Ultra Large Scale DNA Sequence Analysis

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CSHL In-House Symposium XXIV

Outline

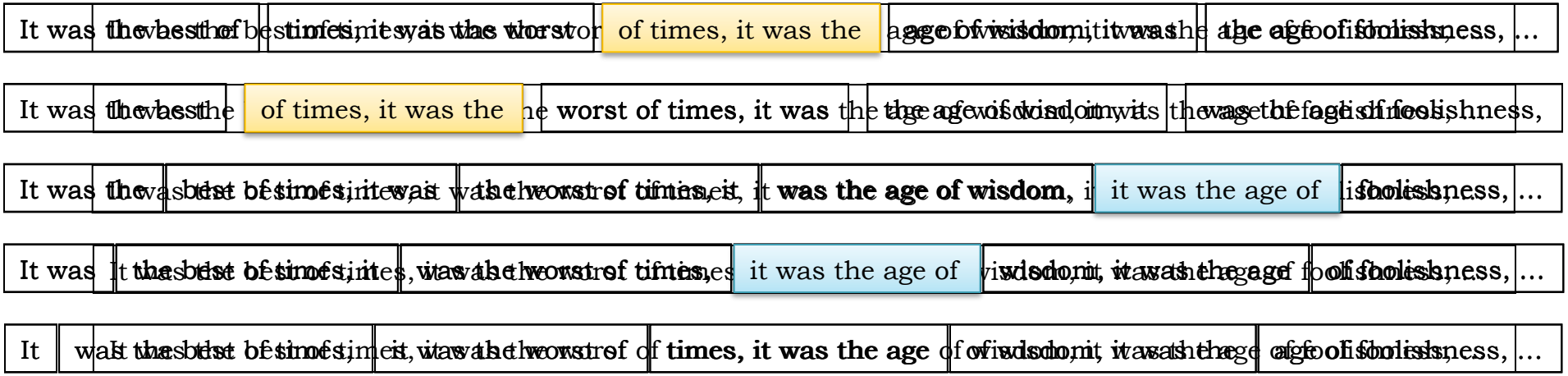
1. Genome Assembly by Analogy

2. DNA Sequencing and Genomics

3. Sequence Analysis Projects
    1. Mapping & Genotyping
    2. Microsatellite Profiling
    3. De novo assembly

Shredded Book Reconstruction

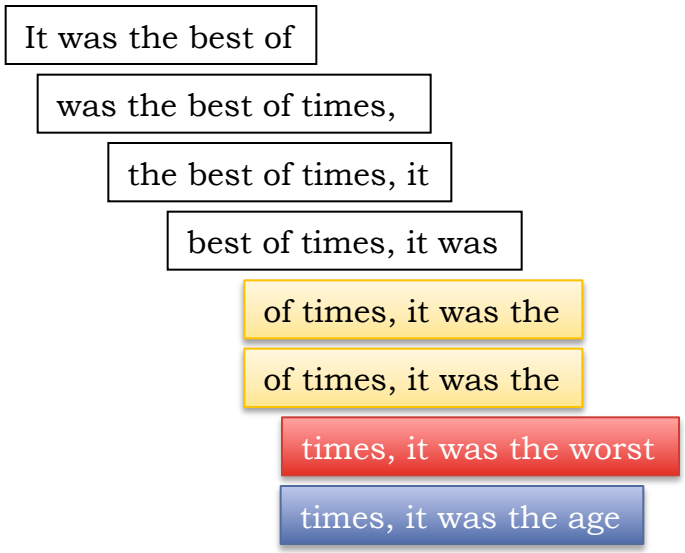
- Dickens accidentally shreds the first printing of A Tale of Two Cities
  - Text printed on 5 long spools



- How can he reconstruct the text?
  - 5 copies x 138, 656 words / 5 words per fragment = 138k fragments
  - The short fragments from every copy are mixed together
  - Some fragments are identical

Greedy Reconstruction

It was the best of
age of wisdom, it was
best of times, it was
it was the age of
it was the age of
it was the worst of
of times, it was the
of times, it was the
of wisdom, it was the
the age of wisdom, it
the best of times, it
the worst of times, it
times, it was the age
times, it was the worst
was the age of wisdom,
was the age of foolishness,
was the best of times,
was the worst of times,
wisdom, it was the age
worst of times, it was



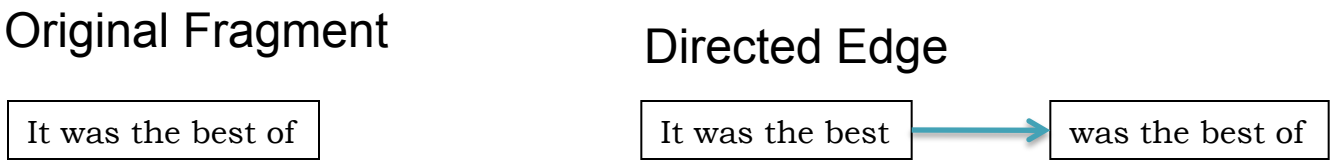
The repeated sequence make the correct reconstruction ambiguous

- It was the best of times, it was the [worst/age]

Model sequence reconstruction as a graph problem.

de Bruijn Graph Construction

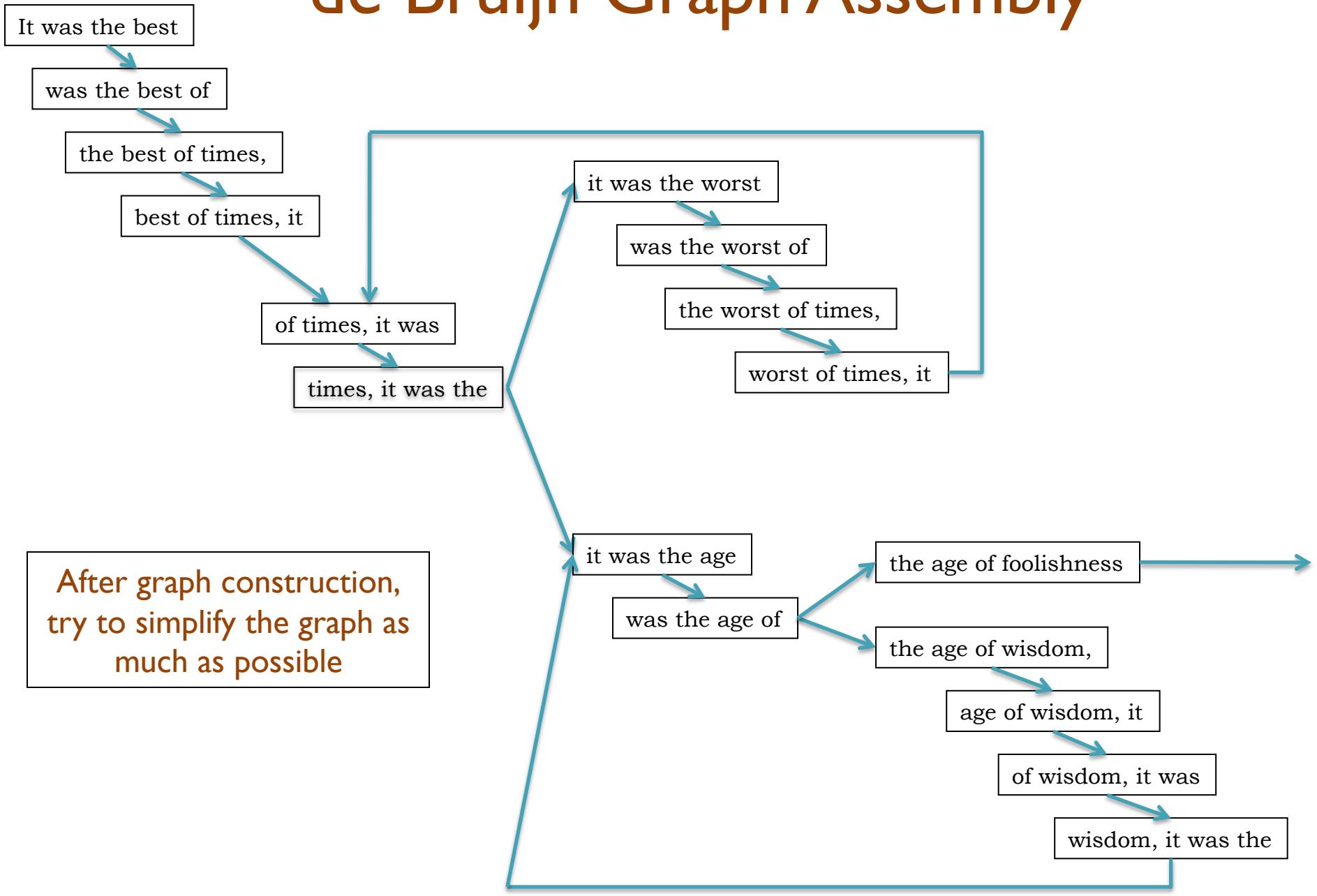
- $D_k = (V,E)$
  - V = All length-k subfragments (k < l)
  - E = Directed edges between consecutive subfragments
    - Nodes overlap by k-1 words



