



Bioinformatics and its applications

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Term Bioinformatics

Term **Bioinformatics** was invented by Paulien Hogeweg (Полина Хогевег) and Ben Hesper in 1970 as "the study of informatic processes in biotic systems".

Paulien Hogeweg is a Dutch theoretical biologist and complex systems researcher studying biological systems as dynamic information processing systems at many interconnected levels.

Definitions of what is Bioinformatics:

Bioinformatics is the use of IT in biotechnology for the data storage, data warehousing and analyzing the DNA sequences. In Bioinfomatics knowledge of many branches are required like biology, mathematics, computer science, laws of physics & chemistry, and of course sound knowledge of IT to analyze biotech

data. Bioinform Bioinformatics is an interdisciplinary field that develops and improves upon used to solve n methods for storing, retrieving, organizing and analyzing biological data. A major activity in bioinformatics is to develop software tools to generate useful

The mathematical, statistical and computing methods that

aim to solve biological problems using DNA and amino acid

Bioinformatics develop of computer to analy biology for example ingredients and metable

sequences and related informal Bioinformatics: Research, development, or application of computational tools and approaches for expanding the use of biological, medical, behavioral or health data, including those to acquire, store, organize, archive, analyze, or visualize such data.

http://www.bisti.nih.gov/CompuBioDef.pdf

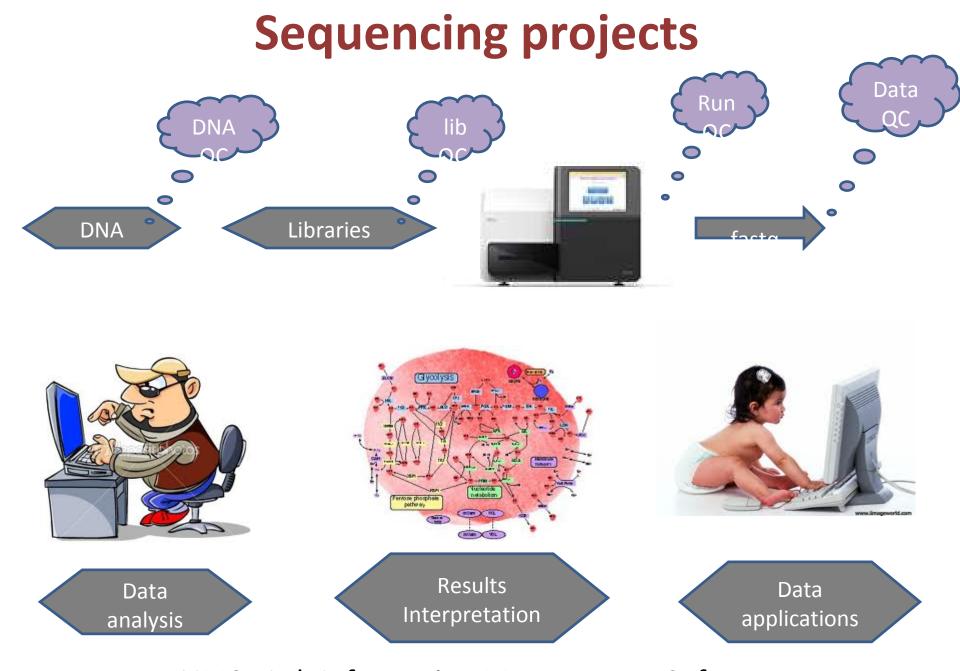
My additions:

- 1. Bioinformatics is a SCIENCE
- 2. Not only to develop algorithms, store, retrieve, organize and analyze biological data but to CURATE data

Bioinformatics is being used in following fields:

- Microbial genome applications
- Molecular medicine
- Personalised medicine
- **→** Preventative medicine
- Gene therapy
- Drug development
- Antibiotic resistance
- Evolutionary studies
- → Waste cleanup
- Biotechnology

- **→** Climate change Studies
- **→** Alternative energy sources
- Crop improvement
- **→** Forensic analysis
- **→** Bio-weapon creation
- **→** Insect resistance
- Improve nutritional quality
- Development of Drought resistant varieties
- **→** Vetinary Science



