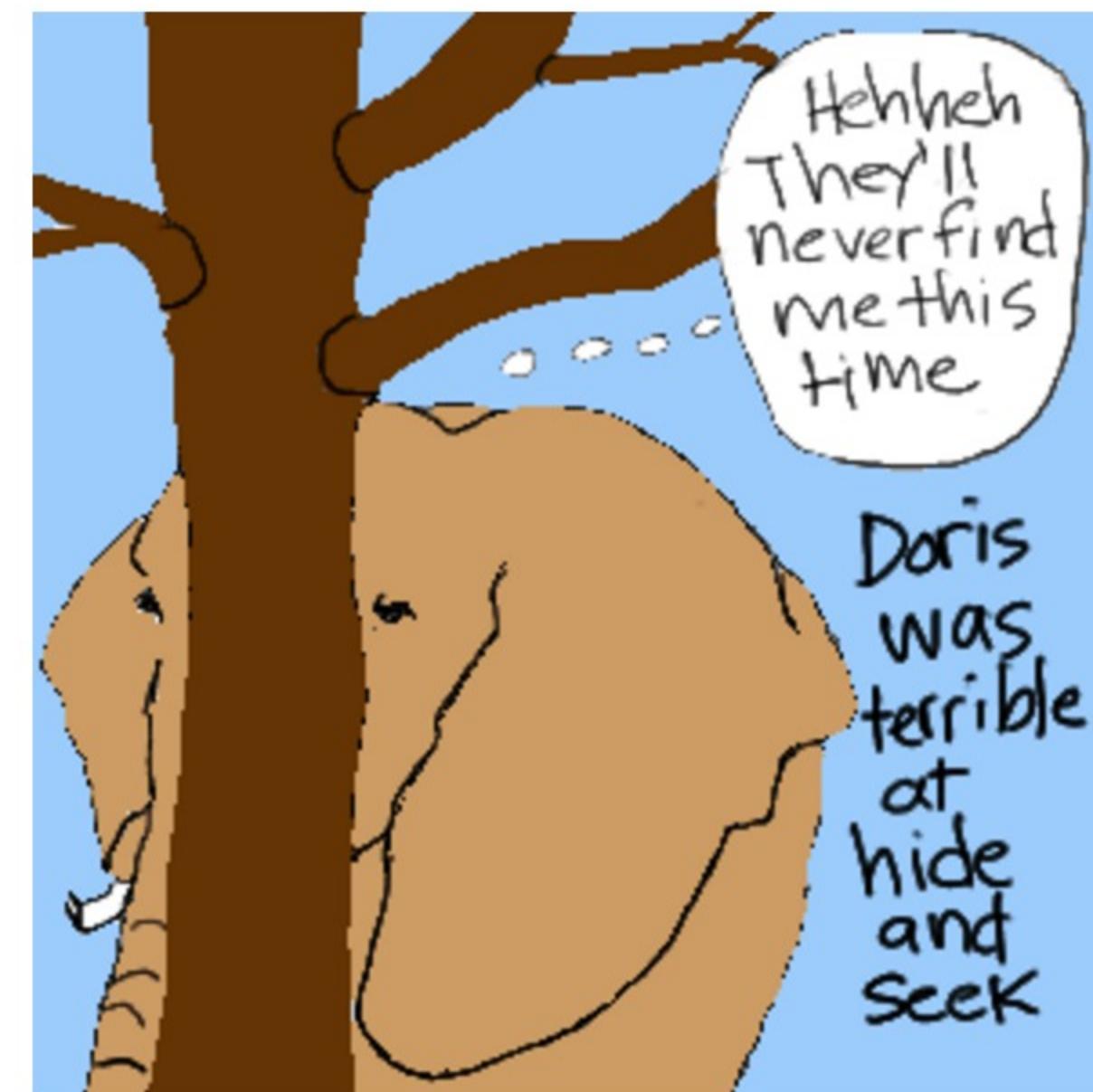


Finding Hidden Patterns in DNA

- What makes searching for frequent subsequences hard?
 - Allowing for errors?
 - All the places they could be hiding?