- Locally constructed graph reveals the global sequence structure
  - Overlaps between sequences implicitly computed

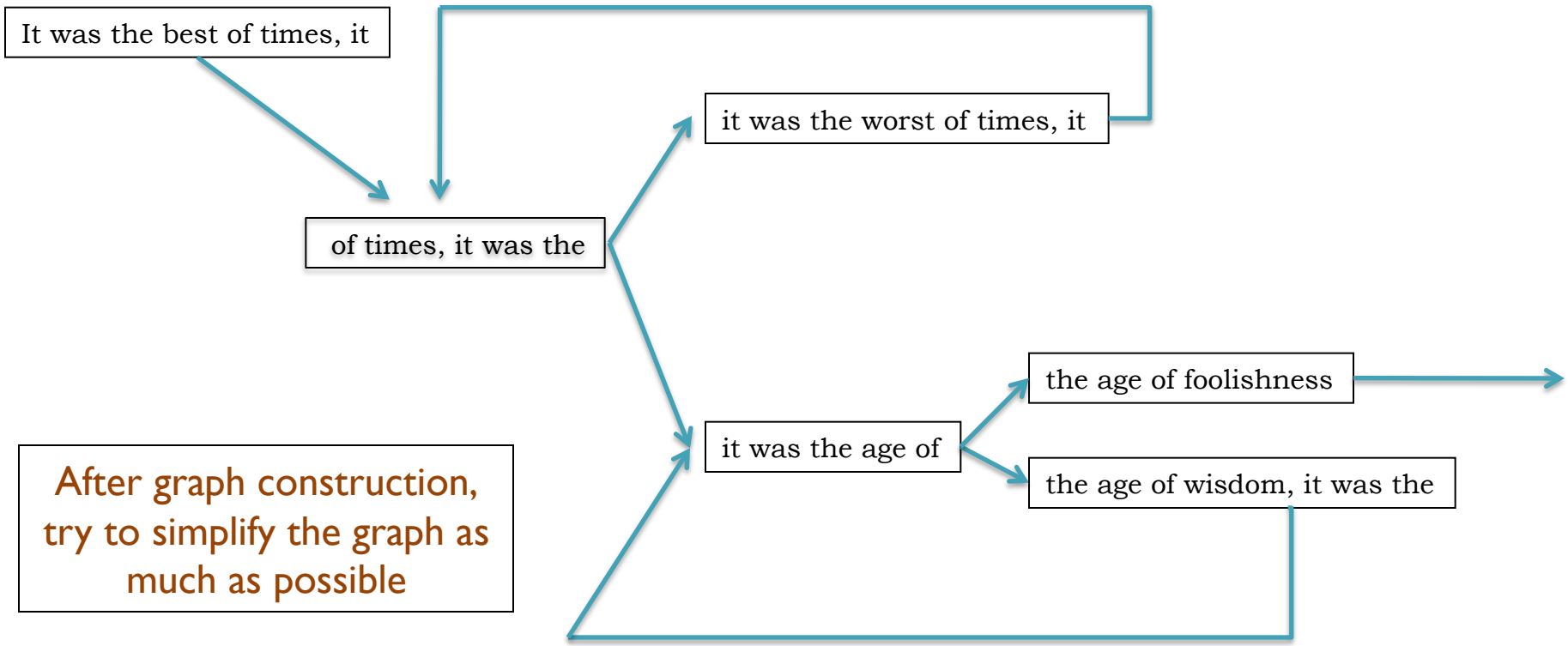
de Bruijn, 1946
Idury and Waterman, 1995
Pevzner, Tang, Waterman, 2001

de Bruijn Graph Assembly



After graph construction, try to simplify the graph as much as possible

de Bruijn Graph Assembly

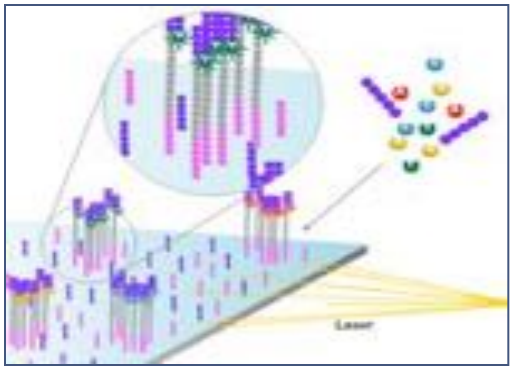


After graph construction, try to simplify the graph as much as possible

Genomics & DNA Sequencing

Genome of an organism encodes the genetic information in long sequence of 4 DNA nucleotides: ACGT

- Bacteria: ~3 million bp
- Humans: ~3 billion bp



Current DNA sequencing machines can sequence millions of short (25-500bp) reads from random positions of the genome

- Per-base error rate estimated at 1-2% (Simpson *et al*, 2009)