LIMS - Lab Information Management Software

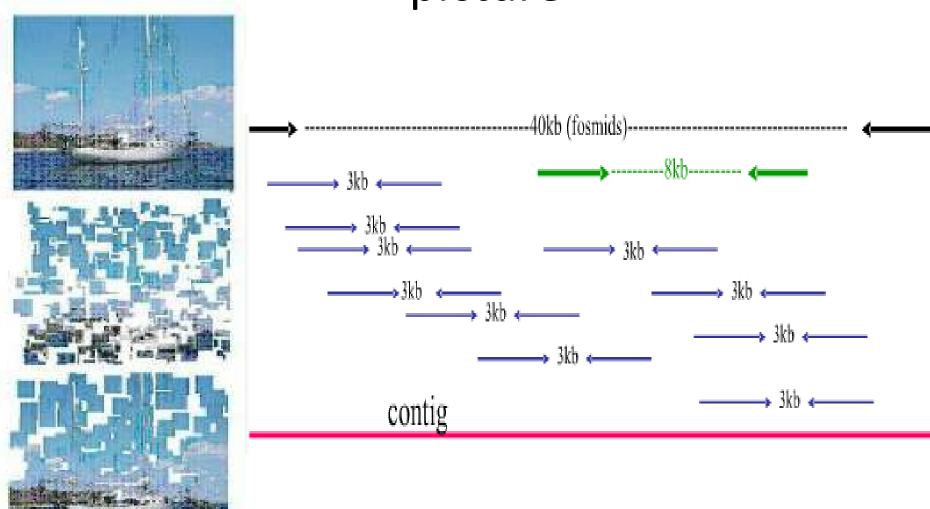
Microbial genome applications

- Genome assembly
- → Re-sequencing
- Comparative analysis
- Evolutionary studies
- Antibiotic resistance
- Waste cleanup
- Biotechnology

Genome Assembly

- Genome assembly is a very complex computational problem due to enormous amount of data to put together and some other reasons reasons.
- Ideally an assembly program should produce one contig for every chromosome of the genome being sequenced. But because of the complex nature of the genomes, the ideal conditions just never possible, thus leading to gaps in the genome.

De Novo assembly - puzzle without the picture

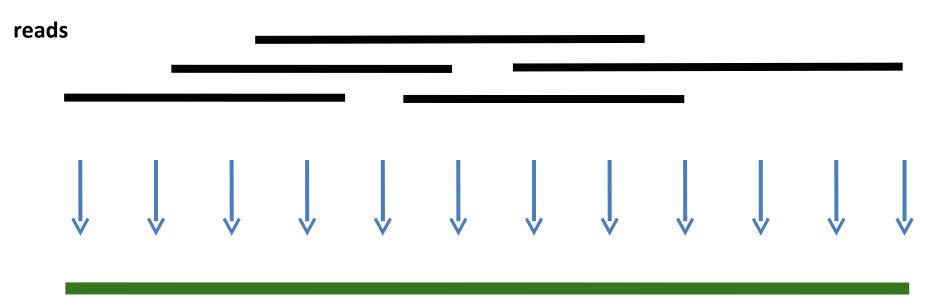


Assembly Challenges

- Presence of **repeats**. Repeats are identical sequences that occur in the genome in different locations and are often seen in varying lengths and in the multiple copies. There are several types of repeats: tandem repeats or interspersed repeats. The read's originating from different copies of the repeat appear identical to the assembler, causing errors in the assembly.
- Contaminants in samples (eg. from Bacteria or Human).
- PCR artefacts (eg. Chimeras and Mutations)
- **Sequencing errors**, such as "*Homopolymer*" errors when eg. 2+ run of same base.
- MID's (multiplex indexes), primers/adapters still in the raw reads.
- polyploid genomes

Assembly algorithms

Overlap-Layout-Consensus - Find overlaps between all reads



Consensus

Problems caused by new sequencing technologies:

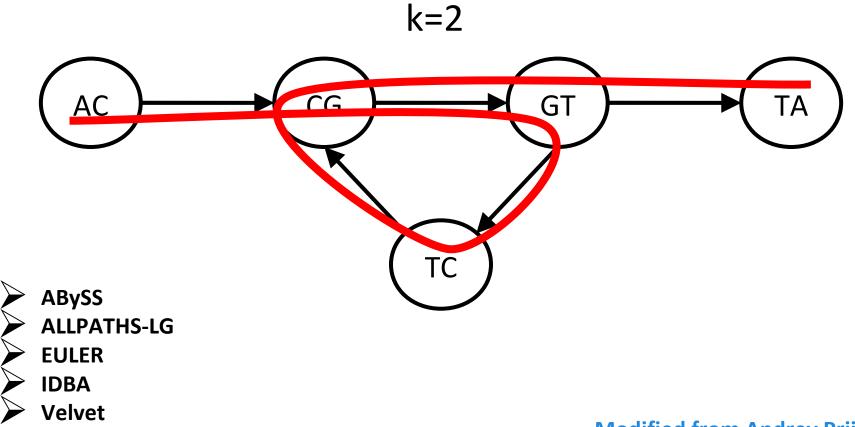
Hard to find overlaps between short reads



