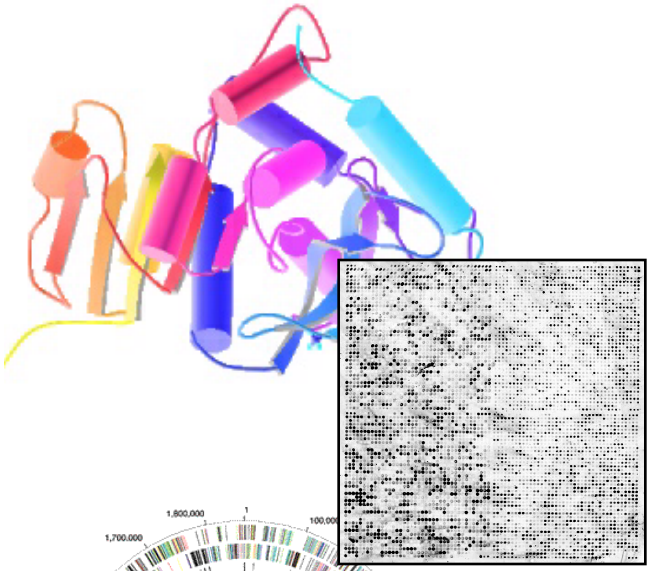
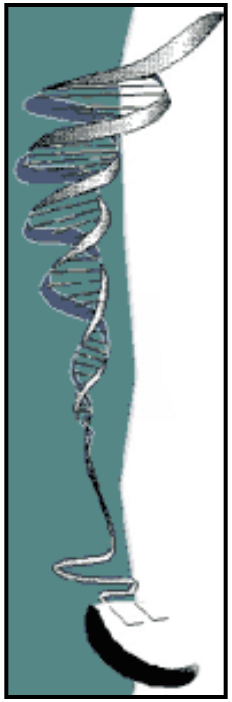
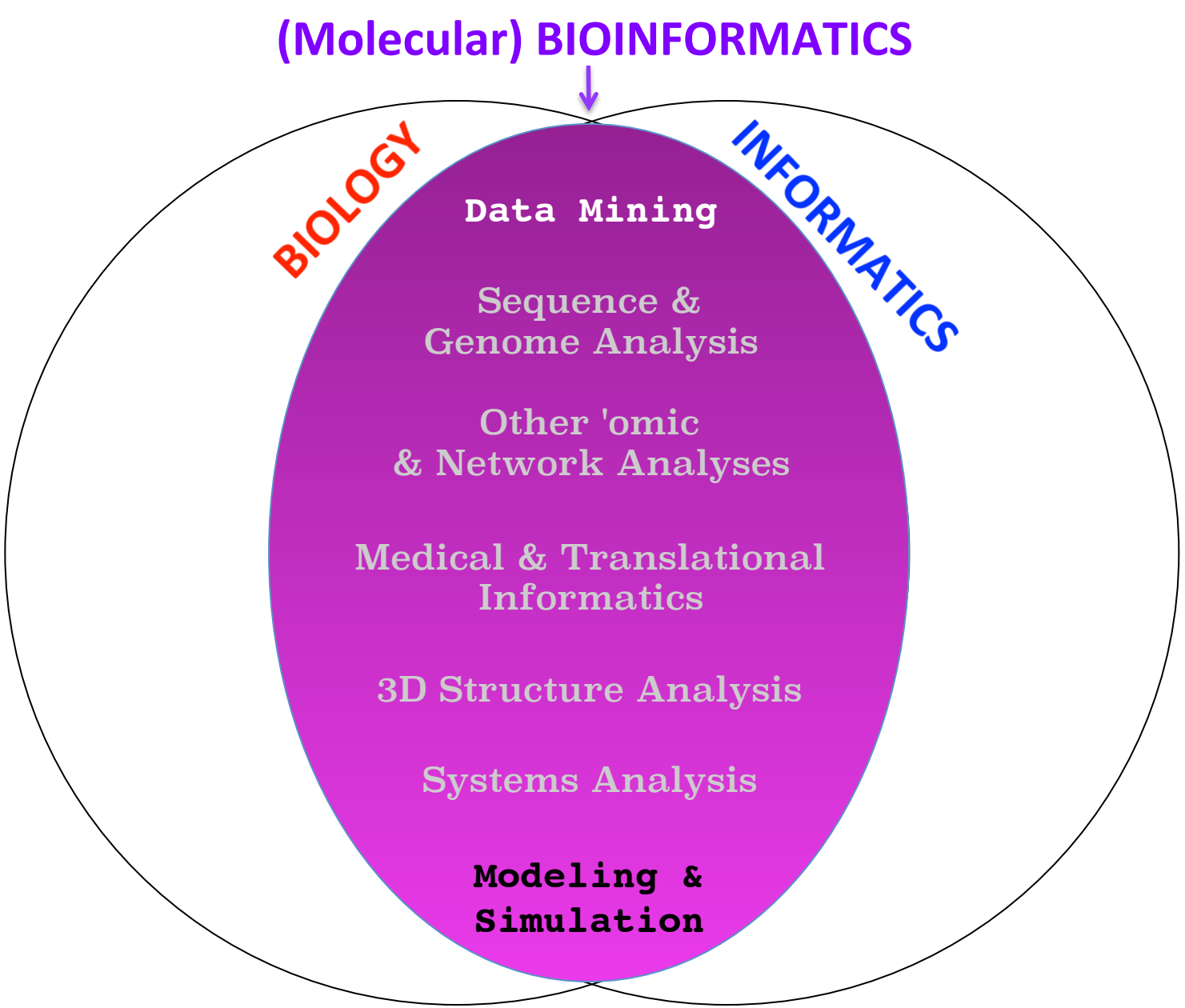
BIOINFORMATICS
Introduction