ATCTGATAAGTCCCAGGACTTCAGT

GCAAGGCAAACCCGAGCCCAGTTT

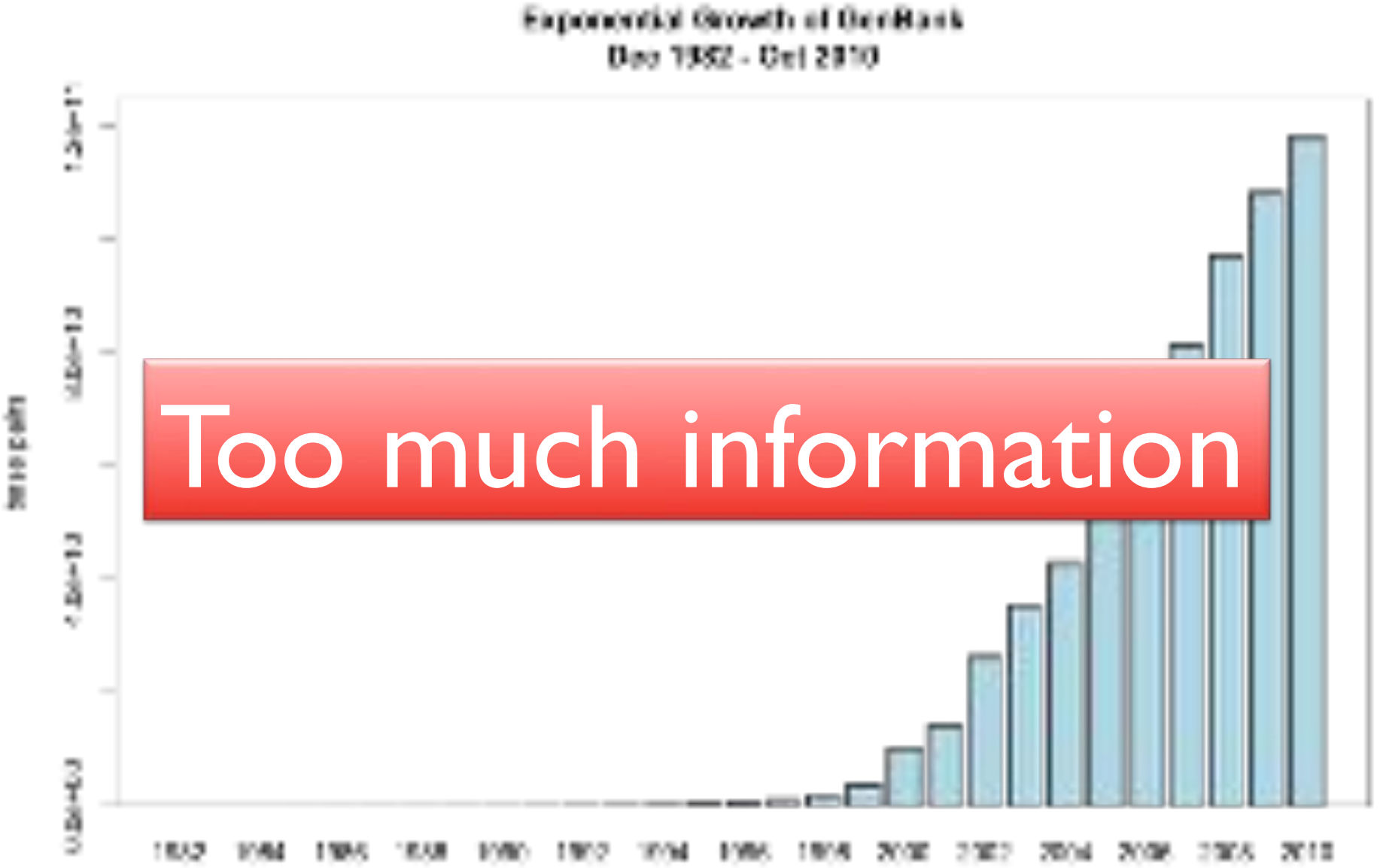
TCCAGTTCTAGAGTTTCACATGATC

GGAGTTAGTAAAAGTCCACATTGAG

Like Dickens, we can only sequence small fragments of the genome at once.

- A single human genome requires ~100 GB of raw data
- We need extremely scalable systems and algorithms

The DNA Deluge



ftp://ftp.ncbi.nih.gov/genbank/gbrel.txt

Hadoop MapReduce

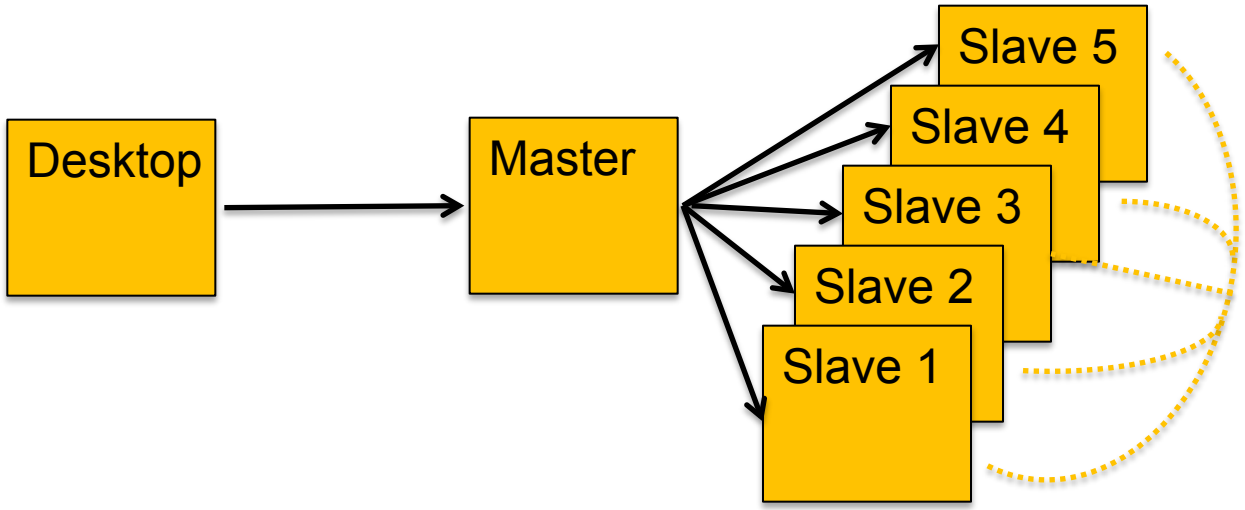
http://hadoop.apache.org

- MapReduce is Google's framework for large data computations
  - Data and computations are spread over thousands of computers
    - Indexing the Internet, PageRank, Machine Learning, etc… (Dean and Ghemawat, 2004)
    - 946,460 TB processed in May 2010 (Jeff Dean at Stanford, 11.10.2010)
  - Hadoop is the leading open source implementation
    - Developed and used by Yahoo, Facebook, Twitter, Amazon, etc
    - GATK is an alternative implementation specifically for NGS

- Benefits
  - Scalable, Efficient, Reliable
  - Easy to Program
  - Runs on commodity computers

- Challenges
  - Redesigning / Retooling applications
    - Not Condor, Not MPI
    - Everything in MapReduce

System Architecture

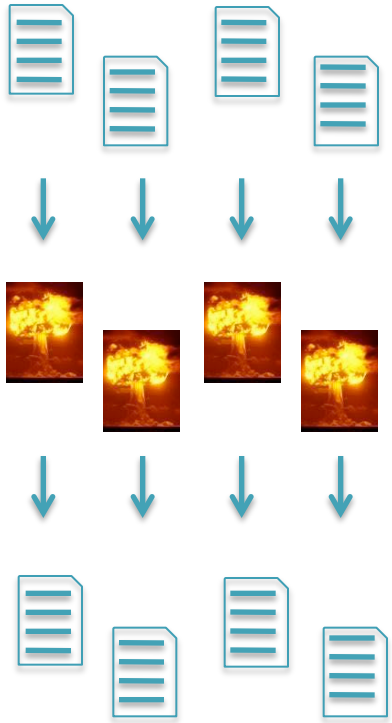


- Hadoop Distributed File System (HDFS)
  - Data files partitioned into large chunks (64MB), replicated on multiple nodes
  - Computation moves to the data, rack-aware scheduling

- Hadoop MapReduce system won the 2009 GreySort Challenge
  - Sorted 100 TB in 173 min (578 GB/min) using 3452 nodes and 4x3452 disks