Impossible to scale up

De Bruijn graph

ACGTCGTA



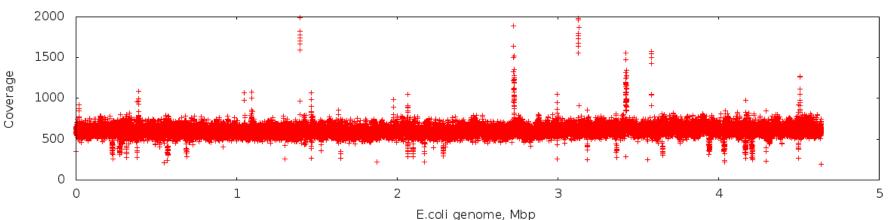
Single-cell dataset

• E. coli isolate dataset

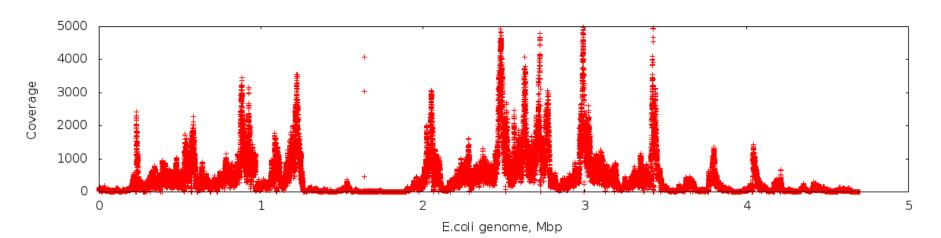
- IDBA-UD
- SPAdes





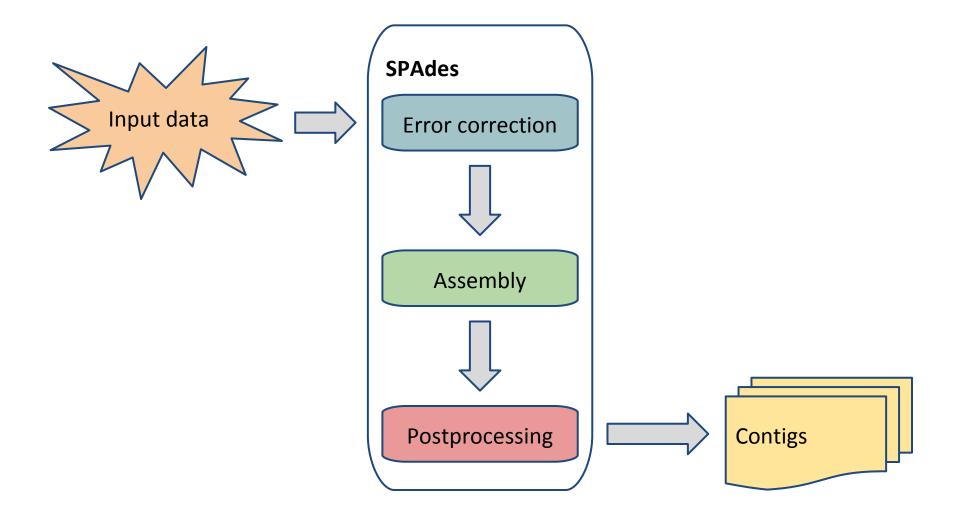


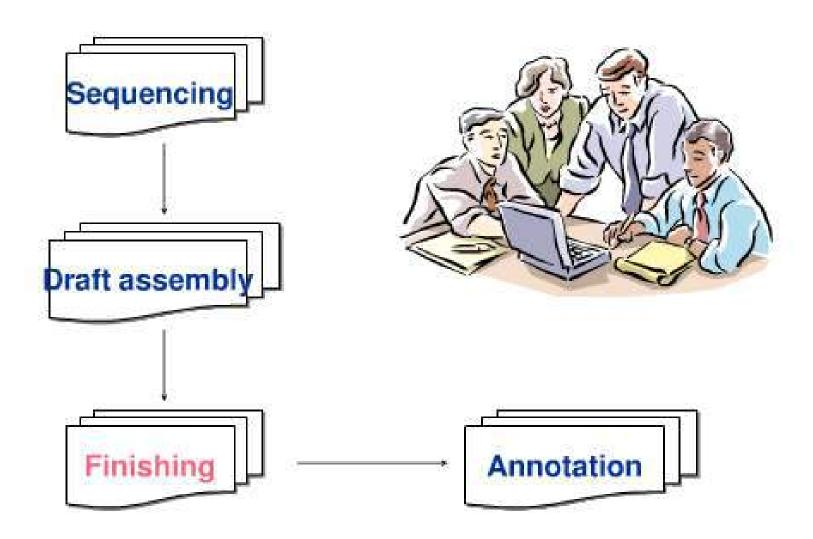
• E.coli single-cell dataset



SPAdes pipeline







Gene Prediction and Genome Annotation



<u>Based on similarity to</u> <u>known genes</u> – blastX (NCBI)

Gene finding programs

- Glimmer for most procaryotic genomes
- GenMark for both procaryotic genomes and eucaryotic genomes



INTEGRATED MICROBIAL GENOMES

IMG Home

Find Genomes

Find Genes

Find Functions

Compare Genomes

Analysis Cart

My IMG

Companion Systems

Using IMG

IMG Content

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	Datasets
Bacteria	6120
Archaea	248
<u>Eukarya</u>	<u>183</u>
Plasmids	1193
Viruses	2809

Genome Fragments 579

Total Datasets 11132

GEBA <u>245</u> Last updated: 2013-07-05

IMG 4.0 is dedicated to the memory of our colleague, lain Anderson

Genome by Metadata Project Map Content History System Requirements About IMG FAQ



Hands on training available at

Microbial Genomics & Metagenomics Workshop The Integrated Microbial Genomes (IMG) system (Nucleic Acids Research, Vol 40, 2012) serves as a community resource for comparative analysis and annotation of all publicly available genomes from three domains of life in a uniquely integrated context. Plasmids that are not part of a specific microbial genome sequencing project and phage genomes are also included into IMG in order to increase its genomic context for comparative analysis.

For details, see <u>IMG Release Notes</u> (Dec. 12, 2012), in particular the workspace and background computation capabilities available to IMG registered users.