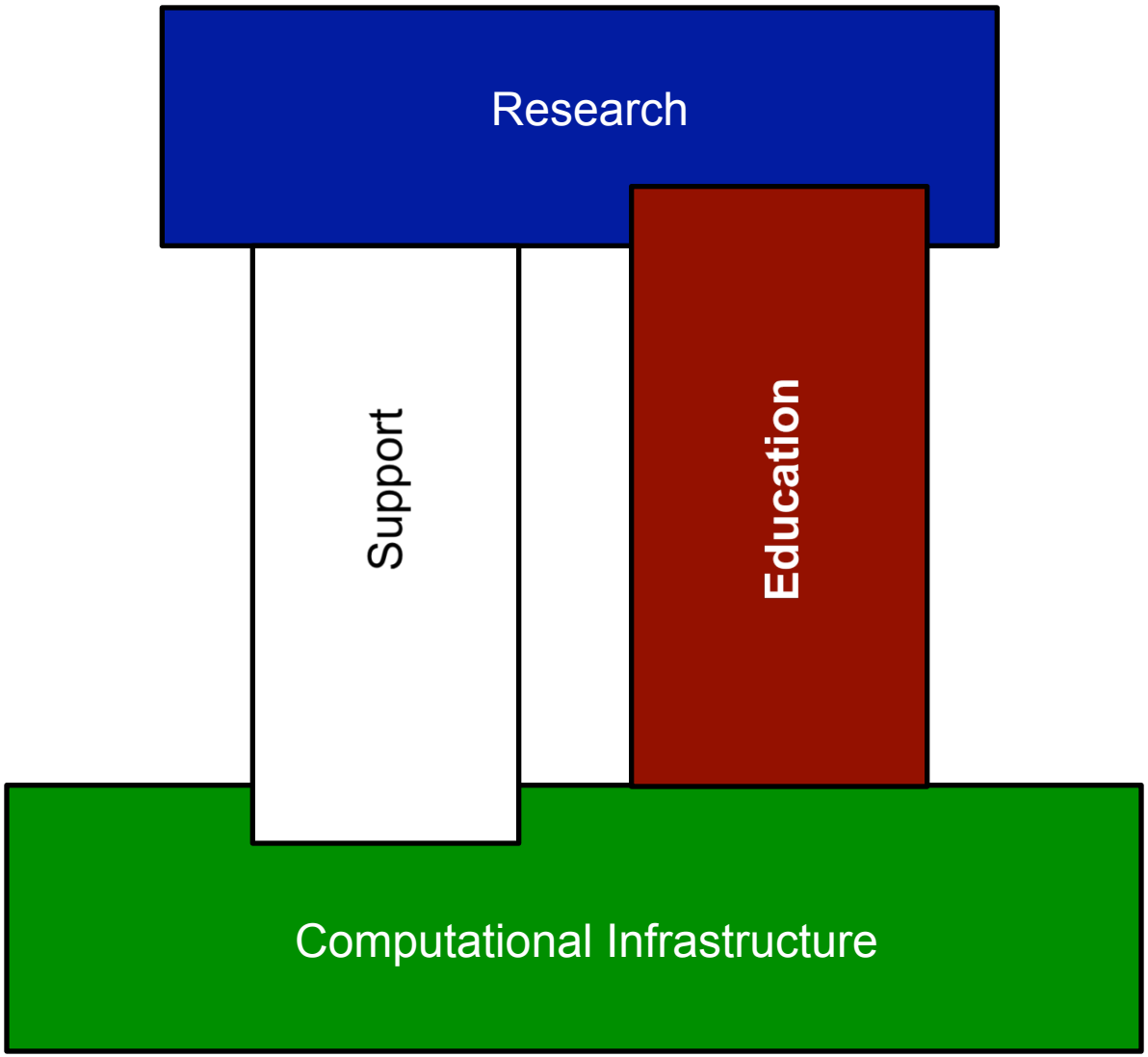


Mark Gerstein, Yale University
GersteinLab.org/courses/452
(last edit in spring ’15)



[Luscombe et al. ('01). *Methods Inf Med* 40: 346]

Elements of Bioinformatics as a discipline



What is Bioinformatics?

- *(Molecular)* **Bio - `informatics`**
- One idea for a definition?
  Bioinformatics is conceptualizing **biology in terms of molecules** (in the sense of physical-chemistry) and then applying **"informatics" techniques** (derived from disciplines such as applied math, CS, and statistics) to **organize, mine, model & understand the information associated** with these molecules, **on a large-scale.**
- Bioinformatics is a practical discipline with many **applications**.

[Luscombe et al. ('01). *Methods Inf Med* 40: 346]

What **Information** to Organize?

- **Sequences** (DNA & Protein)
- 3D Structures
- Network & Pathway Connectivity
- Phylogenetic tree relationships
- Large-scale gene expression & functional genomics data
- Phenotypic data & medical records....

What is the **Information?**

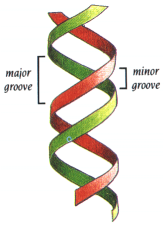
Molecular Biology as an Information Science

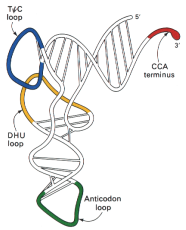
- Central Dogma of Molecular Biology

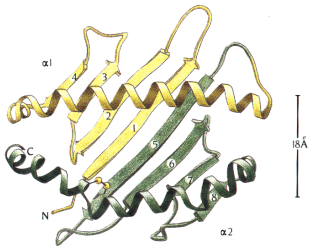
```
DNA
 -> RNA
  -> Protein
   -> Phenotype
    -> DNA
```

- Central Paradigm for Bioinformatics

```
Genomic Sequence Information
 -> mRNA (level)