Programming Models

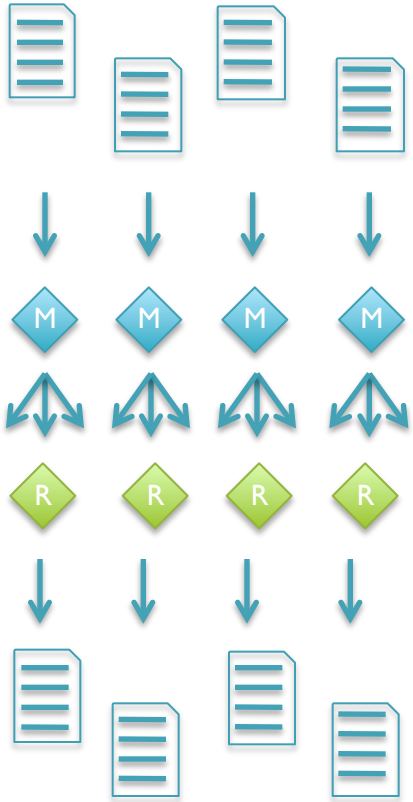
Embarrassingly Parallel



Map-only

Each item is Independent
Traditional Batch Computing

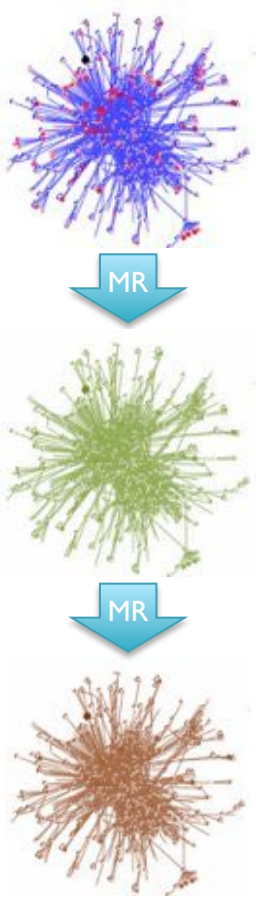
Loosely Coupled



MapReduce

Independent-Shuffle-Independent
Batch Computing + Data Exchange

Tightly Coupled



Iterative MapReduce

Nodes interact with other nodes
Big Data MPI

Short Read Mapping

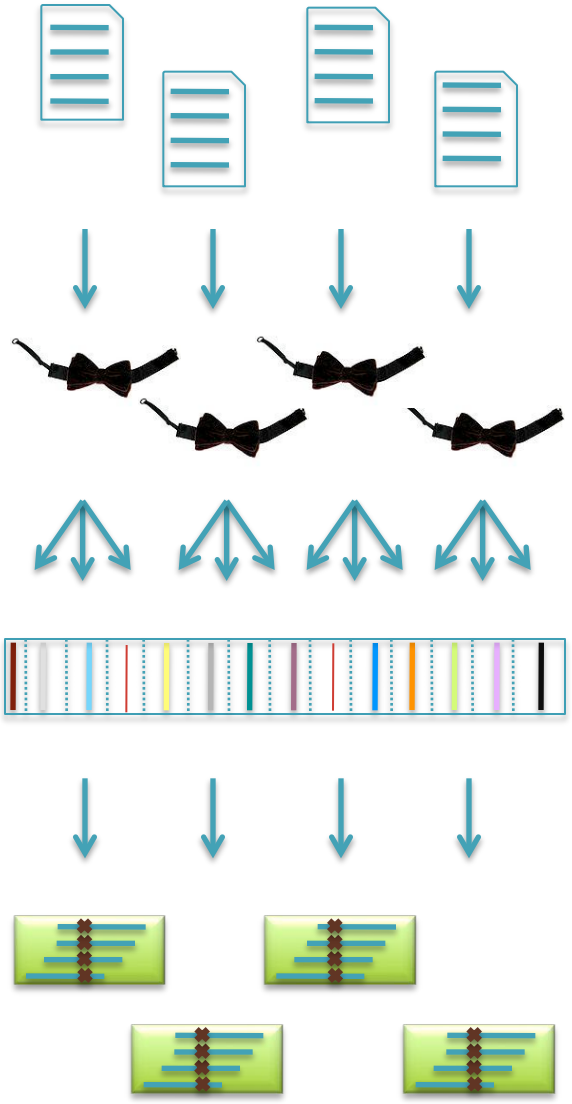
- Given a reference and many subject reads, report one or more "good" end-to-end alignments per alignable read
  - Find where the read most likely originated
  - Fundamental computation for many assays
    - Genotyping
    - Structural Variations
    - RNA-Seq
    - Chip-Seq
    - Methyl-Seq
    - Hi-C-Seq

- Desperate need for scalable solutions
  - Single human requires >1,000 CPU hours / genome

Crossbow

http://bowtie-bio.sourceforge.net/crossbow

- Align billions of reads and find SNPs
  - Reuse software components: Hadoop Streaming
- Map: Bowtie (Langmead *et al.*, 2009)
  - Find best alignment for each read
  - Emit (chromosome region, alignment)
- Shuffle: Hadoop
  - Group and sort alignments by region
- Reduce: SOAPsnp (Li *et al.*, 2009)
  - Scan alignments for divergent columns
  - Accounts for sequencing error, known SNPs



Performance in Amazon EC2

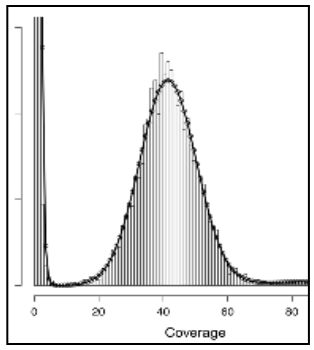
http://bowtie-bio.sourceforge.net/crossbow

	Asian Individual Genome		
Data Loading	3.3 B reads	106.5 GB	$10.65
Data Transfer	1h :15m	40 cores	$3.40
Setup	0h : 15m	320 cores	$13.94
Alignment	1h : 30m	320 cores	$41.82
Variant Calling	1h : 00m	320 cores	$27.88
End-to-end	4h : 00m		$97.69

Analyze an entire human genome for ~$100 in an afternoon.
Accuracy validated at >99%

Searching for SNPs with Cloud Computing.
Langmead B, Schatz MC, Lin J, Pop M, Salzberg SL (2009) *Genome Biology*. **10**:R134

Hadoop for NGS Analysis



Quake

Quality-aware error correction of short reads

Correct 97.9% of errors with 99.9% accuracy

http://www.cbcb.umd.edu/software/quake/

(Kelley, Schatz, Salzberg, 2010*)

CloudBurst

Highly Sensitive Short Read Mapping with MapReduce

100x speedup mapping on 96 cores @ Amazon

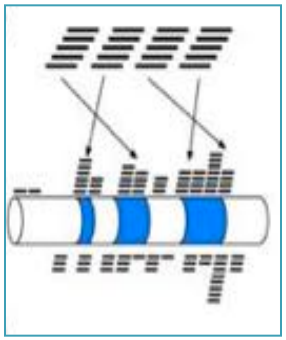
http://cloudburst-bio.sf.net

(Schatz, 2009)

Myrna

Cloud-scale differential gene expression for RNA-seq

Expression of 1.1 billion RNA-Seq reads in ~2 hours for ~$66



(Langmead, Hansen, Leek, 2010)

http://bowtie-bio.sf.net/myrna/

AMOS

Searching for SNPs in the Turkey Genome

Scan the de novo assembly to find 920k hetrozygous alleles

(Dalloul et al, 2010)

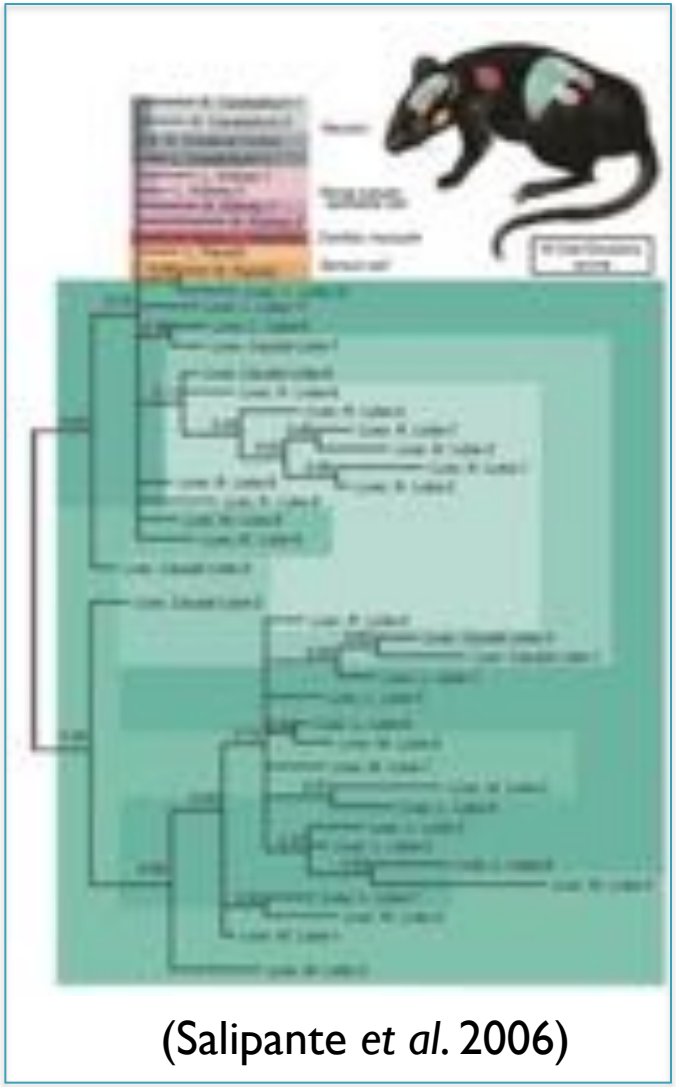
http://amos.sf.net

SeqMS: NextGen Microsatellite Profiling

Mitchell Bekritsky, WSBS

- Class of simple sequence repeats
  - …GCACACACACAT… = …G(CA)$_5$T…
  - Created and mutate primarily through slippage during replication
  - Highly variable & ubiquitous

