theta_{1})\tau_{1} + f_{2}(x_{i}|\theta_{2})\tau_{2}$

Complete Data Likelihood

Recall:

$$z_{1j} = \left\{ egin{array}{ccc} 1 & ext{if } x_1 ext{ drawn from } f_j \ 0 & ext{otherwise} \end{array}
ight.$$

so, correspondingly,

$$L(x_1, z_{1j} \mid \theta) = \begin{cases} \tau_1 f_1(x_1 \mid \theta) & \text{if } z_{11} = 1 \\ \tau_2 f_2(x_1 \mid \theta) & \text{otherwise} \end{cases}$$

Formulas with "if's" are messy; can we blend more smoothly? Yes, many possibilities. Idea 1:

$$L(x_1, z_{1j} \mid \theta) = z_{11} \cdot \tau_1 f_1(x_1 \mid \theta) + z_{12} \cdot \tau_2 f_2(x_1 \mid \theta)$$

Idea 2:

$$L(x_1, z_{1j} \mid \theta) = (\tau_1 f_1(x_1 \mid \theta))^{z_{11}} \cdot (\tau_2 f_2(x_1 \mid \theta))^{z_{12}}$$

M-step Details

(For simplicity, assume $\sigma_1 = \sigma_2 = \sigma; \tau_1 = \tau_2 = .5 = \tau$)

$$\begin{split} L(\vec{x}, \vec{z} \mid \theta) &= \prod_{1 \le i \le n} \frac{\tau}{\sqrt{2\pi\sigma^2}} \exp\left(-\sum_{1 \le j \le 2} z_{ij} \frac{(x_i - \mu_j)^2}{2\sigma^2}\right) \\ E[\log L(\vec{x}, \vec{z} \mid \theta)] &= E\left[\sum_{1 \le i \le n} \left(\log \tau - \frac{1}{2} \log 2\pi\sigma^2 - \sum_{1 \le j \le 2} z_{ij} \frac{(x_i - \mu_j)^2}{2\sigma^2}\right)\right] \\ &= \sum_{1 \le i \le n} \left(\log \tau - \frac{1}{2} \log 2\pi\sigma^2 - \sum_{1 \le j \le 2} E[z_{ij}] \frac{(x_i - \mu_j)^2}{2\sigma^2}\right) \end{split}$$

Find θ maximizing this as before, using $E[z_{ij}]$ found in E-step. Result:

 $|\mu_j = \sum_{i=1}^n E[z_{ij}]x_i / \sum_{i=1}^n E[z_{ij}]|$ (intuit: avg, weighted by subpop prob)

2 Component Mixture $\sigma_1 = \sigma_2 = 1; \ \tau = 0.5$

		mu1	-20.00		-6.00		-5.00		-4.99
		mu2	6.00		0.00		3.75		3.75
									l
x1	-6	z11		5.11E-12		1.00E+00		1.00E+00	
x2	-5	z21		2.61E-23		1.00E+00		1.00E+00	
x3	-4	x31		1.33E-34		9.98E-01		1.00E+00	Î
x4	0	z41		9.09E-80		1.52E-08		4.11E-03	l
x5	4	z51		6.19E-125		5.75E-19		2.64E-18	1
x6	5	z61		3.16E-136		1.43E-21		4.20E-22	
x7	6	z71		1.62E-147		3.53E-24		6.69E-26	Î

EM Summary

- Fundamentally a max likelihood parameter estimation problem
- Useful if analysis is more tractable when 0/1 hidden data z known
- Iterate:
 - E-step: estimate E(z) given θ
 - M-step: estimate θ maximizing E(likelihood) given E(z)

EM Issues

- Under mild assumptions (sect 11.6), EM is guaranteed to increase likelihood with every E-M iteration, hence will converge.
- But may converge to *local*, not global, max. (Recall the 4-bump surface...)
- Issue is probably intrinsic, since EM is often applied to NP-hard problems (including clustering, above, and motif-discovery, soon)
- Nevertheless, widely used, often effective

Gene Expression & Regulation

Gene Expression

- Recall a gene is a DNA sequence for a prot
- To say a gene is *expressed* means that it
 I. is *transcribed* from DNA to RNA
 - 2. the mRNA is processed in various ways
 - 3. is exported from the nucleus (eukaryotes)
 - 4. is translated into protein
- A key point: not all genes are expressed all the time, in all cells, or at equal levels

Transcription

- RNA polymerase complex
 - E. coli: 5 proteins (2α, β, β', σ)
 σ is *initiation factor*; finds promoter, then released/replaced by *elongation factors*
 - Eukaryotes: 3 pols, each >10 subunits
- attaches to DNA, melts helix, makes RNA copy (5' → 3') of template (3' → 5') at ~30nt/sec





5' Processing: Capping

- methylated G added to 5' end, and methyl added to ribose of 1st nucleotide of transcript
- probably helps distinguish protein-coding mRNAs from other RNA junk
 - prevents degradation
 - facilitates start of translation

3' Processing: Poly A (Eukaryotes)

- Transcript cleaved after AAUAAA (roughly)
- pol keeps running (until it falls off) but no 5' cap added to strand downstream of poly A site, so it's rapidly degraded
- I0s I00s of A's added to 3' end of transcript - its "poly A tail"

More processing: Splicing

- Also in eukaryotes, most genes are spliced: protein coding exons are interrupted by non-coding introns, which are cut out & degraded, exons spliced together
- More details about this when we get to gene finding



Nuclear Export

 In eukaryotes, mature mRNAs are actively transported out of the nucleus & ferried to specific destinations (e.g., mitochondria, ribosomes)

Regulation

- In most cells, pro- or eukaryote, easily a 10,000-fold difference between least- and most-highly expressed genes
- Regulation happens at all steps. E.g., some transcripts can be sequestered then released, or rapidly degraded, some are weakly translated, some are very actively translated, some are highly transcribed, some are not transcribed at all
- Below, focus on 1st step only: transcriptional regulation

DNA Binding Proteins

A variety of DNA binding proteins ("transcription factors"; a significant fraction, perhaps 5-10%, of all human proteins) modulate transcription of protein coding genes

Summary

- Learning from data:
 - MLE: Max Likelihood Estimators
 - EM: Expectation Maximization (MLE w/hidden data)
- Expression & regulation
 - Expression: creation of gene products
 - Regulation: when/where/how much of each gene product; complex and critical
- Next: using MLE/EM to find regulatory motifs in biological sequence data