Count	Total		
DNA, number of bases	60,476,662,654		
Total Genes	25,395,838		
Total Genomes	11,132		



IMG Statistics

IMG ER Account Request

All Genomes



Genome Count

Status	Bacteria	Archaea	Eukaryota	Plasmids	Viruses	Genome Fragments	Total
Finished	<u>2131</u>	<u>154</u>	<u>37</u>	<u>1190</u>	2809	<u>579</u>	6900
Draft	2407	<u>28</u>	<u>146</u>	<u>3</u>	0	0	<u>2584</u>
Permanent Draft	<u>1582</u>	<u>66</u>	0	0	0	0	<u>1648</u>
Total	<u>6120</u>	<u>248</u>	<u>183</u>	<u>1193</u>	2809	<u>579</u>	11132

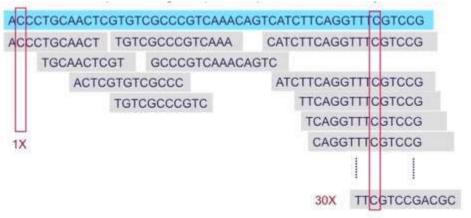
Re-sequencing

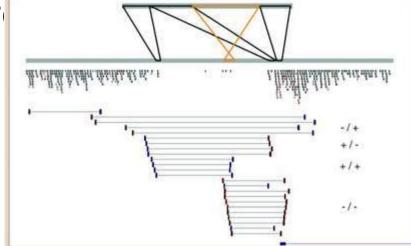
Projects aimed at characterizing the genetic variations of species or populations

Resequencing of bacterial and archaeal isolates etc is possible if reference genomes are available

This approach can help to better understand bacterial community structure, gene function in

bacteria under selective pr





Climate change Studies

Increasing levels of carbon dioxide emission are thought to contribute to global climate change.

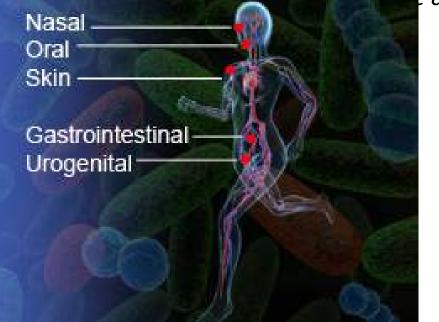
One way to decrease atmospheric carbon dioxide is to study the genomes of microbes that use carbone dioxidet as their sole carbon source

Human microbiome

MetaHIT - Europe Human Microbiome Project –US

The human microbiome includes viruses, fungi and bacteria, their genes and their environmental interactions, and is known to influence human physiology.

There's very broad variation in these bacteria in different people and that severely limits our ability to create a "normal" microflora profile for comparison among healthy people and those with any kind of health issues.



Children with **autism** harbor significantly fewer types of gut bacteria than those who are not affected by the disorder, researchers have found.

Prevotella species were most dramatically reduced among samples from autistic children—especially *P. copri*. (helps the breakdown of protein and carbohydrate foods)

Bioinformatics combining biology with computer science

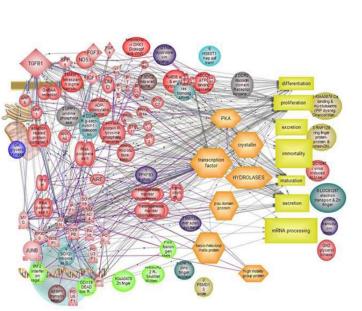
- it can explore the causes of diseases at the molecular level
- explain the phenomena of the diseases on the gene/pathway level
- make use of computer techniques (data mining, machine learning etc), to analyze and interpret data faster
- to enhance the accuracy of the results





Reduce the cost and time of drug discovery

To improve drug discovery we need to discover (read "develop") efficient bioinformatics algorithms and approaches for