  -> Protein Sequence
   -> Protein Structure
    -> Biological Function
     -> Organismal Phenotype
```







•Genetic material

•Information transfer (mRNA)
•Protein synthesis (tRNA/mRNA)
•Some catalytic activity

Molecular Biology Information - DNA

- Raw DNA Sequence
  - ◊ Coding or Not?
  - ◊ Parse into genes?
  - ◊ 4 bases: AGCT
  - ◊ ~1 K in a gene, ~2 M in genome
  - ◊ ~3 Gb Human

```
atggcaattaaaattggtatcaatggttttggtcgtatcggccgtatcgtattccgtgca
gcacaacaccgtgatgacattgaagttgtaggtattaacgacttaatcgacgttgaatac
atggcttatatgttgaaatatgattcaactcacggtcgtttcgacggcactgttgaagtg
aaagatggtaacttagtggttaatggtaaaactatccgtgtaactgcagaacgtgatcca
gcaaacttaaactggggtgcaatcggtgttgatatcgctgttgaagcgactggtttattc
ttaactgatgaaactgctcgtaaacatatcactgcaggcgcaaaaaaagttgtattaact
ggcccatctaaagatgcaacccctatgttcgttcgtggtgtaaacttcaacgcatacgca
ggtcaagatatcgtttctaacgcatcttgtacaacaaactgtttagctcctttagcacgt
gttgttcatgaaactttcggtatcaaagatggtttaatgaccactgttcacgcaacgact
gcaactcaaaaaactgtggatggtccatcagctaaagactggcgcggcggccgcggtgca
tcacaaaacatcattccatcttcaacaggtgcagcgaaagcagtaggtaaagtattacct
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. . .   caaaaatagggttaatatgaatctcgatctccattttgttcatcgtattcaa
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aatacagcccagcaagcagaatttatcctaaatcacgccgatgtaaaaattctcttcgtc
ggcgatcaagagcaatacgatcaaacattggaaattgctcatcattgtccaaaattacaa
aaaattgtagcaatgaaatccaccattcaattacaacaagatcctctttcttgcacttgg
```