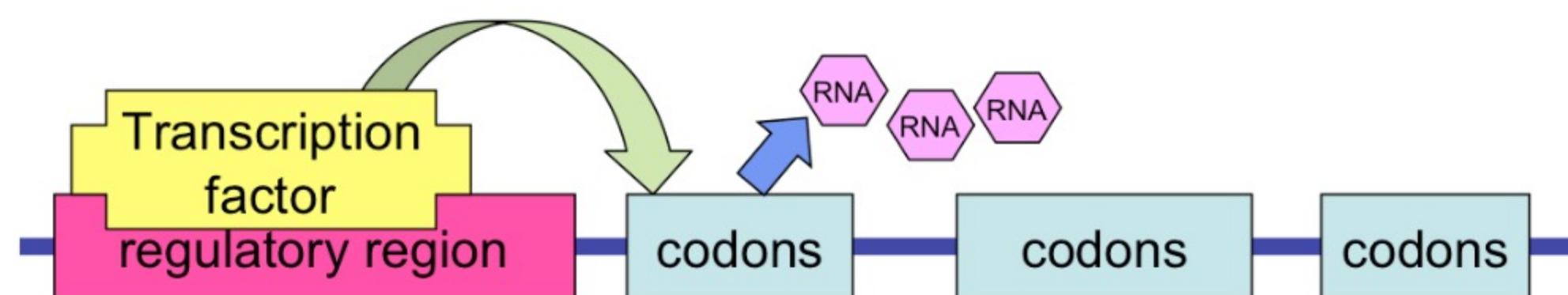
Initiating Transcription

- As a precursor to transcription (the reading of DNA to construct RNAs, that eventually leading to protein synthesis) special proteins bind to the DNA, and separate it to enable its reading.
- How do these proteins know where the coding genes are in order to bind?
- Genes are relatively rare
 - $O(1,000,000,000)$ bases/genome
 - $O(10000)$ genes/genome
 - $O(1000)$ bases/gene
- Approximately 1% of DNA codes for genes ($10^3 10^4 / 10^9$)



Regulatory Regions

- RNA polymerases seek out *regulatory* or *promoting* regions located 100-1000 bp upstream from the coding region
- They work in conjunction with special proteins called *transcription factors* (*TFs*) whose presence enables gene expression
- Within these regions are the *Transcription Factor Binding Sites (TFBS)*, special DNA sequence patterns known as *motifs* that are specific to a given transcription factor
- A Single TF can influence the expression of many genes. Through biological experiments one can infer, at least a subset of these affected genes.



Transcription Factor Binding Sites

- A TFBS can be located anywhere within the regulatory region.
- TFBS may vary slightly across different regulatory regions since non-essential bases could mutate
- Transcription factors are robust (they will still bind) in the presence of small sequence differences by a few bases



Identifying Motifs: Complications

- We don't know the motif sequence for every TF
- We don't know where it is located relative to a gene's start
- Moreover, motifs can differ slightly from one gene to the next
- We *only* know that it occurs somewhere near genes that share a TF
- How to discern a Motif's frequent *similar* pattern from *random* patterns?
- How is this problem different than finding frequent k-mers from Lecture 2?



Let's look for an *Easy* Motif

```
1 tagggctttgagttagatccgaaggaaagtattccaccagttcgggtcacccagcagggcagggtacttaat  
2 cgcgactcggcgctcacagttatcgacgttagacaaaacggagttagatccgaaactggagttaatcggagtcctt  
3 gttacttgtgagcctggtagatccgaaataataattttggctgcatagcggagctgacatacgagtagggaaatgcgt  
4 aacatcaggcttgattaaacaatttaagcacgttagatccgaattgacctgatgacaatacggAACATGCCGGCTCCGGG  
5 accaccggataggctgctttagatccgaaaggtagtatcgtaataatggctcagccatgtcaatgtgcggcattccac  
6 tagatccgaatcgatcggtttccctctgtggtaacgaggggtccgaccctgctcgatgtgccgaacttgtaccc  
7 gaaatggttcggtgcgatatcaggccgttcttaacttggcggttagatccgaaacgtctggaggggtcgctgccta  
8 atgtatactagacattctaacgctcgcttattggcgagaccattgctccactacaagaggctactgttagatccgaa  
9 ttcttacaccctttagatccgaacctgttggccatcttccgtacccatggctgtacccatgtac  
10 ctacctatgtaaaacaacatctactaacgttagtccggtcttcctgatctggccctaacctacaggttagatccgaaattcg
```

