#### Exact Matching & CS Fundamentals Michael Schatz

Bioinformatics Lecture I Quantitative Biology 2010



#### Computer Science & Computational Biology

"Computer science is no more about computers than astronomy is about telescopes." Edsger Dijkstra

- Computer Science = Science of Computation
  - Solving problems, designing & building systems
  - Thinking recursively about data, across levels of abstraction
  - Reasoning that your methods are fast & correct
- Computer Science >> Computer Programming
  - Computers are very, very dumb, but we can instruct them
    - Build complex systems out of simple components
    - They will perfectly & repeatedly execute instructions forever
- CompBio = Thinking Computationally about Biology
  - Processing: Make more powerful instruments, analyze results
  - Designing & Understanding: protocols, procedures, systems

# Sequence Alignment Applications

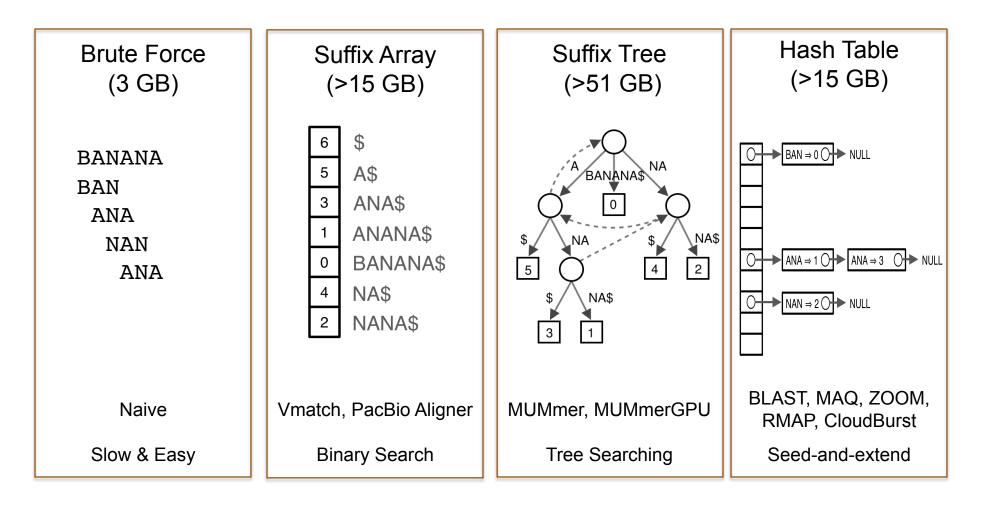
- A very common problem in computational biology is to find occurrences of one sequence in another sequence
  - Genome Assembly
  - Gene Finding
  - Comparative Genomics
  - Functional analysis of proteins
  - Motif discovery
  - SNP analysis
  - Phylogenetic analysis
  - Primer Design

. . .

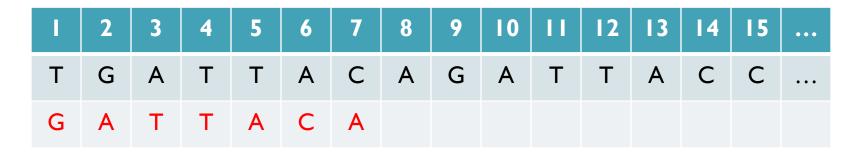
- Personal Genomics

#### Exact Matching Overview

Where is GATTACA in the human genome?

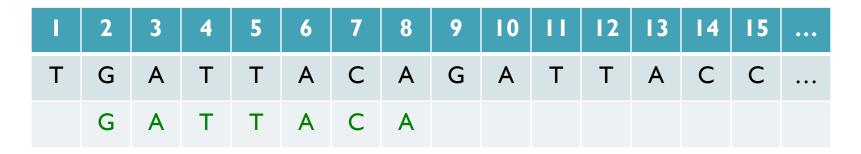


- Where is GATTACA in the human genome?
- Strategy I: Brute Force



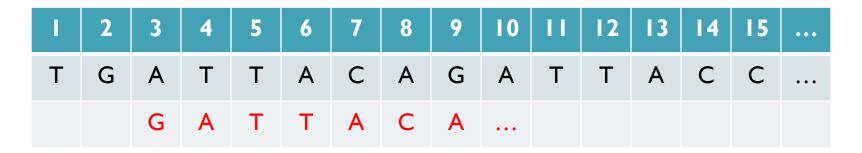
No match at offset I

- Where is GATTACA in the human genome?
- Strategy I: Brute Force



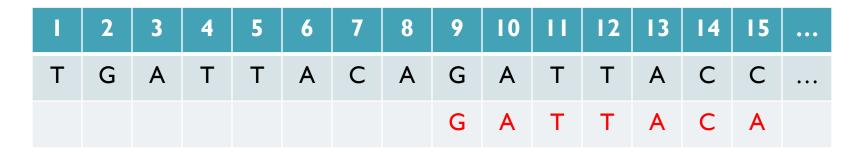
Match at offset 2

- Where is GATTACA in the human genome?
- Strategy I: Brute Force



No match at offset 3...