- Genotyping with SeqMS
  - Rapidly detect MS sequences
  - Map reads using a new MS-mapper
  - Analyze profiles in cells, across cells, & across populations
    - Loss of heterozygosity
    - Development of somatic & cancer cells
    - Relations across strains, across species
    - etc…

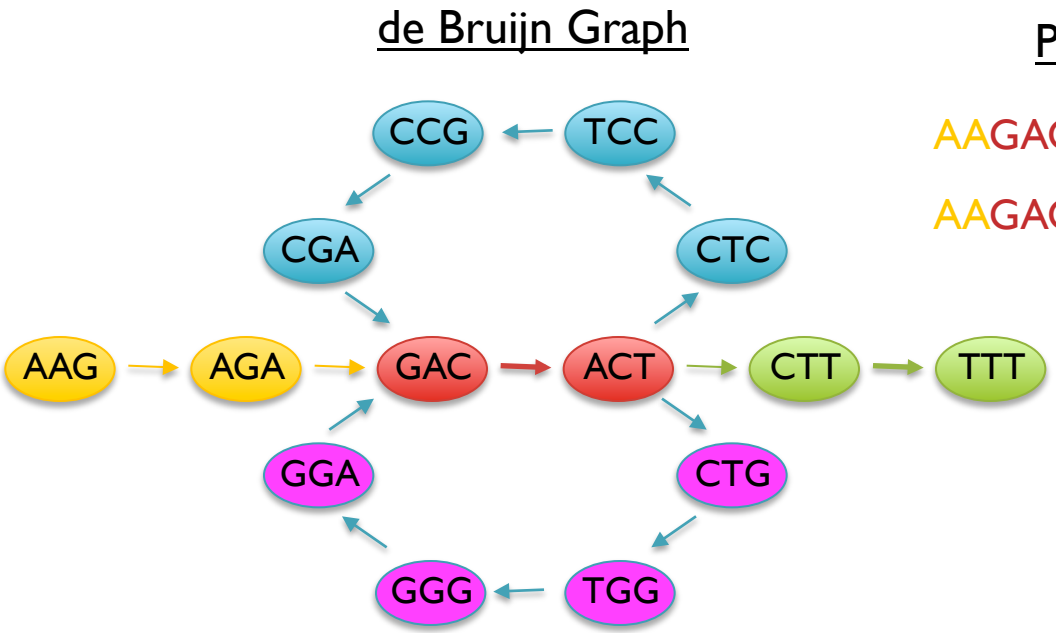


(Salipante *et al.* 2006)

Short Read Assembly

Reads

AAGA
ACTT
ACTC
ACTG
AGAC
CCGA
CGAC
CTCC
CTGG
CTTT
…



Potential Genomes

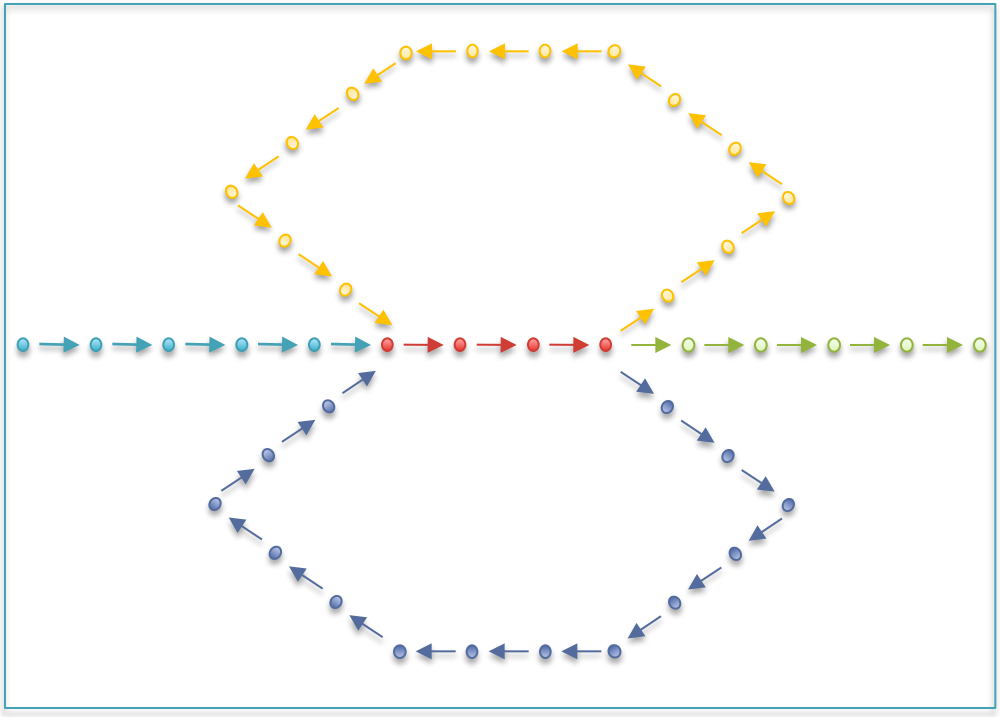
AAGACTCCGACTGGGACTTT

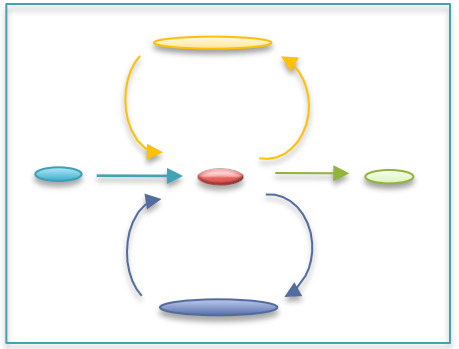
AAGACTGGGACTCCGACTTT

- Genome assembly as finding an Eulerian tour of the de Bruijn graph
  - Human genome: >3B nodes, >10B edges
- The new short read assemblers require tremendous computation
  - Velvet (Zerbino & Birney, 2008) serial: > 2TB of RAM
  - ABySS (Simpson *et al.*, 2009) MPI: 168 cores x ~96 hours
  - SOAPdenovo (Li *et al.*, 2010) pthreads: 40 cores x 40 hours, >140 GB RAM

Graph Compression

- After construction, many edges are unambiguous
  - Merge together compressible nodes
  - Graph randomly distributed over hundreds of computers





Design Patterns for Efficient Graph Algorithms in MapReduce.
Lin, J, Schatz, MC (2010) *Workshop on Mining and Learning with Graphs Workshop (KDD/MLG-2010)*