Molecular Biology Information: Protein Sequence

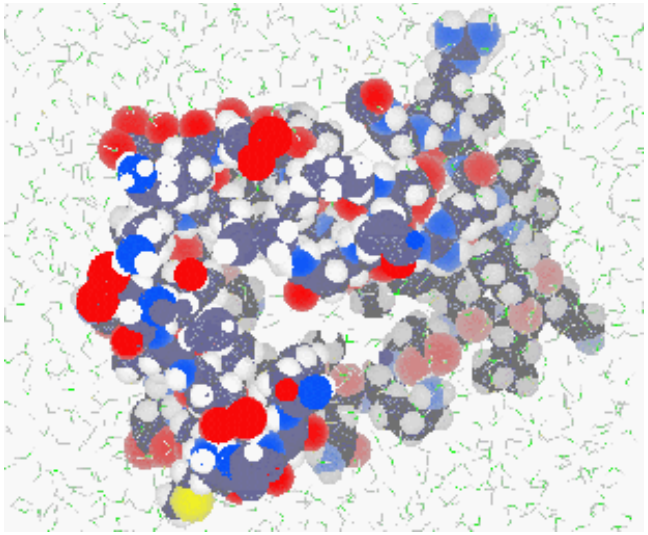
- 20 letter alphabet
  - ◊ ACDEFGHIKLMNPQRSTVWY but not BJOUXZ
- Strings of ~300 aa in an average protein (in bacteria), ~200 aa in a domain
- >12 M known protein sequences (uniprot, http://www.ebi.ac.uk/uniprot/TrEMBLstats/, 2011)

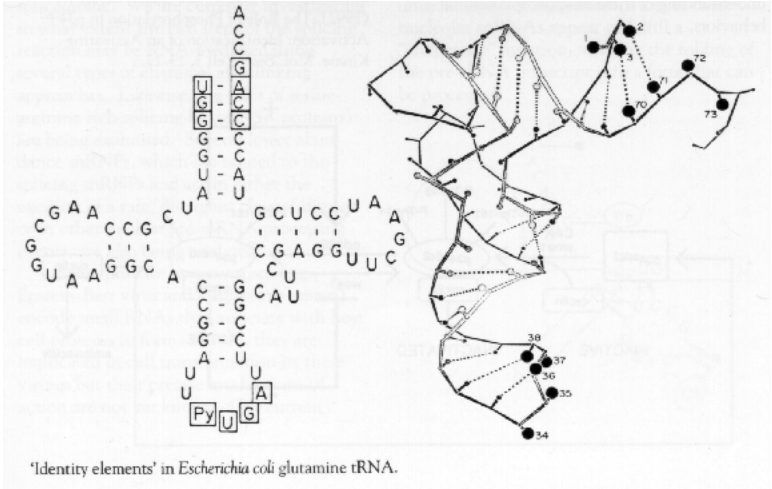
```
d1dhfa_ LNCIVAVSQNMGIGKNGDLPWPPLRNEFRYFQRMTTTSSVEGKQ-NLVIMGKKTWFSI
d8dfr__ LNSIVAVCQNMGIGKDGNLPWPPLRNEYKYFQRMTSTSHVEGKQ-NAVIMGKKTWFSI
d4dfra_ ISLIAALAVDRVIGMENAMPWN-LPADLAWFKRNTL--------NKPVIMGRHTWESI
d3dfr__ TAFLWAQDRDGLIGKDGHLPWH-LPDDLHYFRAQTV--------GKIMVVGRRTYESF

d1dhfa_ LNCIVAVSQNMGIGKNGDLPWPPLRNEFRYFQRMTTTSSVEGKQ-NLVIMGKKTWFSI
d8dfr__ LNSIVAVCQNMGIGKDGNLPWPPLRNEYKYFQRMTSTSHVEGKQ-NAVIMGKKTWFSI
d4dfra_ ISLIAALAVDRVIGMENAMPW-NLPADLAWFKRNTLD--------KPVIMGRHTWESI
d3dfr__ TAFLWAQDRNGLIGKDGHLPW-HLPDDLHYFRAQTVG--------KIMVVGRRTYESF

d1dhfa_ VPEKNRPLKGRINLVLSRELKEPPQGAHFLSRSLDDALKLTEQPELANKVDMVWIVGGSSVYKEAMNHP
d8dfr__ VPEKNRPLKDRINIVLSRELKEAPKGAHYLSKSLDDALALLDSPELKSKVDMVWIVGGTAVYKAAMEKP
d4dfra_ ---G-RPLPGRKNIILS-SQPGTDDRV-TWVKSVDEAIAACGDVP------EIMVIGGGRVYEQFLPKA
d3dfr__ ---PKRPLPERTNVVLTHQEDYQAQGA-VVVHDVAAVFAYAKQHLDQ----ELVIAGGAQIFTAFKDDV

d1dhfa_ -PEKNRPLKGRINLVLSRELKEPPQGAHFLSRSLDDALKLTEQPELANKVDMVWIVGGSSVYKEAMNHP
d8dfr__ -PEKNRPLKDRINIVLSRELKEAPKGAHYLSKSLDDALALLDSPELKSKVDMVWIVGGTAVYKAAMEKP
d4dfra_ -G---RPLPGRKNIILSSSQPGTDDRV-TWVKSVDEAIAACGDVPE-----  IMVIGGGRVYEQFLPKA
d3dfr__ -P--KRPLPERTNVVLTHQEDYQAQGA-VVVHDVAAVFAYAKQHLD----QELVIAGGAQIFTAFKDDV
```