How would you go about finding a 10-mer that appears in *every one* of these strings?

Sneak Peek at the Answer

1 tagggctttgagtg**TAGATCCGAA**ggaaaagtattccaccagttcgggtcaccagcaggcagggtacttaat
2 cgcaactcgccgtcacagttatcgacgttagacaaaacggagt**TAGATCCGAA**actggagttaatcgagtcctt
3 gttacttgtgagcctgg**TAGATCCGAA**atataatttgtggctgcatacgagctgacatacgatggaaatgcgt
4 aacatcaggcttgattaaacaatttaagcacg**TAGATCCGAA**ttgacctgatgacaatacggaacatgccggctccgg
5 accaccggataggctgcttat**TAGATCCGAA**aggttagtatcgtaataatggctcagccatgtcaatgtgcggcattccac
6 **TAGATCCGAA**tgcgtgtttccctctgtggtaacgagggtccgacctgctcgcatgtgccgaacttgtaccc
7 gaaatggttcggtgcgatatcaggccgtcttaacttggcggtg**TAGATCCGAA**cgtctctggagggcgtgccta
8 atgtataactagacattctaaccgtcgcttattggcgagaccattgctccactacaagaggctactgtg**TAGATCCGAA**
9 ttcttacacccttctt**TAGATCCGAA**cctgttggccatcttcttcgagtcctgtacccattgctctgatgac
10 ctacctatgtaaaacaacatctactaacgttagtccggctttcctgatctgccctaacctacagg**TAGATCCGAA**attcg

Now that you've seen the answer, how would you find it?

Meet Mr *Brute Force*

- He's often the best starting point when approaching a problem
- He'll also serve as a straw-man when designing new approaches
- Though he's seldom elegant, he gets the job done
- Often, we can't afford to wait for him



For our current problem a brute force solution would consider every k-mer position in all strings and see if they match. Given M sequences of length N, there are:

$$(N - k + 1)^M$$

position combinations to consider.

A Library of Helper Functions

- There's a tendency to approach this problem with a series of nested for-loops, while the approach is valid, it doesn't generalize. It assumes a specific number of sequences.
- What we need is an *iterator* that generates all permutations of a sequence.
- This nested-for-loop iterator is called a *Cartesian Product* over sets.
- Python has a library to accomplish this

Using *itertools*

```
import itertools

for number in itertools.product("01", repeat=3):
    print ''.join(number)
```

000
001
010
011
100
101
110
111

All permutations of items from a list

```
N = 0
for number in itertools.product(range(3), repeat=3):
    print number,
    N += 1
    if (N % 5 == 0):
        print
```

```
(0, 0, 0) (0, 0, 1) (0, 0, 2) (0, 1, 0) (0, 1, 1)
(0, 1, 2) (0, 2, 0) (0, 2, 1) (0, 2, 2) (1, 0, 0)
(1, 0, 1) (1, 0, 2) (1, 1, 0) (1, 1, 1) (1, 1, 2)
(1, 2, 0) (1, 2, 1) (1, 2, 2) (2, 0, 0) (2, 0, 1)
(2, 0, 2) (2, 1, 0) (2, 1, 1) (2, 1, 2) (2, 2, 0)
(2, 2, 1) (2, 2, 2)
```

