Inferring Models of cis-Regulatory Modules using Information Theory

BMI/CS 776 www.biostat.wisc.edu/bmi776/ Spring 2018 Anthony Gitter gitter@biostat.wisc.edu

These slides, excluding third-party material, are licensed under <u>CC BY-NC 4.0</u> by Mark Craven, Colin Dewey, and Anthony Gitter

Overview

- Biological question
 - What is causing differential gene expression?
- Goal

- Find regulatory motifs in the DNA sequence

Solution

- FIRE (Finding Informative Regulatory Elements)

Goals for Lecture

Key concepts:

- Entropy
- Mutual information (MI)
- Motif logos
- Using MI to identify cis-regulatory module elements

A Common Type of Question



What causes this set of yeast genes to be up-regulated in stress conditions?

Experiments / Conditions

Figure from Gasch et al., Mol. Biol. Cell, 2000

cis-Regulatory Modules (CRMs)

 Co-expressed genes are often controlled by specific configurations of binding sites



- Problem
 - Create a code to communicate information
- Example
 - Need to communicate the manufacturer of each bike



- Four types of bikes
- Possible code

Туре	code	
Trek	11	
Specialized	10	
Cervelo	01	
Serotta	00	

 Expected number of bits we have to communicate: 2 bits/bike

- Can we do better?
- Yes, if the bike types aren't equiprobable

Type, probability	# bits	code
P(Trek) = 0.5	1	1
P(Specialized) = 0.25	2	01
P(Cervelo) = 0.125	3	001
P(Serotta) = 0.125	3	000

• Optimal code uses $-\log_2 P(c)$ bits for event with probability P(c)

Type, probability	# bits	code
P(Trek) = 0.5	1	1
P(Specialized) = 0.25	2	01
P(Cervelo) = 0.125	3	001
P(Serotta) = 0.125	3	000

Expected number of bits we have to communicate:
 1.75 bits/bike

$$-\sum_{c=1}^{|\mathcal{C}|} P(c) \log_2 P(c)$$

Entropy

- Entropy is a measure of uncertainty associated with a random variable
- Can be interpreted as the expected number of bits required to communicate the value of the variable

$$H(C) = -\sum_{c=1}^{|C|} P(c) \log_2 P(c)$$



 $\Pr(X=1)$

How is entropy related to DNA sequences?



- Typically represent a binding site
- Height of each <u>character</u> c is proportional to P(c)



Height of <u>logo</u> at a given position determined by decrease in entropy (from maximum possible)

$$H_{\max} - H(C) = \log_2 N - \left(-\sum_c P(c)\log_2 P(c)\right)$$

of characters in alphabet

13



Mutual Information

• *Mutual information* quantifies how much knowing the value of one variable tells about the value of another



Elemento et al., *Molecular Cell* 2007

- Finding Informative Regulatory Elements (FIRE)
- **Given** a set of sequences grouped into clusters
- **Find** motifs, and relationships, that have high *mutual information* with the clusters
- Applicable when sequences have continuous values instead of cluster labels





Continuous



Orientation bias



Mutual Information in FIRE

• We can compute the mutual information between a motif and the clusters as follows

$$I(M;C) = \sum_{m=0}^{1} \sum_{c=1}^{|C|} P(m,c) \log_2 \frac{P(m,c)}{P(m)P(c)}$$

m=0, 1 represent absence/presence of motif

c ranges over the cluster labels

Finding Motifs in FIRE

- Motifs are represented by regular expressions; initially each motif is represented by a strict *k*-mer (e.g. TCCGTAC)
- 1. Test all *k*-mers (*k*=7 by default) to see which have significant mutual information with the cluster label
- 2. Filter *k*-mers using a significance test to obtain motif seeds
- 3. Generalize each motif seed
- 4. Filter motifs using a significance test

Key Step in Generalizing a Motif in FIRE

- Randomly pick a position in the motif
- Generalize in all ways consistent with current value at position
- Score each by computing mutual information
- Retain the best generalization



Generalizing a Motif in FIRE

given: k-mer, n

 $best \leftarrow null$ repeat *n* times
motif \leftarrow *k*-mer
repeat
motif \leftarrow GeneralizePosition(motif) // shown on previous slide
until convergence (no improvement at any position)
if score(motif) > score(best)
best \leftarrow motif

return: best



Avoiding Redundant Motifs

- Different seeds could converge to similar motifs
 TCCGTAC
 TCCCTAC
 TCC[CG]TAC
- Use mutual information to test whether new motif is unique and contributes new information

$$\frac{I(M; C \mid M')}{I(M; M')} > r$$

M' previous motif

M new candidate motif

C expression clusters

Characterizing Predicted Motifs in FIRE

- Mutual information is also used to assess various properties of found motifs
 - orientation bias
 - position bias
 - interaction with another motif

Using MI to Determine Orientation Bias

I(S;C) C indicates cluster S=1 indicates motif present on transcribed strand S=0 otherwise (not present or not on transcribed strand)



Also compute MI where *S*=1 indicates motif present on complementary strand

Using MI to Determine Position Bias

I(P;O) *P* ranges over position bins O=0, 1 indicates whether or not the motif is over-represented in a sequence's cluster



Only sequences containing the motif are considered for this calculation

Using MI to Determine Motif Interactions

 $I(M_1; M_2)$ $M_1=0, 1$ indicates whether or not a sequence has the motif **and** is in a cluster for which the motif is over-represented; similarly for M_2



Using MI to Determine Motif Interactions



Yeast motif-motif interactions

White: positive association Dark red: negative association Blue box: DNA-DNA Green box: DNA-RNA Plus: spatial co-localization

Discussion of FIRE

- FIRE
 - mutual information used to identify motifs and relationships among them
 - motif search is based on generalizing informative kmers
- Consider advantages and disadvantages of k-mers versus PWMs
- In contrast to many motif-finding approaches, FIRE takes advantage of *negative* sequences
- FIRE returns all informative motifs found

Mutual Information for Gene Networks

- Mutual information and conditional mutual information can also be useful for reconstructing biological networks
- Build gene-gene network where edges indicate high MI in genes' expression levels
- Algorithm for the Reconstruction of Accurate Cellular Networks (ARACNE)

ARACNE

- Gaussian kernel estimator to estimate mutual information
 - No binning or histograms
- Data processing inequality
 - Prune indirect edges



Margolin et al. BMC Bioinformatics 2006