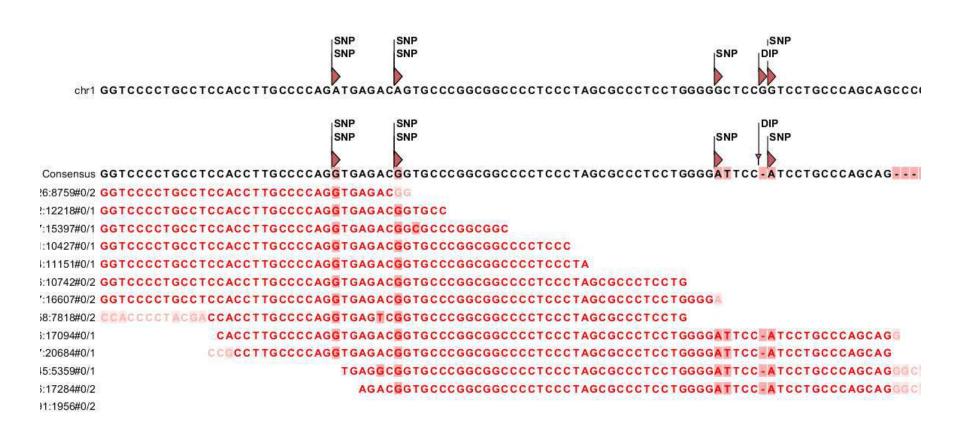


target identification
target validation
lead identification
lead optimization

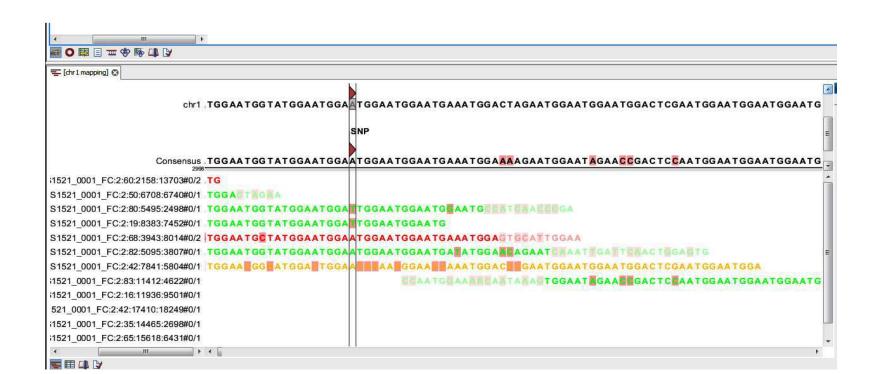
Advantages of detecting mutations with next-generation sequencing

- High throughput
 - Test many genes at once
- Systematic, unbiased mutation detection
 - All mutation types
 - Single nucleotide variants (SNV), copy number variation (CNV)-insertions, deletions and translocations
- Digital readout of mutation frequency
 - Easier to detect and quantify mutations in a heterogeneous sample
- Cost effective **precision** medicine
 - "Right drug at right dose to the right patient at the right time"

Homozygous SNPs and indel



Poor alignment



Missed SNP?