Permutations of mixed types

```
for section in itertools.product(("I", "II", "III", "IV"), "ABC", range(1, 3)):  
    print section
```

```
('I', 'A', 1)  
('I', 'A', 2)  
('I', 'B', 1)  
('I', 'B', 2)  
('I', 'C', 1)  
('I', 'C', 2)  
('II', 'A', 1)  
('II', 'A', 2)  
('II', 'B', 1)  
('II', 'B', 2)  
('II', 'C', 1)  
('II', 'C', 2)  
('III', 'A', 1)  
('III', 'A', 2)  
('III', 'B', 1)  
('III', 'B', 2)  
('III', 'C', 1)  
('III', 'C', 2)  
('IV', 'A', 1)  
('IV', 'A', 2)  
('IV', 'B', 1)  
('IV', 'B', 2)  
('IV', 'C', 1)
```

Now let's try some *Brute Force* code

```
sequences = [
    'tagtgtctttgagtgtagatccgaaggaaagtattccaccagttcgggtcacccagcagggcagggtgacttaat',
    'cgcgactcggcgctcacagttatcgcacgttagacaaaacggagttagatccgaaactggagttaatcgagtcctt',
    'gttacttgtgagcctggtagatccgaaataattttttggctgcatacgagctgacatacgacttagggaaatgcgt',
    'aacatcaggcttgattaaacaatttaagcacgttagatccgaattgacctgatgacaatacggAACATGCCGGCTCCGGG',
    'accaccggataggctgcttatttagatccgaaaggtagtatcgtaataatggctcagccatgtcaatgtgcggcattccac',
    'tagatccgaatcgatcggtttctccctctgtgggttaacgagggtccgacctgctcgcatgtgccgaacttgttaccc',
    'gaaatggttcggtgcgatatcaggccgttcttaacttggcggttagatccgaaacgtctcgactggagggtcgctgcgcta',
    'atgtataactagacattctaacgctcgcttattggcgagaccattgctccactacaagaggctactgttagatccgaa',
    'ttcttacaccctttagatccgaaacctgttggcgccatcttgcgtcctgtacccattgctctgatgac',
    'ctacctatgtaaaacaacatctaactaacgttagtccggctttcgatctgccctaacctacaggttagatccgaaattcg']

def bruteForce(dna, k):
    """Finds a *k*-mer common to all sequences from a
       list of *dna* fragments with the same length"""
    t = len(dna)      # how many sequences
    N = len(dna[0])   # length of sequences
    for offset in itertools.product(range(N-k+1), repeat=t):
        for i in xrange(1, len(offset)):
            if dna[0][offset[0]:offset[0]+k] != dna[i][offset[i]:offset[i]+k]:
                break
        else:
            return offset, dna[0][offset[0]:offset[0]+10]
```

Test and then time it

```
N = 4
position, motif = bruteForce(sequences[0:N], 10)
print position, motif
print
for i in xrange(N):
    p = position[i]
    print sequences[i][:p]+sequences[i][p:p+10].upper()+sequences[i][p+10:]
print

%timeit bruteForce(sequences[0:N], 10)
# you can try N = 5, but be prepared to wait
```

(17, 47, 18, 33) tagatccgaa

tagtgttttagtgTAGATCCGAAggaaaagtattccaccagttcgggtcacccagcaggcagggtgacttaat
cgcgactcggcgctcacagttatgcacgttagacaaaacggagtTAGATCCGAAactggagttaatcgagtcctt
gttacttgtgagcctggTAGATCCGAAatataattgtggctgcatggagctgacatacgagtagggaaatgcgt
aacatcaggcttgattaaacaattaaagcacgTAGATCCGAAttgacctgatgacaatacggAACatgccggctccgg