- Where is GATTACA in the human genome?
- Strategy I: Brute Force



No match at offset 9 <- Checking each possible position takes time

#### Brute Force Analysis

- Brute Force:
  - At every possible offset in the genome:
    - Do all of the characters of the query match?
- Analysis
  - Simple, easy to understand
  - Genome length = n
  - Query length = m
  - Comparisons: (n-m+I) \* m
- Overall runtime: O(nm)
  - If we double genome or query size, takes twice as long
  - If we double both, takes 4 times as long

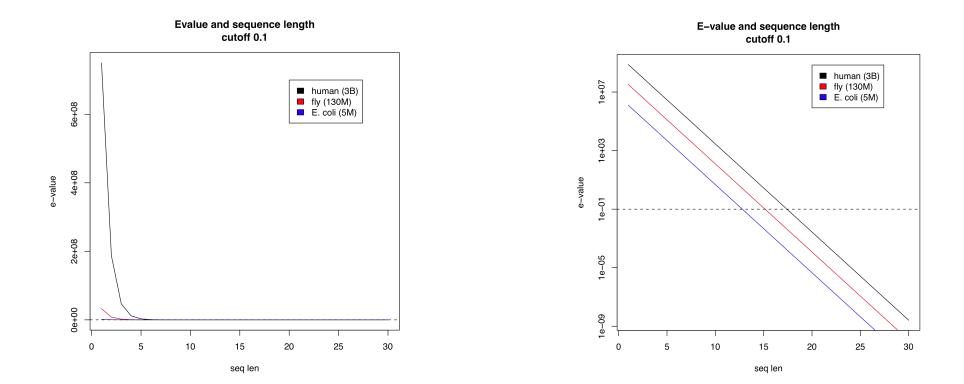
[3B] [7] [21B]

# **Expected Occurrences**

The expected number of occurrences (e-value) of a given sequence in a genome depends on the length of the genome and inversely on the length of the sequence

- I in 4 bases are G, I in 16 positions are GA, I in 64 positions are GAT
- I in 16,384 should be GATTACA
- $E=(n-m+1)/(4^m)$

[183,105 expected occurrences]



#### Brute Force in Matlab

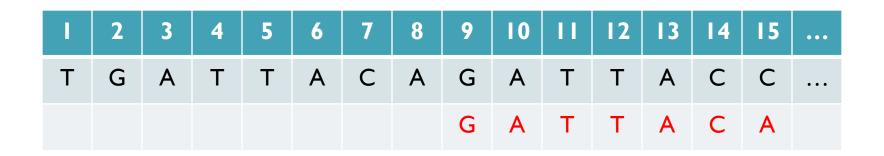
```
query = 'GATTACA';
genome = 'TGATTACAGATTACC';
nummatches=0;
% At every possible offset
for offset=1:length(genome)-length(guery)+1
    % Do all of the characters match?
   if (genome(offset:offset+length(query)-1) == query)
          disp(['Match at offset ', num2str(offset)])
          nummatches = nummatches+1;
    else
          %Uncomment to see every non-match
          %disp(['No match at offset ', num2str(offset)])
    end
end
disp(['Found ', num2str(nummatches),' matches of ', query, ' in genome of length ',
   num2str(length(genome))])
disp(['Expected number of occurrences: ', num2str((length(genome)-length(guery)+1)/
    (4<sup>^</sup>length(query)))))
```

#### **Brute Force Reflections**

Why check every position?

- GATTACA can't start at position 15

[WHY?]



- Improve runtime to O(n + m)

[3B + 7]

- If we double both, it just takes twice as long
- Knuth-Morris-Pratt, 1977
- Boyer-Moyer, 1977, 1991
- For one-off scans, this is the best we can do (optimal performance)
  - We have to read every character of the genome, and every character of the query
  - For short queries, runtime is dominated by the length of the genome

#### 2. Suffix Arrays

- What if we need to check many queries?
  - We don't need to check every page of the phone book to find 'Schatz'
  - Sorting alphabetically lets us immediately skip 96% (25/26) of the book without any loss in accuracy
- Sorting the genome: Suffix Array (Manber & Myers, 1991)
  - Sort every suffix of the genome



Split into n suffixes Sort suffixes alphabetically

[Challenge Question: How else could we split the genome?]

- Strategy 2: Binary search
  - Compare to the middle, refine as higher or lower
- Searching for GATTACA
  - Lo = I; Hi = 15;

Lo	#	Sequence	Pos
->	-	ACAGATTACC	6
	2	ACC	13
	3	AGATTACC	8
	4	ATTACAGATTACC	3
	5	ATTACC	10
	6	C	15
	7	CAGATTACC	7
	8	CC	14
	9	GATTACAGATTACC	2
	10	GATTACC	9
	11	TACAGATTACC	5
	12	TACC	12
	13	TGATTACAGATTACC	I
	14	TTACAGATTACC	4
Hi	15	TTACC	11

- Strategy 2: Binary search ۲
  - Compare to the middle, refine as higher or lower
- Searching for GATTACA ٠
  - Lo = 1; Hi = 15; Mid = (1+15)/2 = 8
  - Middle = Suffix[8] = CC

Lo	#	Sequence	Pos
->	I	ACAGATTACC	6
	2	ACC	13
	3	AGATTACC	8
	4	ATTACAGATTACC	3
	5	ATTACC	10
	6	C	15
	7	CAGATTACC	7
	8	сс	14
	9	GATTACAGATTACC	2
	10	GATTACC	9
	11	TACAGATTACC	5
	12	TACC	12
	13	TGATTACAGATTACC	I
	14	TTACAGATTACC	4
Hi	15	TTACC	11

- Strategy 2: Binary search ۲
  - Compare to the middle, refine as higher or lower
- Searching for GATTACA •
  - Lo = 1; Hi = 15; Mid = (1+15)/2 = 8
  - Middle = Suffix[8] = CC => Higher: Lo = Mid + I

Lo	#	Sequence	Pos
-	I	ACAGATTACC	6
	2	ACC	13
	3	AGATTACC	8
	4	ATTACAGATTACC	3
	5	ATTACC	10
	6	C	15
	7	CAGATTACC	7
	8	CC	14
	9	GATTACAGATTACC	2
	10	GATTACC	9
	11	TACAGATTACC	5
	12	TACC	12
	13	TGATTACAGATTACC	I
	14	TTACAGATTACC	4
Hi	15	TTACC	11

- Strategy 2: Binary search
  - Compare to the middle, refine as higher or lower
- Searching for GATTACA
  - Lo = 1; Hi = 15; Mid = (1+15)/2 = 8
  - Middle = Suffix[8] = CC
     => Higher: Lo = Mid + I
  - Lo = 9; Hi = 15;