Fast Path Compression

Challenges

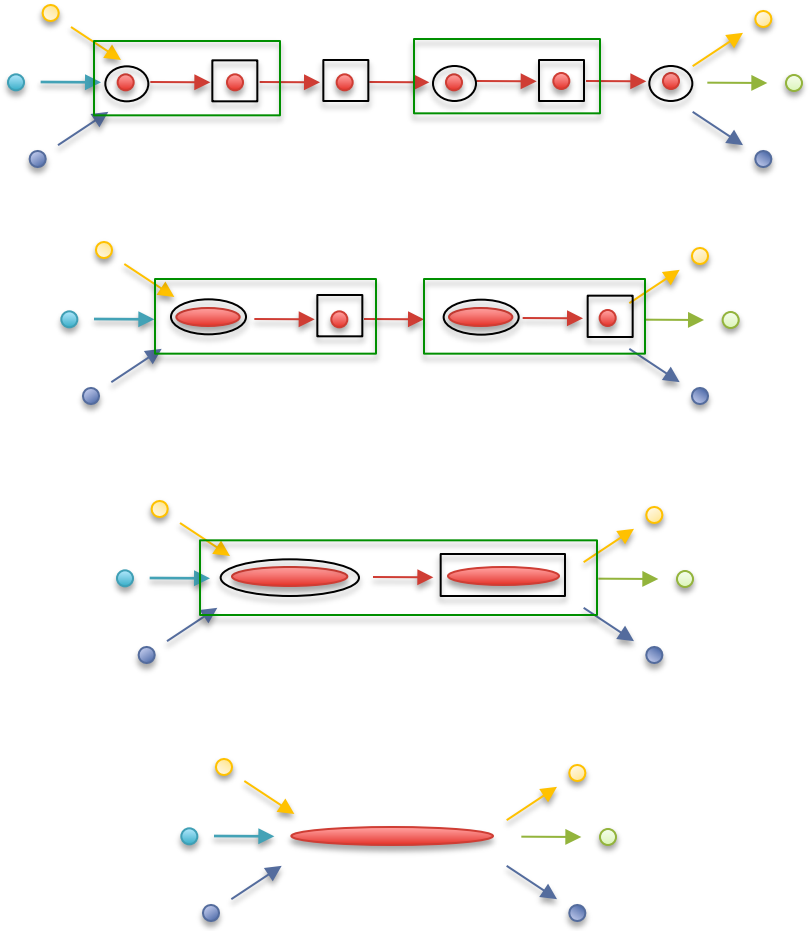
- Nodes stored on different computers
- Nodes can only access direct neighbors

Randomized List Ranking

- Randomly assign Ⓗ / [T] to each compressible node
- Compress Ⓗ → [T] links

Performance

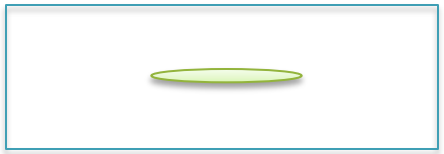
- Compress all chains in log(S) rounds
- <30 rounds for human genome



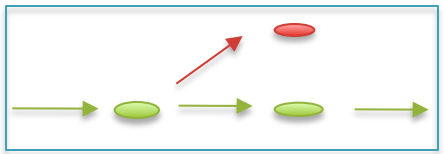
Randomized Speed-ups in Parallel Computation.
Vishkin U. (1984) *ACM Symposium on Theory of Computation. 230-239.*

Node Types

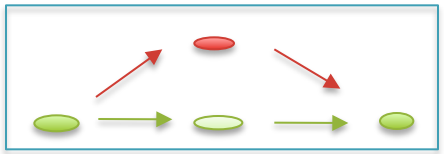
(Chaisson, 2009)



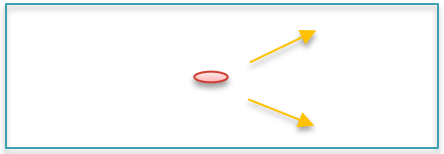
Isolated nodes (10%)



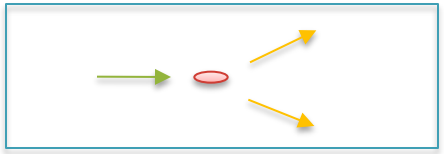
Tips (46%)



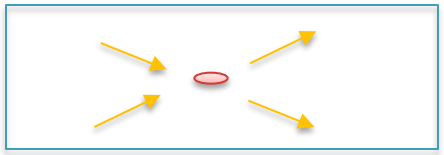
Bubbles/Non-branch (9%)



Dead Ends (.2%)



Half Branch (25%)



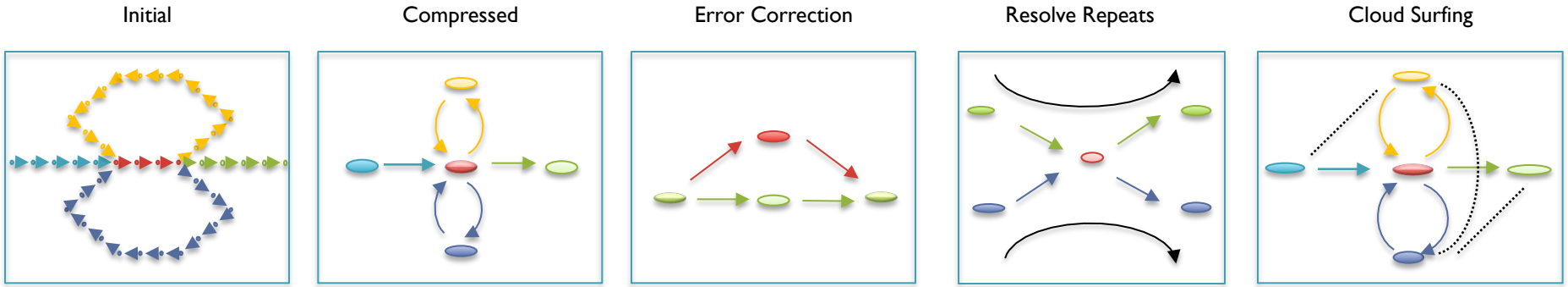
Full Branch (10%)

Contrail

http://contrail-bio.sourceforge.net

De novo bacterial assembly

- *Genome: E. coli* K12 MG1655, 4.6Mbp
- *Input:* 20.8M 36bp reads, 200bp insert (~150x coverage)
- *Preprocessor*: Quake Error Correction



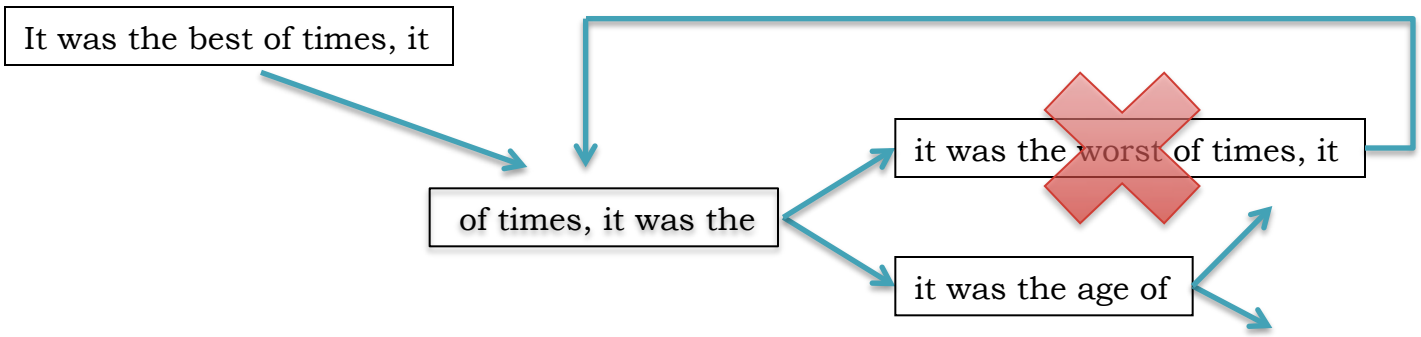
N	5.1 M	245,131	2,769	1,909	300
Max	27 bp	1,079 bp	70,725 bp	90,088 bp	149,006 bp
N50	27 bp	156 bp	15,023 bp	20,062 bp	54,807 bp

Assembly of Large Genomes with Cloud Computing.
Schatz MC, Sommer D, Kelley D, Pop M, *et al. In Preparation.*

E. coli Assembly Quality

Incorrect contigs: Align at < 95% identity or < 95% of their length

Assembler	Contigs ≥ 100bp	N50 (bp)	Incorrect contigs
Contrail PE	300	54,807	4
Contrail SE	529	20,062	0
SOAPdenovo PE	182	89,000	5
ABySS PE	233	45,362	13
Velvet PE	286	54,459	9
EULER-SR PE	216	57,497	26
SSAKE SE	931	11,450	38
Edena SE	680	16,430	6