1 loop, best of 3: 5.97 s per loop

Approximate Matching

Now let's consider a more realistic motif finding problem, where the binding sites do not need to match exactly.

```
1 tagggctttgagtTAGATCTGAAggaaagtattccaccagtcgggtcaccagcaggcagggacttaat  
2 cgcactcggcgctcacagttatcgacgttagacaaaacggagtTGGATCCGAAactggagttaatcgagtcctt  
3 gttacttgtgagcctggtTAGACCCGAAatataatttgtggctgcatagcgagctgacatacgagtagggaaatgcgt  
4 aacatcaggcttgattaaacaatttaagcacgTAAATCCGAAttgacctgatgacaatacggaacatgccggctccgg  
5 accaccggataggctgcttatTAGGTCCAAAaggttagtatcgtaataatggctcagccatgtcaatgtcgccattccac  
6 TAGATTCGAAtcgatcggtttccctctgtgggttaacgagggtccgacctgctcgatgtgccgaacttgtaccc  
7 gaaatggttcggtgcgatatcaggccgttcttaacttggcggtgCAGATCCGAAcgtcttggaggggtcgctgccta  
8 atgtataactagacattctaacgctcgcttattggcgagaccattgctccactacaagaggctactgtgTAGATCCGTA  
9 ttcttacacccttcttTAGATCCAAAcctgttggcgccatctttcgagtccattgtacccattgtctgatgac  
10 ctacctatgtaaaacaacatctactaacgttagtccggctttcctgatctggccctaacc tacaggTCGATCCGAAattcg
```

Actually, none of the sequences have an unmodified copy of the original motif

Profile and Consensus

- Align candidate motifs by their start indexes

$$s = (s_1, s_2, \dots, s_t)$$

- Construct a matrix profile with the frequencies of each nucleotide in columns
- Consensus nucleotide in each position has the highest score in each column

| Alignment | | | | | | | |
|-----------|---|---|---|---|---|---|---|
| a | G | g | t | a | c | T | t |
| C | c | A | t | a | c | g | t |
| a | c | g | t | T | A | g | t |
| a | c | g | t | C | C | A | t |
| C | c | g | t | a | c | g | G |

| Profile | A | 3 | 0 | 1 | 0 | 3 | 1 | 1 | 0 |
|---------|---|---|---|---|---|---|---|---|---|
| C | 2 | 4 | 0 | 0 | 1 | 4 | 0 | 0 | 0 |
| G | 0 | 1 | 4 | 0 | 0 | 0 | 3 | 1 | 0 |
| T | 0 | 0 | 0 | 5 | 1 | 0 | 1 | 4 | 0 |

| Consensus | A | C | G | T | A | C | G | T |
|-----------|---|---|---|---|---|---|---|---|
| | | | | | | | | |

Consensus

- One can think of the consensus as an ***ancestor*** motif, from which mutated motifs emerged
- The *distance* between an actual motif and the consensus sequence is generally less than that for any two actual motifs
- *Hamming distance* is number of positions that differ between two strings