	#	Sequence	Pos
	Ι	ACAGATTACC	6
	2	ACC	13
	З	AGATTACC	8
	4	ATTACAGATTACC	3
	5	ATTACC	10
	6	C	15
	7	CAGATTACC	7
Lo →	8	CC	14
	9	GATTACAGATTACC	2
	10	GATTACC	9
	11	TACAGATTACC	5
	12	TACC	12
	13	TGATTACAGATTACC	I
	14	TTACAGATTACC	4
Hi	15	TTACC	11

- Strategy 2: Binary search
  - Compare to the middle, refine as higher or lower
- Searching for GATTACA
  - Lo = 1; Hi = 15; Mid = (1+15)/2 = 8
  - Middle = Suffix[8] = CC
     => Higher: Lo = Mid + 1
  - Lo = 9; Hi = 15; Mid = (9+15)/2 = 12
  - Middle = Suffix[12] = TACC

	#	Sequence	Pos
	I	ACAGATTACC	6
	2	ACC	13
	3	AGATTACC	8
	4	ATTACAGATTACC	3
	5	ATTACC	10
	6	C	15
	7	CAGATTACC	7
Lo	8	CC	14
$\rightarrow$	9	GATTACAGATTACC	2
	10	GATTACC	9
	11	TACAGATTACC	5
	12	TACC	12
	13	TGATTACAGATTACC	I
	14	TTACAGATTACC	4
Hi	15	TTACC	11

- Strategy 2: Binary search •
  - Compare to the middle, refine as higher or lower
- Searching for GATTACA •
  - Lo = 1; Hi = 15; Mid = (1+15)/2 = 8
  - Middle = Suffix[8] = CC = Higher: Lo = Mid + I
  - Lo = 9; Hi = 15; Mid = (9+15)/2 = 12
  - Middle = Suffix[12] = TACC => Lower: Hi = Mid - I
  - Lo = 9; Hi = 11;

	#	Sequence	Pos
	I	ACAGATTACC	6
	2	ACC	13
	3	AGATTACC	8
	4	ATTACAGATTACC	3
	5	ATTACC	10
	6	C	15
	7	CAGATTACC	7
Lo	8	CC	14
-	9	GATTACAGATTACC	2
	10	GATTACC	9
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- Strategy 2: Binary search
  - Compare to the middle, refine as higher or lower
- Searching for GATTACA
  - Lo = 1; Hi = 15; Mid = (1+15)/2 = 8
  - Middle = Suffix[8] = CC
     => Higher: Lo = Mid + 1
  - Lo = 9; Hi = 15; Mid = (9+15)/2 = 12
  - Middle = Suffix[12] = TACC
     => Lower: Hi = Mid 1
  - Lo = 9; Hi = 11; Mid = (9+11)/2 = 10
  - Middle = Suffix[10] = GATTACC

	#	Sequence	Pos
	I	ACAGATTACC	6
	2	ACC	13
	3	AGATTACC	8
	4	ATTACAGATTACC	3
	5	ATTACC	10
	6	C	15
	7	CAGATTACC	7
Lo	8	CC	14
→,	9	GATTACAGATTACC	2
	10	GATTACC	9
Hi	11	TACAGATTACC	5
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	13	TGATTACAGATTACC	I
	14	TTACAGATTACC	4
	15	TTACC	

- Strategy 2: Binary search
  - Compare to the middle, refine as higher or lower
- Searching for GATTACA
  - Lo = 1; Hi = 15; Mid = (1+15)/2 = 8
  - Middle = Suffix[8] = CC
     => Higher: Lo = Mid + 1
  - Lo = 9; Hi = 15; Mid = (9+15)/2 = 12
  - Middle = Suffix[12] = TACC
     => Lower: Hi = Mid 1
  - Lo = 9; Hi = 11; Mid = (9+11)/2 = 10
  - Middle = Suffix[10] = GATTACC
     => Lower: Hi = Mid I
  - Lo = 9; Hi = 9;

#	Sequence	Pos
Ι	ACAGATTACC	6
2	ACC	13
3	AGATTACC	8
4	ATTACAGATTACC	3
5	ATTACC	10
6	C	15
7	CAGATTACC	7
8	CC	14
9	GATTACAGATTACC	2
10	GATTACC	9
	TACAGATTACC	5
12	TACC	12
13	TGATTACAGATTACC	I
14	TTACAGATTACC	4
15	TTACC	

Lo

Hi

- Strategy 2: Binary search
  - Compare to the middle, refine as higher or lower
- Searching for GATTACA
  - Lo = 1; Hi = 15; Mid = (1+15)/2 = 8
  - Middle = Suffix[8] = CC
     => Higher: Lo = Mid + 1
  - Lo = 9; Hi = 15; Mid = (9+15)/2 = 12
  - Middle = Suffix[12] = TACC
     => Lower: Hi = Mid 1
  - Lo = 9; Hi = 11; Mid = (9+11)/2 = 10
  - Middle = Suffix[10] = GATTACC
     => Lower: Hi = Mid I
  - Lo = 9; Hi = 9; Mid = (9+9)/2 = 9
  - Middle = Suffix[9] = GATTACA...
     => Match at position 2!

	#	Sequence	Pos
	Ι	ACAGATTACC	6
	2	ACC	13
	З	AGATTACC	8
	4	ATTACAGATTACC	3
	5	ATTACC	10
	6	C	15
	7	CAGATTACC	7
Lo	8	СС	14
HÌ	9	GATTACAGATTACC	2
-	10	GATTACC	9
		TACAGATTACC	5
	12	TACC	12
	13	TGATTACAGATTACC	I
	14	TTACAGATTACC	4
	15	TTACC	11

## **Binary Search Analysis**

Binary Search

Initialize search range to entire list mid = (hi+lo)/2; middle = suffix[mid] if query matches middle: done else if query < middle: pick low range else if query > middle: pick hi range Repeat until done or empty range

#### [WHEN?]

- Analysis
  - More complicated method
  - How many times do we repeat?
    - How many times can it cut the range in half?
    - Find smallest x such that:  $n/(2^x) \le 1$ ;  $x = \lg_2(n)$  [32]
- Total Runtime: O(m lg n)
  - More complicated, but much faster!
  - Looking up a query loops 32 times instead of 3B

[How long does it take to search 6B nucleotides?]