chrGCAGAGGCCAAGCCAGAGGTTCCAGGCTTAAACCCCCAGCCCTGCCCTGCCCAGTCCA

ConsensuGCAGAGGCCAAGCCAGAGGTTCCAGGCTTAAACCCCAGCCCTGCCCTGCCCAGTCCA

3866:4795#0/GCA

1407:2153#0/

1308:3912#0/GCAGAGGCCAAGCCAGAGGTTCCAGGCTTAAA

3914:7870#0/: GCCAAGCCAGAGGTTCCAGGCTTAAACCCCCAGCCCTGCCCTGCCCAGTCCA

5555:2114#0/ CAAGCCAGAGGTTCCAGGCTTAAACCCCCAGCCCTGCCCAGTCCA

579:16341#0/ CCAGGCTTAACCCCCAGCCCTGCCCAGTCCA

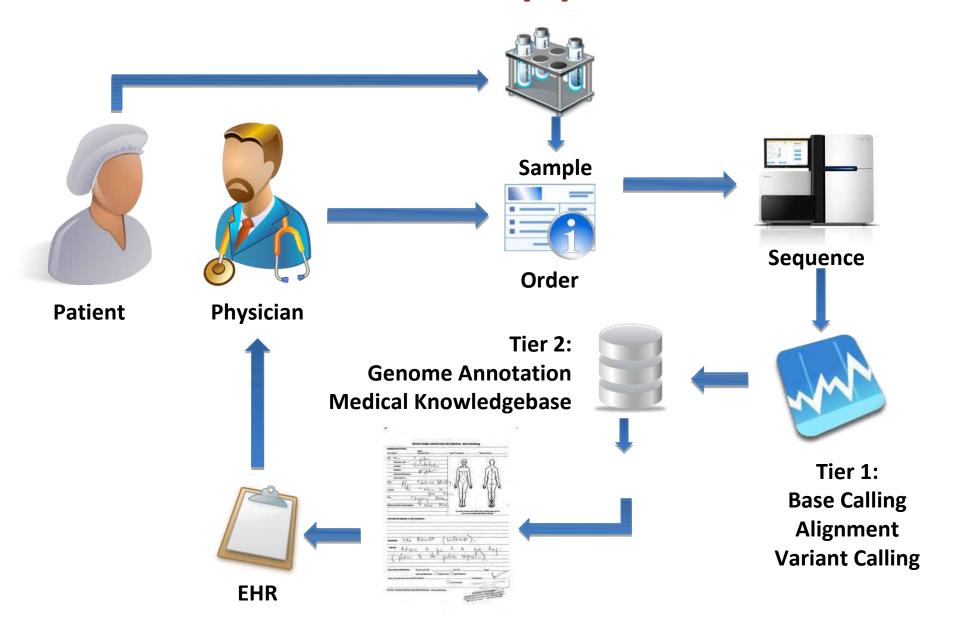
3944:9734#0/: GGCTTAACCCCCAGCCCTGCCCAGTCCA

108:13945#0/

Bioinformatics and Health Informatics

If bioinformatics is the study of the flow of information in biological sciences, Health Informatics is the study of the information in patient care