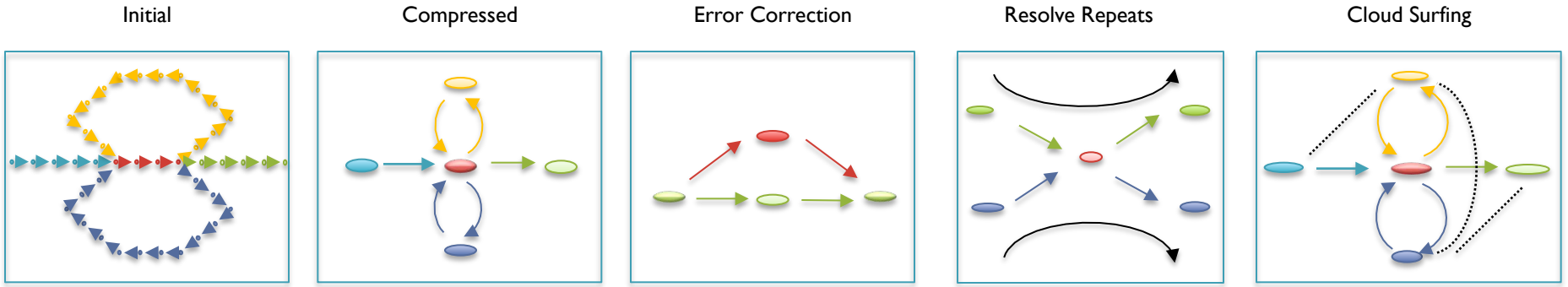


Contrail

http://contrail-bio.sourceforge.net

De novo assembly of the Human Genome

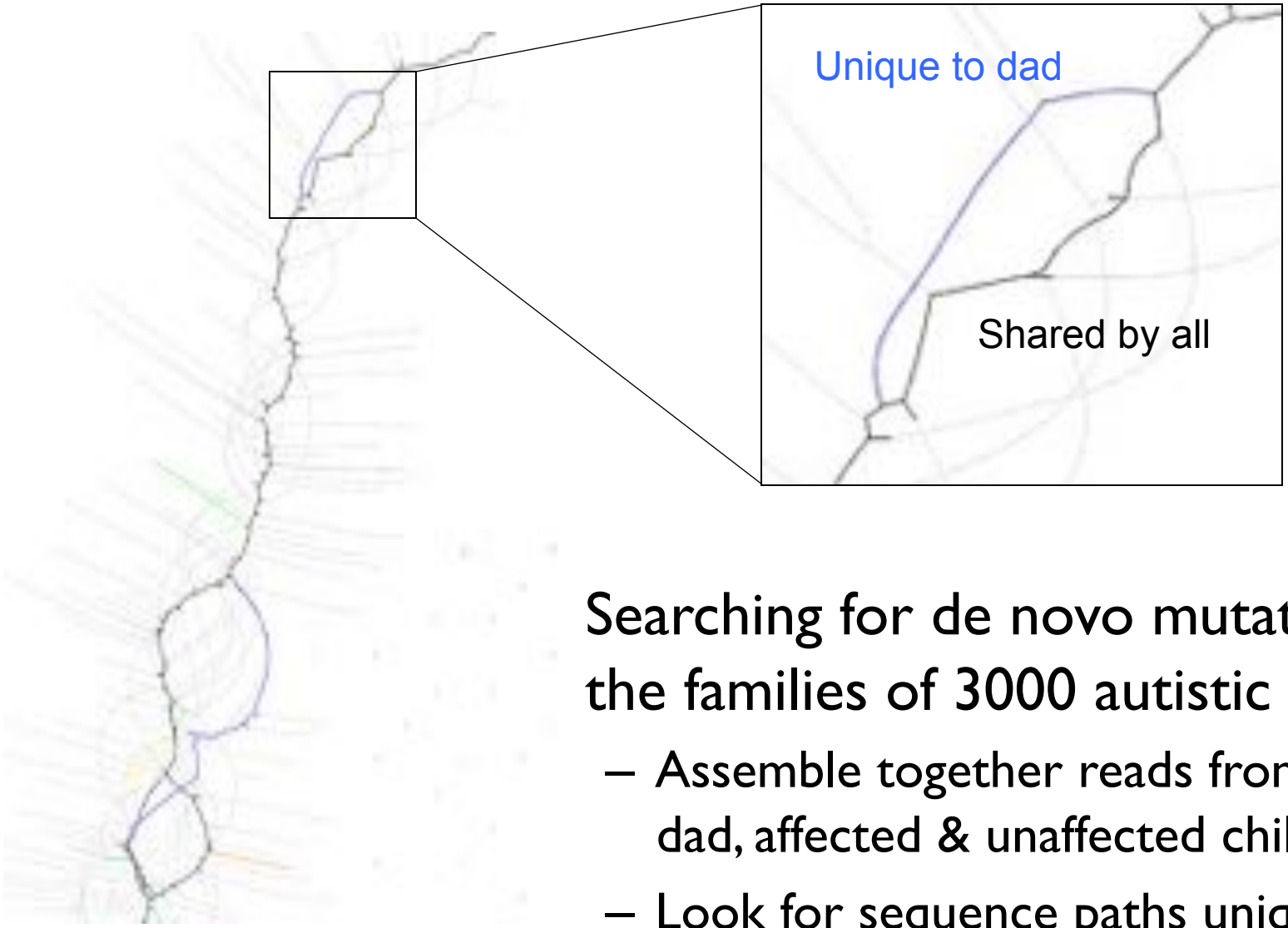
- *Genome:* African male NA18507 (SRA000271, Bentley *et al.*, 2008)
- *Input: 3.5*B 36bp reads, 210bp insert (~40x coverage)



N	>7 B	>1 B	4.2 M	4.1 M	3.3 M
Max	27 bp	303 bp	20,594 bp	20,594 bp	20,594 bp
N50	27 bp	< 100 bp	995 bp	1,050 bp	1,427 bp*

Assembly of Large Genomes with Cloud Computing.
Schatz MC, Sommer D, Kelley D, Pop M, *et al. In Preparation.*

Variations and de Bruijn Graphs



Searching for de novo mutations in the families of 3000 autistic children.

- Assemble together reads from mom, dad, affected & unaffected children
- Look for sequence paths unique to affected child

Summary

- Staying afloat in the data deluge means computing in parallel
  - Hadoop + Cloud computing is an attractive platform for large scale sequence analysis and computation
- Significant obstacles ahead
  - Price
  - Transfer time
  - Privacy / security requirements
  - Time and expertise required for development

(Schatz *et al.*, Nature Biotechnology, 2010)

- Emerging technologies are a great start, but we need continued research
  - A word of caution: new technologies are new

Acknowledgements

CSHL

Mike Wigler
Ivan Iossifov
Mike Ronemus
Jude Kendall
Dan Levy

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Dick McCombie
Doreen Ware

Mitch Bekritsky

Matt Titmus

Univ. of Maryland

Steven Salzberg
Mihai Pop
Carl Kingsford
Art Delcher
Jimmy Lin
Dan Sommer
David Kelley

JHU

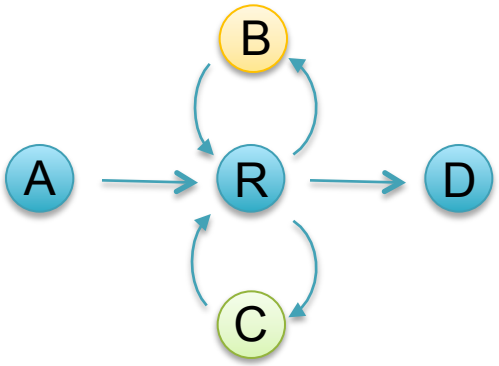
Ben Langmead

Thank You!

http://schatzlab.cshl.edu

@mike_schatz

Counting Eulerian Tours



ARBRCRD
or
ARCRBRD

Often an astronomical number of possible assemblies

– Value computed by application of the BEST theorem (Hutchinson, 1975)

$$\mathcal{W}(G,t) = (\det L)\left\{\prod_{u\in V}(r_u-1)!\right\}\left\{\prod_{(u,v)\in E} a_{uv}!\right\}^{-1}$$