# Binary Search in Matlab

```
%% create our sorted list of 100 numbers
seq=1:100;
%% seq=sort(floor(rand(100)*100));
query=33;
%% initialize search range
lo=1;
hi=length(seq);
steps=0;
%% search
while (lo<=hi)
  steps = steps+1;
  mid=floor((lo+hi)/2);
  middle=seq(mid);
  disp(['Step ', num2str(steps), ' checking seq[', num2str(mid), ']=', num2str(middle)])
  if (query == middle)
   disp(['Found at ', num2str(mid), ' in ', num2str(steps), ' steps'])
   break
  elseif (query < middle)</pre>
    disp(['less than ', num2str(middle)])
    hi=mid-1;
  else
    disp(['greater than ', num2str(middle)])
    lo=mid+1;
  end
```

```
end
```

## Suffix Array Construction

- Searching the array is very fast, but it takes time to construct
  - This time will be amortized over many, many searches
  - Run it once "overnight" and save it away for all future queries
- How do we store the suffix array?
  - Explicitly storing all n strings is not feasible

$$S = 1 + 2 + 3 + \dots + n = \sum_{i=1}^{n} i = \frac{n(n+1)}{2} = O(n^2)$$

For human genome S = 9 billion billion characters

- Instead use implicit representation
  - Keep I copy of the genome, and a list of sorted offsets
  - Storing 3 billion offsets requires a big server (12GB)
    - Build a separate index for each chromosome

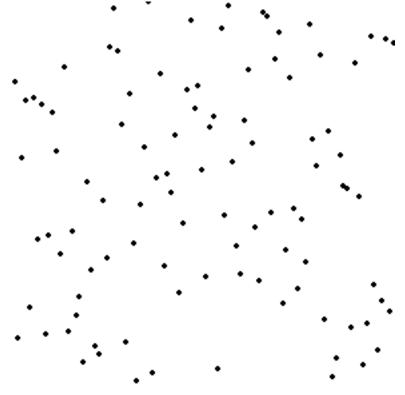
TGATTACAGATTACC

#### Sorting

Sort these numbers into ascending order: 14, 29, 6, 31, 39, 64, 78, 50, 13, 63, 61, 19

[How do you do it?]

6, 13, 14, 29, 31, 39, 64, 78, 50, 63, 61, 19 6, 13, 14, 29, 31, 39, 64, 78, 50, 63, 61, 19 6, 13, 14, 19, 29, 31, 39, 64, 78, 50, 63, 61 6, 13, 14, 19, 29, 31, 39, 64, 78, 50, 63, 61 6, 13, 14, 19, 29, 31, 39, 64, 78, 50, 63, 61 6, 13, 14, 19, 29, 31, 39, 50, 64, 78, 63, 61 6, 13, 14, 19, 29, 31, 39, 50, 61, 64, 78, 63 6, 13, 14, 19, 29, 31, 39, 50, 61, 63, 64, 78 6, 13, 14, 19, 29, 31, 39, 50, 61, 63, 64, 78 6, 13, 14, 19, 29, 31, 39, 50, 61, 63, 64, 78 6, 13, 14, 19, 29, 31, 39, 50, 61, 63, 64, 78 6, 13, 14, 19, 29, 31, 39, 50, 61, 63, 64, 78 6, 13, 14, 19, 29, 31, 39, 50, 61, 63, 64, 78 6, 13, 14, 19, 29, 31, 39, 50, 61, 63, 64, 78 6, 13, 14, 19, 29, 31, 39, 50, 61, 63, 64, 78



http://en.wikipedia.org/wiki/Selection\_sort

#### Selection Sort Analysis

• Selection Sort (Input: list of n numbers)

```
for pos = 1 to n

// find the smallest element in [pos, n]

smallest = pos

for check = pos+1 to n
```

if (list[check] < list[smallest]): smallest = check</pre>

// move the smallest element to the front
tmp = list[smallest]
list[pos] = list[smallest]
list[smallest] = tmp

• Analysis

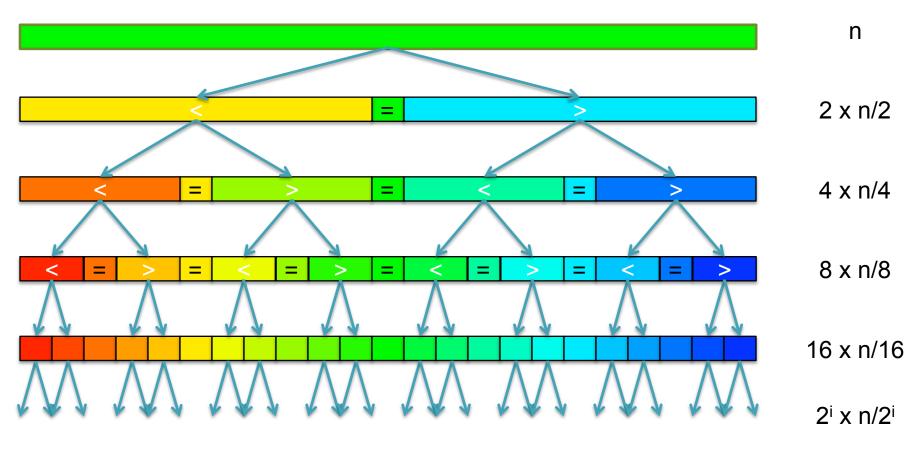
$$T = n + (n - 1) + (n - 2) + \dots + 3 + 2 + 1 = \sum_{i=1}^{n} i = \frac{n(n + 1)}{2} = O(n^2)$$

- Outer loop: pos = I to n
- Inner loop: check = pos to n
- Running time: Outer \* Inner =  $O(n^2)$  [9 Billion Billion]

[Challenge Questions: Why is this slow? / Can we sort any faster?]