Medicine: Informatics pipeline workflow



Tier 3: Clinical Report

Huge need in bioinformatics tools

Simple pipelines/protocols and easy to read reports



Sample sequencing



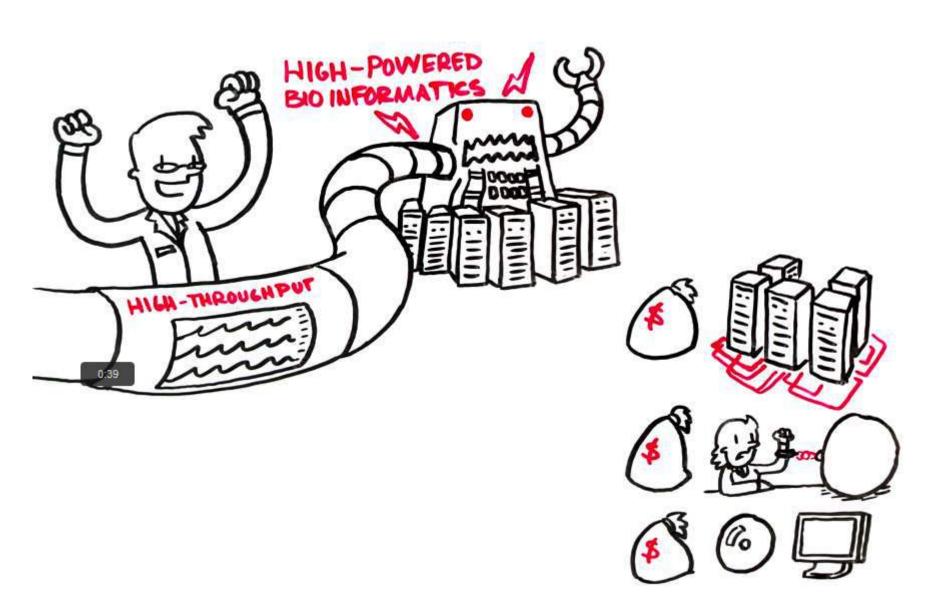
Data Analysis



Patients treatment

Team work to set up cancer sequencing





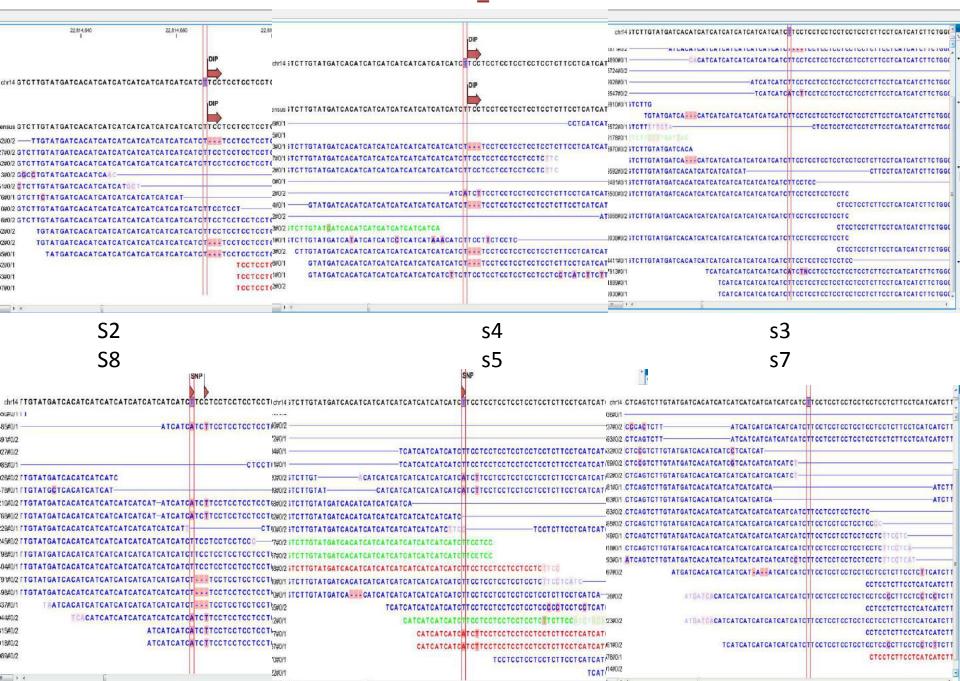
Ion Torrent: Torrent Suite Software





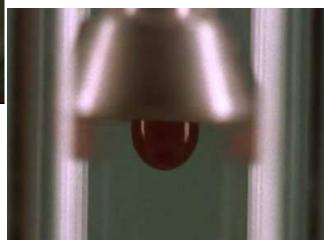


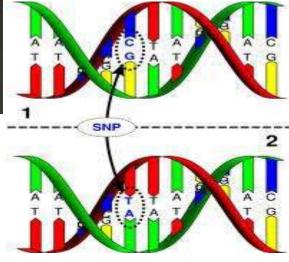
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Each baby to be sequenced at birth: personal reference

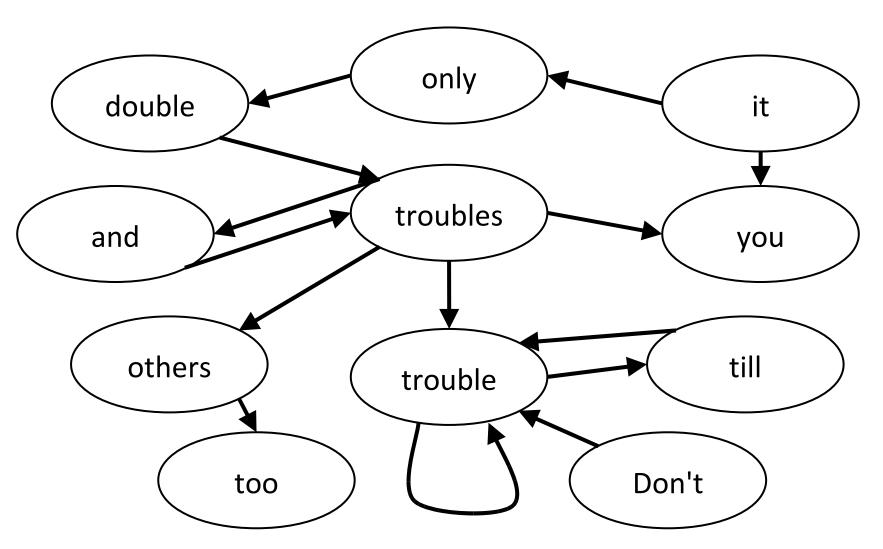






Funny De Bruijn graph









THANK YOU!