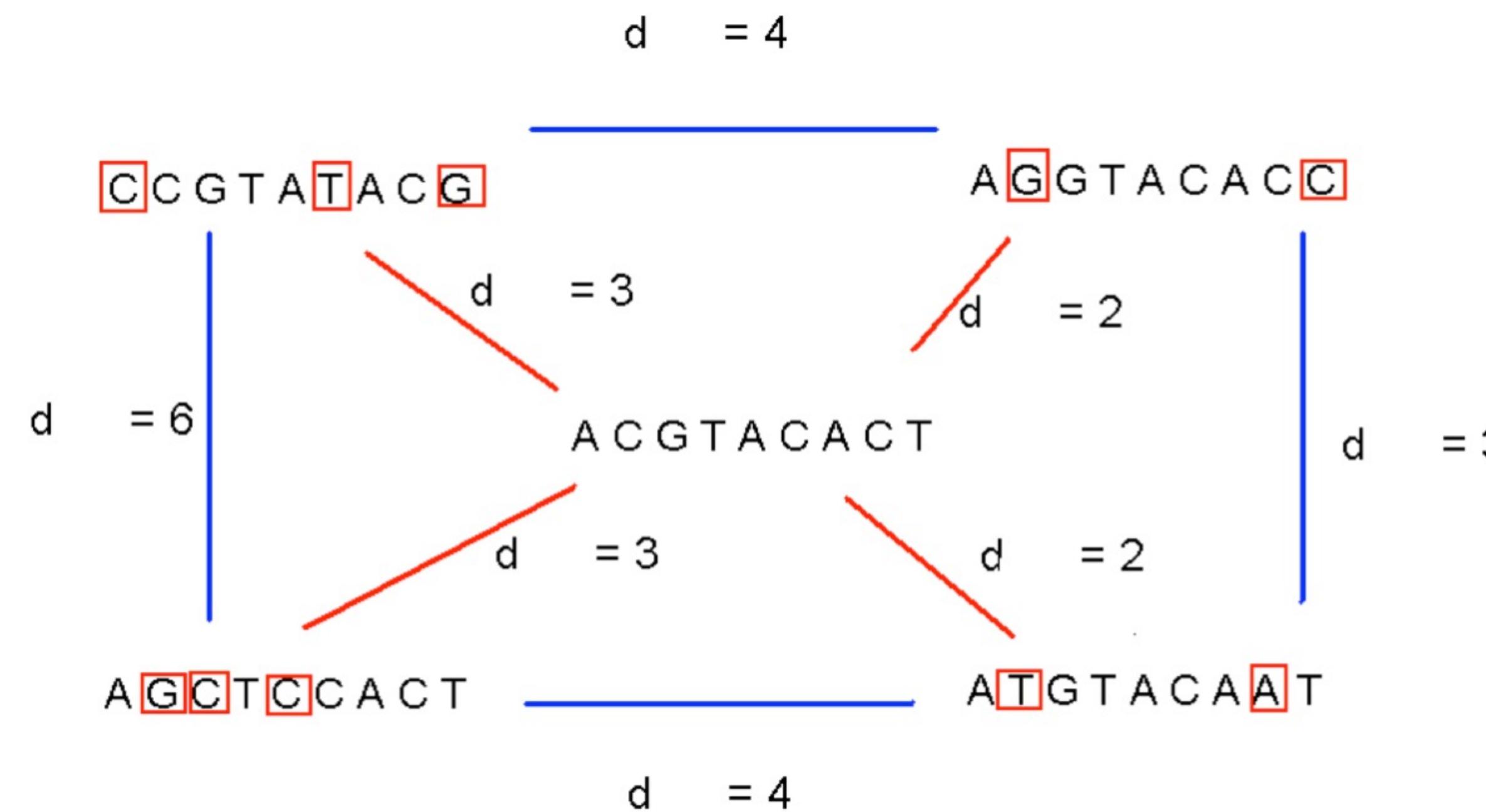
| | | | | | | | | |
|---|---|---|---|---|---|---|---|---|
| G | A | G | A | C | T | C | A | T |
| X | | | | | X | | | |
| T | A | G | A | C | G | C | A | T |