#### **Divide and Conquer**

- Selection sort is slow because it rescans the entire list for each element
  - How can we split up the unsorted list into independent ranges?
  - Hint I: Binary search splits up the problem into 2 independent ranges (hi/lo)
  - Hint 2: Assume we know the median value of a list

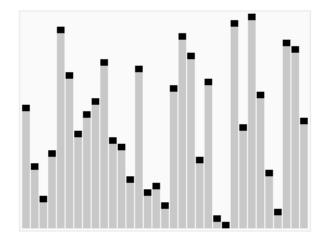


[How many times can we split a list in half?]

#### QuickSort Analysis

QuickSort(Input: list of n numbers)
 // see if we can quit
 if (length(list)) <= 1): return list</li>

```
// split list into lo & hi
pivot = median(list)
lo = {}; hi = {};
for (i = I to length(list))
        if (list[i] < pivot): append(lo, list[i])
        else: append(hi, list[i])</pre>
```



http://en.wikipedia.org/wiki/Quicksort

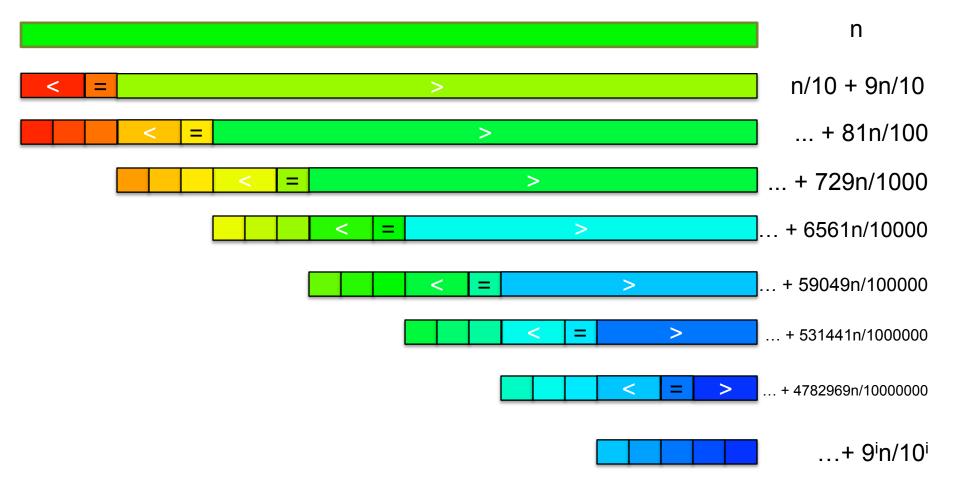
// recurse on sublists
return (append(QuickSort(lo), QuickSort(hi))

• Analysis (Assume we can find the median in O(n))

$$T(n) = \begin{cases} O(1) & \text{if } n \le 1\\ O(n) + 2T(n/2) & \text{else} \end{cases}$$
  
$$T(n) = n + 2(\frac{n}{2}) + 4(\frac{n}{4}) + \dots + n(\frac{n}{n}) = \sum_{i=0}^{lg(n)} \frac{2^{i}n}{2^{i}} = \sum_{i=0}^{lg(n)} n = O(n \lg n) \quad [\sim 94B]$$

## Picking the Median

• What if we miss the median and do a 90/10 split instead?



[How many times can we cut 10% off a list?]

# Randomized Quicksort

- 90/10 split runtime analysis  $T(n) = n + T(\frac{n}{10}) + T(\frac{9n}{10})$   $T(n) = n + \frac{n}{10} + T(\frac{n}{100}) + T(\frac{9n}{100}) + \frac{9n}{10} + T(\frac{9n}{100}) + T(\frac{81n}{100})$   $T(n) = n + n + T(\frac{n}{100}) + 2T(\frac{9n}{100}) + T(\frac{81n}{100})$   $T(n) = \sum_{i=0}^{\log_{10/9}(n)} n = O(n \lg n)$ Find smallest x s.t. (9/10)<sup>x</sup> n \le 1  $(10/9)^{x} \ge n$   $x \ge \log_{10/9} n$
- If we randomly pick a pivot, we will get at least a 90/10 split with very high probability
  - Everything is okay as long as we always slice off a fraction of the list

[Challenge Question:What happens if we slice I element]

# QuickSort in Matlab

sort(seq)

- The goal of software engineering is to build libraries of correct reusable functions that implement higher level ideas
  - Build complex software out of simple components
  - Software tends to be 90% plumbing, 10% research
  - You still need to know how they work
    - Matlab requires an explicit representation of the strings

## Break

#### Sorting in Linear Time

- Can we sort faster than O(n lg n)?
  - No Not if we have to compare elements to each other
  - Yes But we have to 'cheat' and know the structure of the data

Sort these numbers into ascending order: 14, 29, 6, 31, 39, 64, 78, 50, 13, 63, 61, 19

I,2,3,4,5,6,7,8,9,10,11,12,13,14,15,16,17,18,19,20,21,22,23,24,25, 26,27,28,29,30,31,32,33,34,35,36,37,38,39,40,41,42,43,44,45,46,47,48,49,50, 51,52,53,54,55,56,57,58,59,60,61,62,63,64,65,66,67,68,69,70,71,72,73,74,75, 76,77,78,79,80,81,82,83,84,85,86,87,88,89,90,91,92,93,94,95,96,97,98,99,100

#### Sorting in Linear Time

- Can we sort faster than O(n lg n)?
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Sort these numbers into ascending order: 14, 29, 6, 31, 39, 64, 78, 50, 13, 63, 61, 19