L = n x n matrix with r_u-a_{uu} along the diagonal and -a_{uv} in entry uv

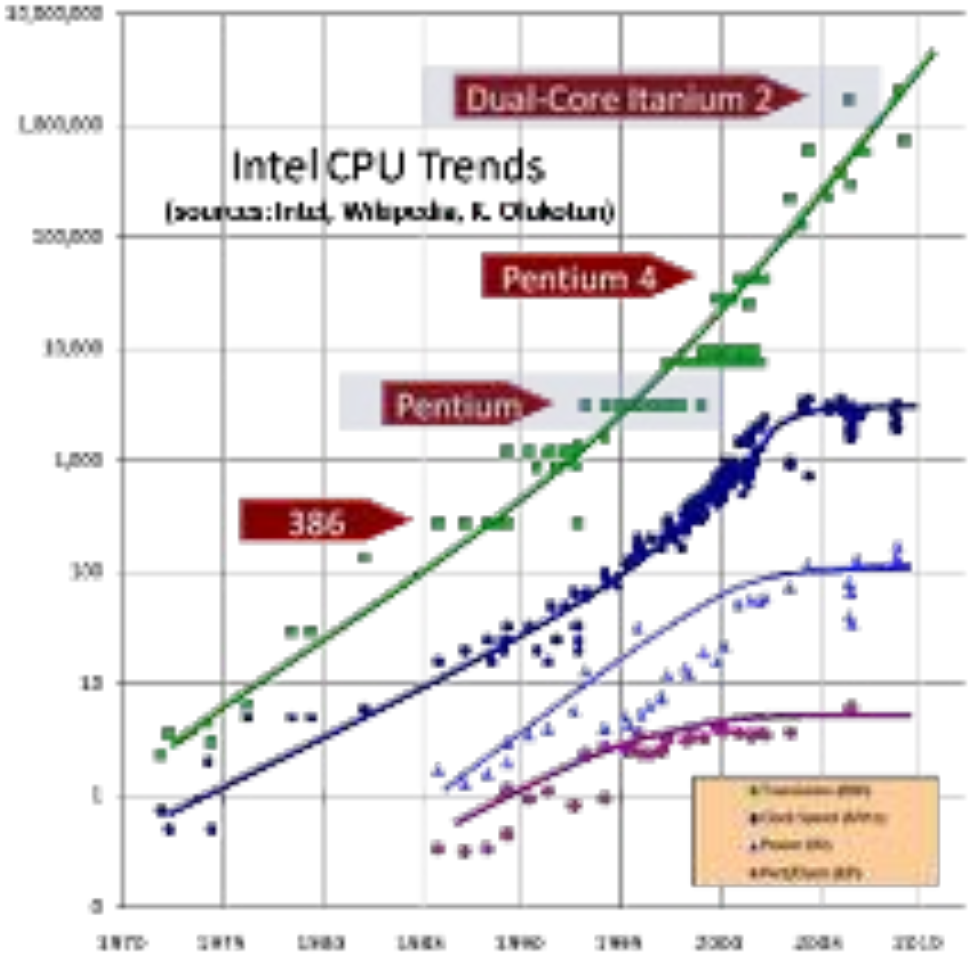
$r_u = d^+(u)+1$ if $u=t$, or $d^+(u)$ otherwise

a_{uv} = multiplicity of edge from u to v

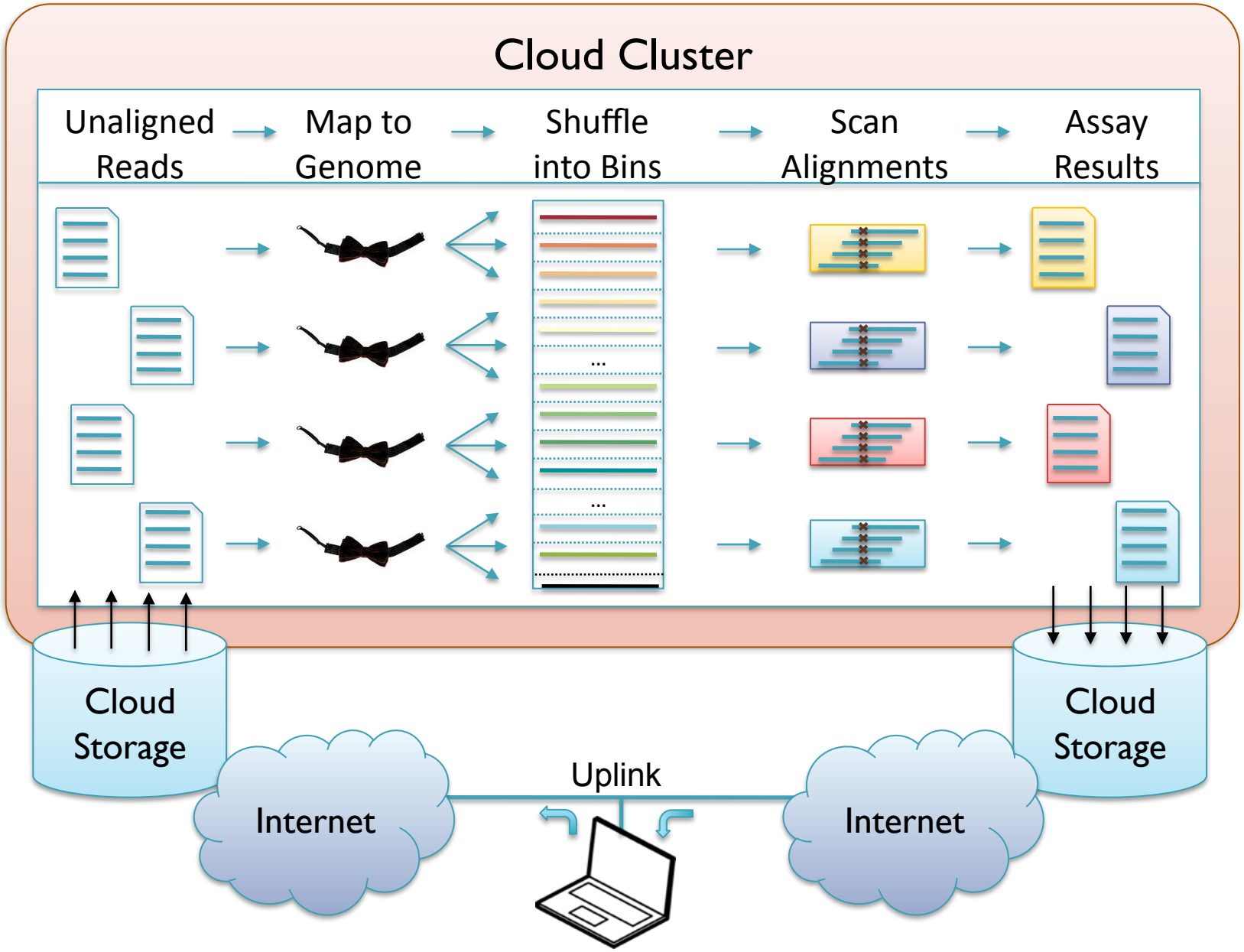
Assembly Complexity of Prokaryotic Genomes using Short Reads.
Kingsford C, Schatz MC, Pop M (2010) *BMC Bioinformatics*.

Why HPC?

- Moore's Law is valid in 2010
  - But CPU speed is flat
  - Vendors adopting parallel solutions instead
- Parallel Environments
  - Many cores, including GPUs
  - Many computers
  - Many disks
- Why parallel
  - Need results faster
  - Doesn't fit on one machine



The Free Lunch Is Over: A Fundamental Turn Toward Concurrency in Software
Herb Sutter, http://www.gotw.ca/publications/concurrency-ddj.htm



Cloud Computing and the DNA Data Race.
Schatz, MC, Langmead, B, Salzberg SL (2010) *Nature Biotechnology*. 28: 691–693