A Hamming
distance of 2

Consensus Properties

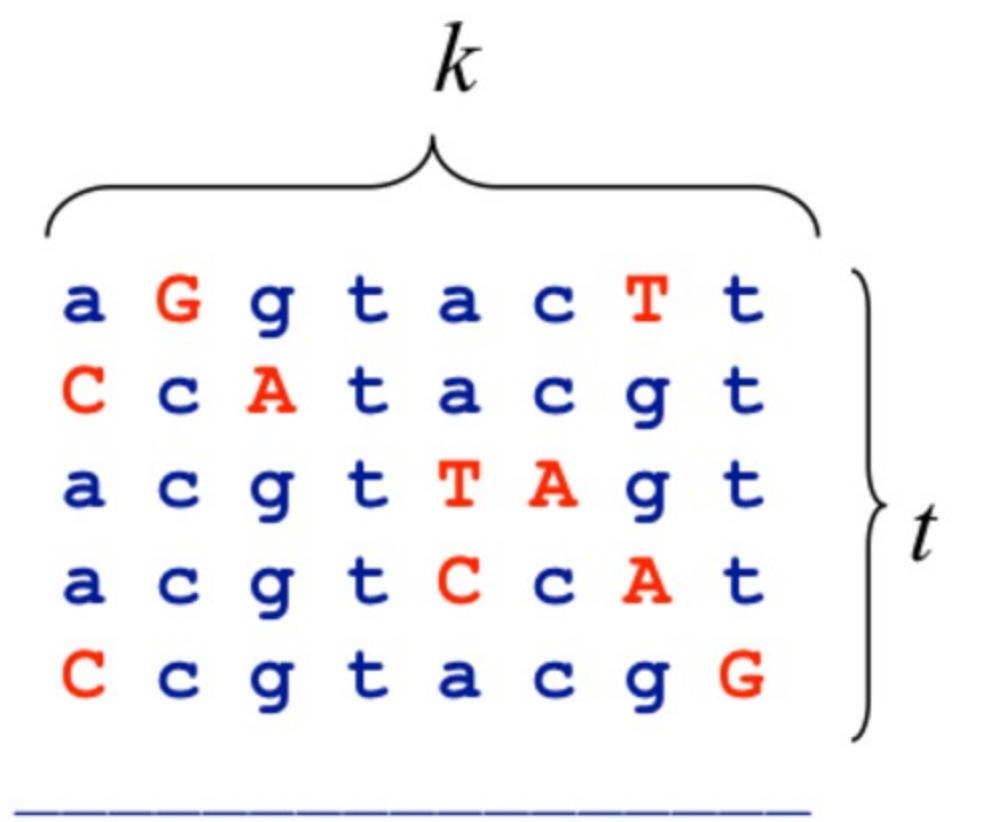
- A consensus string has a minimal hamming distance to all its source strings



Scoring Motifs

- Given $s = (s_1, s_2, \dots, s_t)$ and DNA

$$Score(s, DNA) = \sum_{i=1}^k \max_{j \in A, C, G, T} count(j, i)$$



- So our approach is back to *brute force*
 - We consider every candidate motif in every string
 - Return the set of indices with the highest score

| | | | | | | | | |
|---|---|---|---|---|---|---|---|---|
| A | 3 | 0 | 1 | 0 | 3 | 1 | 1 | 0 |
| C | 2 | 4 | 0 | 0 | 1 | 4 | 0 | 0 |
| G | 0 | 1 | 4 | 0 | 0 | 0 | 3 | 1 |
| T | 0 | 0 | 0 | 5 | 1 | 0 | 1 | 4 |

Consensus a c g t a c g t

Score $3+4+4+5+3+4+3+4=30$

Let's try again, and handle errors this

```
def Score(s, DNA, k):
    """ find the consensus SCORE of a given alignment given offsets into each string.
        s = list of starting indices, DNA = list of sequences, k = Target Motif length """
    score = 0
    for i in xrange(k):
        cnt = dict(zip("acgt", (0,0,0,0)))
        for j, sval in enumerate(s):
            cnt[DNA[j][sval+i]] += 1
        score += max(cnt.values())
    return score

def BruteForceMotifSearch(dna, k):
    t = len(dna)      # how many sequences
    N = len(dna[0])   # length of sequences
    bestScore = 0
    bestAlignment = []
    for offset in itertools.product(range(N-k+1), repeat=t):
        s = Score(offset, dna, k)
        if (s > bestScore):
            bestAlignment = [p for p in offset]
            bestScore = s
    return bestAlignment, bestScore
```

Test and time

```
seqApprox = [  
    'tagtgtttgagtgtagatctgaaggaaagtattccaccagttcggttcacccagcaggcagggtgacttaat',  
    'cgcgactcggcgctcacagttatcgcacgttagacaaaacggagttggatccgaaactggagttaatcgagtcct',  
    'gttacttgcggctggtagaccgaaataattgtggctgcatagcggagctgacatacgagtagggaaatgcgt',  
    'aacatcaggcttgattaaacaatttaagcacgtaaatccgaattgacctgatgacaatacggaacatgccggctccgg',  
    'accacccggataggctgcttatttaggtccaaaaggtagtatcgtaataatggctcagccatgtcaatgtcgccattccac',  
    'tagattcgaatcgatcgatgtttctccctctgtgggttaacgagggtccgacctgctcgatgtgccgaacttgtaccc',  
    'gaaatggttcggtgcgatattcaggccgttcttaacttggcggtgcagatccgaacgtctctggagggtcgctgcta',  
    'atgtataactagacattctaacgctcgcttattggcgagaccattgctccactacaagaggctactgttagatccgta',  
    'ttcttacaccctttagatccaaacctgttggcgccatcttcttgcagtcctgtacctccattgctctgatgac',  
    'ctacctatgtaaaacaacatctaactaacgttagtccggctttcctgatctgccctaacc tacaggtcgatccgaaattcg']  
  
%timeit Score([17, 47, 18, 33, 21, 0, 46, 70, 16, 65], seqApprox, 10)  
%time BruteForceMotifSearch(seqApprox[0:4], 10)
```