I,2,3,4,5,6,7,8,9,10,11,12,13,14,15,16,17,18,19,20,21,22,23,24,25, 26,27,28,29,30,31,32,33,34,35,36,37,38,39,40,41,42,43,44,45,46,47,48,49,50, 51,52,53,54,55,56,57,58,59,60,61,62,63,64,65,66,67,68,69,70,71,72,73,74,75, 76,77,78,79,80,81,82,83,84,85,86,87,88,89,90,91,92,93,94,95,96,97,98,99,100

#### Sorting in Linear Time

- Can we sort faster than O(n lg n)?
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Sort these numbers into ascending order: 14, 29, 6, 31, 39, 64, 78, 50, 13, 63, 61, 19

I,2,3,4,5,6,7,8,9,10,11,12,13,14,15,16,17,18,19,20,21,22,23,24,25, 26,27,28,29,30,31,32,33,34,35,36,37,38,39,40,41,42,43,44,45,46,47,48,49,50, 51,52,53,54,55,56,57,58,59,60,61,62,63,64,65,66,67,68,69,70,71,72,73,74,75, 76,77,78,79,80,81,82,83,84,85,86,87,88,89,90,91,92,93,94,95,96,97,98,99,100

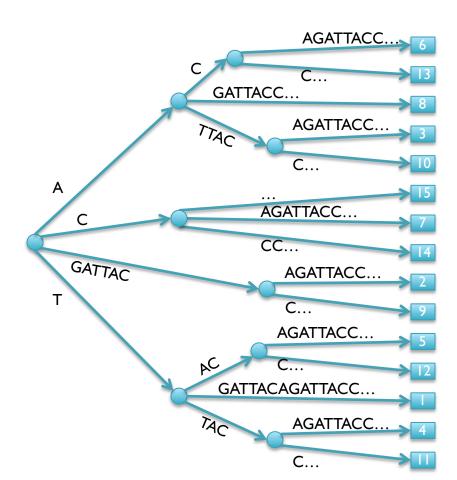
#### 6, 13, 14, 19, 29, 31, 39, 50, 61, 63, 64, 78

for(i = I to I00) { range[i] = 0; }
for(i = I to n) { range[list[i]] = I; }
for(i = I to I00) { if (range[i] == I){print i}}

[3B instead of 94B]

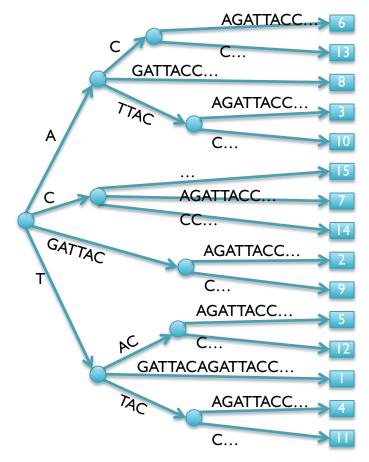
#### 3. Suffix Trees

#	Sequence	Pos
Ι	ACAGATTACC	6
2	ACC	13
3	AGATTACC	8
4	ATTACAGATTACC	3
5	ATTACC	10
6	C	15
7	CAGATTACC	7
8	CC	14
9	GATTACAGATTACC	2
10	GATTACC	9
11	TACAGATTACC	5
12	TACC	12
13	TGATTACAGATTACC	I
14	TTACAGATTACC	4
15	TTACC	11



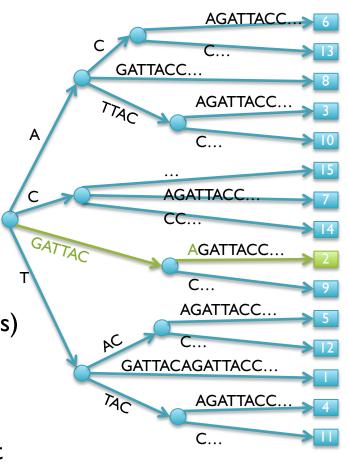
- Suffix Tree = Tree of suffixes (indexes **all** substrings of a sequence)
  - I Leaf (\$) for each suffix, path-label to leaf spells the suffix
  - Nodes have at least 2 and at most 5 children (A,C,G,T,\$)

- Look up a query by "walking" along the edges of the tree
  - GATTACA

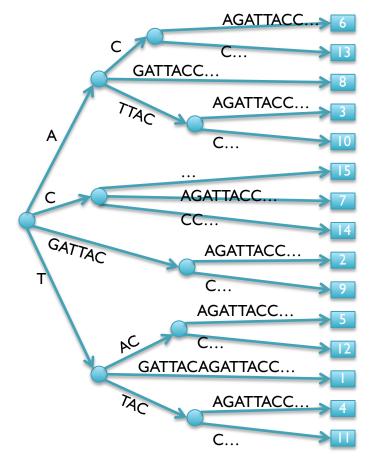


- Look up a query by "walking" along the edges of the tree
  - GATTACA
  - Matches at position 2

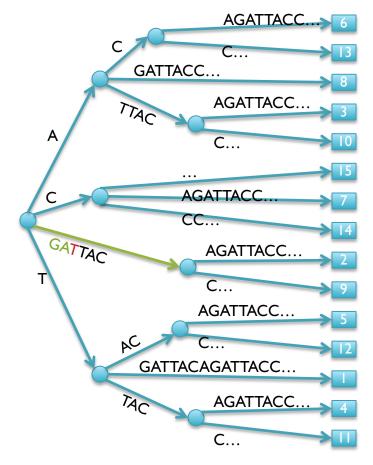
```
WalkTree
   cur = ST.Root; qrypos = 0;
   while (cur)
        // check for partial matches
        // walk the tree
        edge = cur.getEdge(q[qrypos]); edgepos=0
        dist = matchstrings(edge, edgepos, qry, qrypos)
        if (qrypos+dist == length(qry))
             print "end-to-end match"
        else if (dist == length(edge))
             cur=cur.getNode(edge[0]); qrypos+=dist
        else
             print "no match"
```



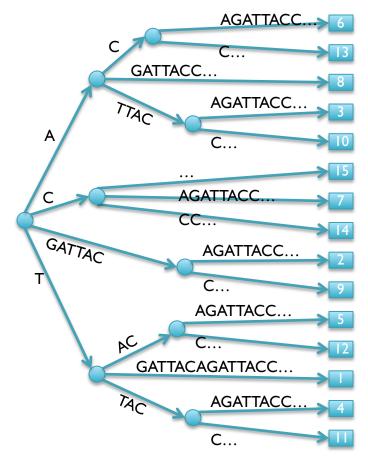
- Look up a query by "walking" along the edges of the tree
  - GACTACA



- Look up a query by "walking" along the edges of the tree
  - GACTACA
  - Fell off tree no match



- Look up a query by "walking" along the edges of the tree
  - ATTAC

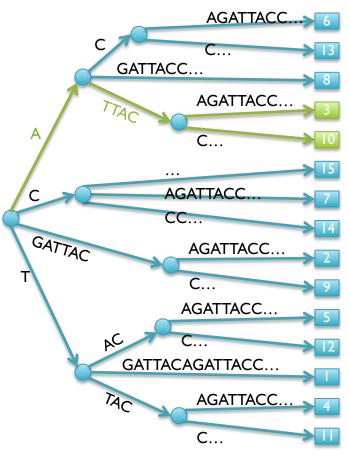


- Look up a query by "walking" along the edges of the tree
  - ATTAC
  - Matches at 3 and 10
  - Query Lookup in 2 phases:
    - I. Walk along edges to find matches
    - 2. Walk subtree to find positions

DepthFirstPrint(Node cur) if cur.isLeaf print cur.pos else foreach child in cur.children

DepthFirstPrint(child)

[What is the running time of DFP => How many nodes does the tree have?]

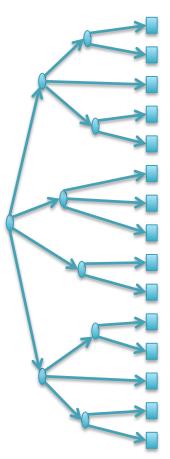


### Suffix Tree Properties & Applications

Properties

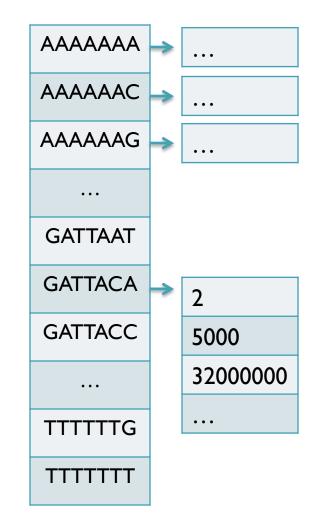
- Number of Nodes/Edges: O(n)
- Tree Size: O(n)
- Max Depth: O(n)
- Construction Time: O(n)
  - Uses suffix links to jump between nodes without rechecking
  - Tricky to implement, prove efficiency
- Applications
- Sorting all suffixes: O(n)
- Check for query: O(m)
- Find all z occurrences of a query O(m + z)
- Find maximal exact matches O(m)
- Longest common substring O(m)
- Used for many string algorithms in linear time
  - Many can be implemented on suffix arrays using a little extra work

[HOW?]



# 4. Hashing

- Where is GATTACA in the human genome?
  - Build an inverted index (table) of every kmer in the genome
- How do we access the table?
  - We can only use numbers to index
    - table[GATTACA] <- error, does not compute</li>
  - Encode sequences as numbers
    - Easy:  $A = I_{10}, C = 2_{10}, G = 3_{10}, T = 4_{10}$ - GATTACA = 3144121<sub>10</sub>
    - Smart:  $A = 00_2$ ,  $C = 01_2$ ,  $G = 10_2$ ,  $T = 11_2$ - GATTACA = 10001111000100\_2 = 9156\_10
  - Running time
    - Construction: O(n)
    - Lookup: O(I) + O(z)
    - Sorts the genome mers in linear time



## Hash Tables and Hash Functions

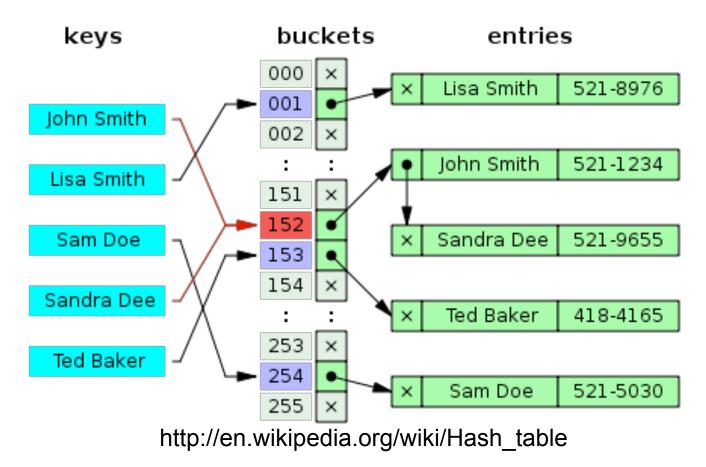
- Number of possible sequences of length  $k = 4^k$ 
  - $-4^7 = 16,384$  (easy to store)
  - $4^{20} = 1,099,511,627,776$  (impossible to directly store in RAM)
    - There are only 3B 20-mers in the genome
      - $\Rightarrow$  Even if we could build this table, 99.7% will be empty

 $\Rightarrow$  But we don't know which cells are empty until we try

- Hash Function: hash(n) -> h
  - Maps a number *n* in [0,R] to h in [0,H] where H << R
    - More than one *n* will have the same *h*
    - A good hash function evenly distributes the values
      - R/H have the same hash value
    - A really good hash function also spreads out the values
      - Pr(hash(n)==hash(m)) = I/H
    - A common (decent) choice is hash(n) = n mod H