10000 loops, best of 3: 76.3 µs per loop
CPU times: user 25min 50s, sys: 10.5 s, total: 26min 1s
Wall time: 25min 49s

([17, 47, 18, 33], 36)

Running Time of BruteForceMotifSearch

- Search $(N - k + 1)$ positions in each of t sequences, by examining $(N - k + 1)^t$ sets of starting positions
- For each set of starting positions, the scoring function makes $O(tk)$ operations, so complexity is

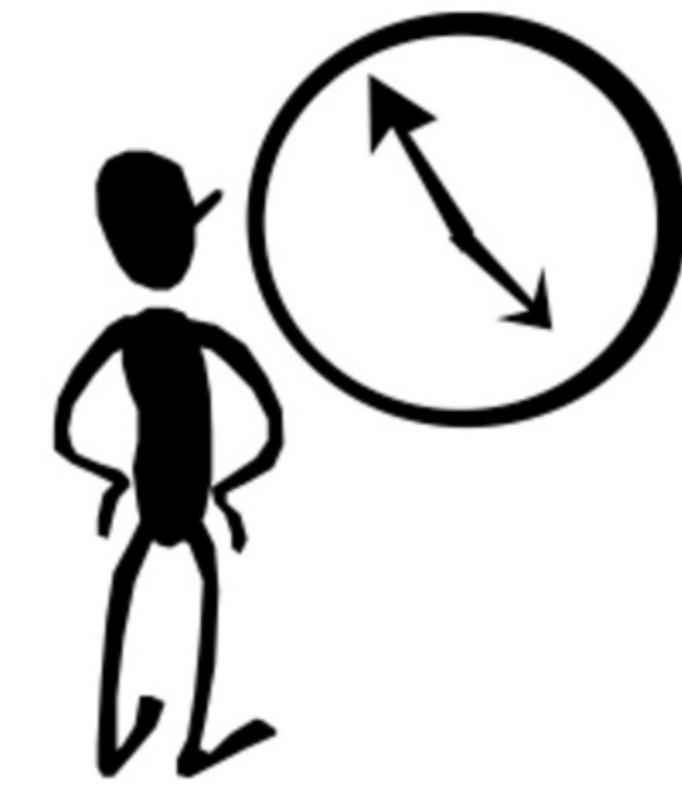
$$tk(N - k + 1)^t = O(tkN^t)$$

- That means that for $t = 10, N = 80, k = 10$ we must perform approximately 10^{21} computations
- Generously assuming 10^9 comps/sec it will require only 10^{12} secs $\frac{10^{12}}{(60*60*24*365)} > 30000$ years
- Want to wait?



How conservative is this estimate?

- For the example we just did $t = 4, N = 80, k = 10$
- So that gives $\approx 1.5 \times 10^9$ operations
- Using our 10^9 operations per second estimate, it should have taken **only 1.5 secs.**
- Instead it took closer to 1500 secs, which suggests we are getting around 1.0 million operations per second.
- So, in reality it will even take longer!



Can we find Motifs in our lifetime?

- Should we give up on Python and write in C? Assembly Language?
- Will biological insights save us this time?
- Are there other ways to find Motifs?
 - Consider that if you knew what motif you were looking for, it would take only

$$k(N-k+1)t = O(kNt)$$

- Is that significantly better?