## Hash Table Lookup

- By construction, multiple keys have the same hash value
  - Store elements with the same key in a bucket chained together
  - Looking up a value scans the entire bucket
    - Slows down the search as a function of the hash table load
    - Warning: This complexity is usually hidden in the hash table code

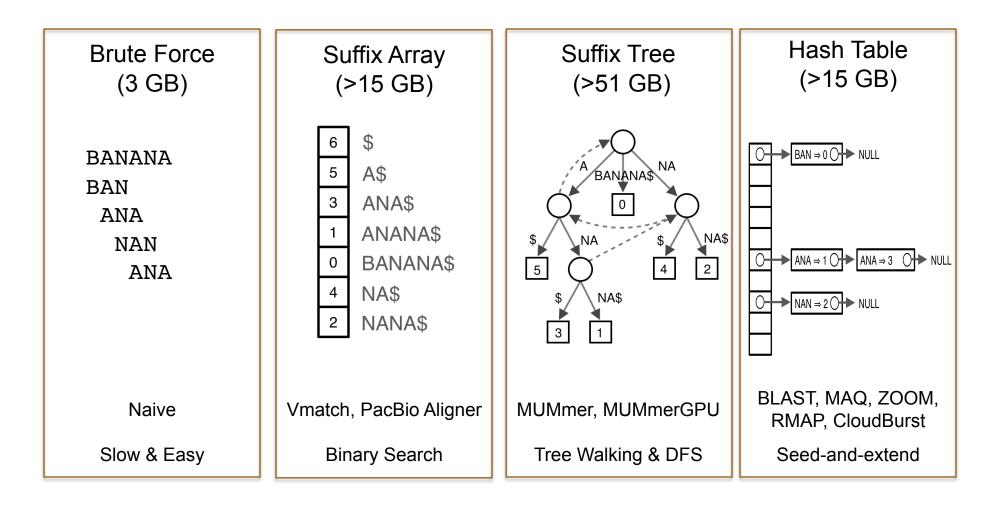


# Variable Length Queries

- Where are GATTACA and GATTACCA in the human genome?
  - s = min(length of all queries)
  - Build an inverted index of all s-mers (seeds) in the genome
    - GATTACA => 2, 5000, 32000000, ...
    - GATTACC => 5500, 10101, 1000000, ...
- Seed-and-extend to find end-to-end exact matches
  - Check every occurrence of the qry seed (first s characters)
    - ~I in 4 are GATTACCA, I in 4 are GATTACCC, etc
  - The specificity of the seed depends on length(q) & s
    - Works best if max(length) =~ min(length)
    - Works best if e-value(m) is << 1</li>

### **Exact Matching Review**

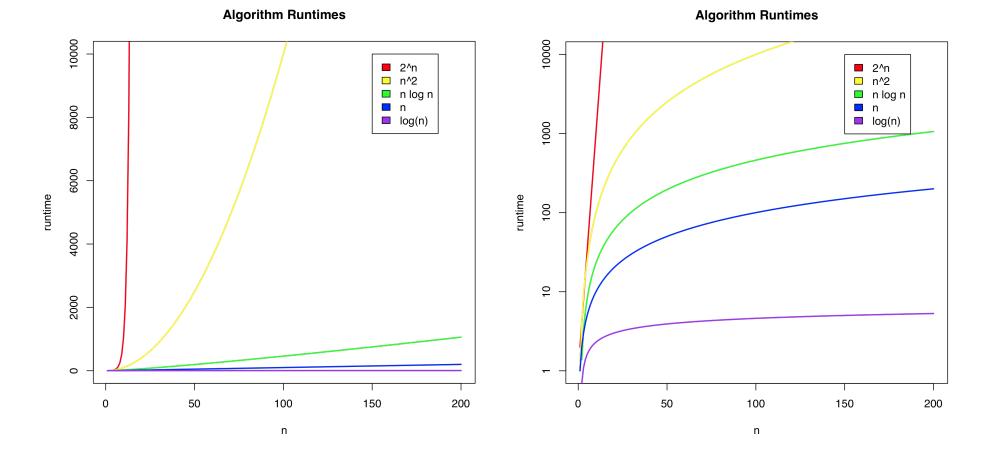
- E-value depends on length of genome and inversely on query length
  - $E = (n-m+1)/4^m$



### Algorithms Summary

- Algorithms choreograph the dance of data inside the machine
  - Algorithms add provable precision to your method
  - A smarter algorithm can solve the same problem with much less work
- Techniques
  - Binary search: Fast lookup in any sorted list
  - Divide-and-conquer: Split a hard problem into an easier problem
  - Recursion: Solve a problem using a function of itself
  - Randomization: Avoid the demon
  - Hashing: Storing sets across a huge range of values
  - Indexing: Focus on the search on the important parts
    - Different indexing schemes have different space/time features
- Data Structures
  - Primitives: Integers, Numbers, Strings
  - Lists / Arrays / Multi-dimensional arrays
  - Trees
  - Hash Table

#### Algorithmic Complexity



What is the runtime as a function of the input size?

## Next Time

- In-exact alignment
  - Smith & Waterman (1981) Identification of Common Molecular Subsequences. J. of Molecular Biology. 147:195-197.
- Sequence Homology
  - Altschul SF, Gish W, Miller W, Myers EW, Lipman DJ (1990). Basic local alignment search tool. J of Molecular Biology. 215 (3): 403–410.
- Whole Genome Alignment
  - A.L. Delcher, S. Kasif, R.D. Fleischmann, J. Peterson, O. White, and S.L. Salzberg (1999) Alignment of Whole Genomes. Nucleic Acids Research (27):11 2369-2376.
- Short Read Mapping
  - Langmead B, Trapnell C, Pop M, Salzberg SL. (2009) Ultrafast and memoryefficient alignment of short DNA sequences to the human genome. Genome Biology. 10:R25.