Molecular Biology Information: Macromolecular Structure

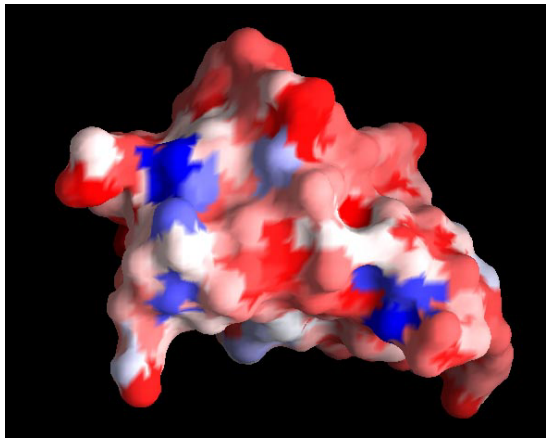
- DNA/RNA/Protein
  - ◊ Almost all protein

(RNA Adapted From D Soll Web Page,
Right Hand Top Protein from M Levitt web page)





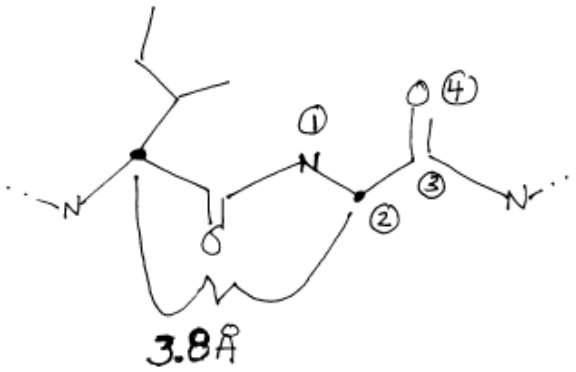
'Identity elements' in *Escherichia coli* glutamine tRNA.



Molecular Biology Information: Protein Structure Details

- Statistics on Number of XYZ triplets
  - ◊ 200 residues/domain –> 200 CA atoms, separated by 3.8 A
  - ◊ Avg. Residue is Leu: 4 backbone atoms + 4 sidechain atoms, 150 cubic A
    - => ~1500 xyz triplets (=8x200) per protein domain
  - ◊ >110K Domains, ~1200 folds (scop 1.75)

```
ATOM      1  C   ACE     0       9.401  30.166  60.595  1.00 49.88      1GKY  67
ATOM      2  O   ACE     0      10.432  30.832  60.722  1.00 50.35      1GKY  68
ATOM      3  CH3 ACE     0       8.876  29.767  59.226  1.00 50.04      1GKY  69
ATOM      4  N   SER     1       8.753  29.755  61.685  1.00 49.13      1GKY  70
ATOM      5  CA  SER     1       9.242  30.200  62.974  1.00 46.62      1GKY  71
ATOM      6  C   SER     1      10.453  29.500  63.579  1.00 41.99      1GKY  72
ATOM      7  O   SER     1      10.593  29.607  64.814  1.00 43.24      1GKY  73
ATOM      8  CB  SER     1       8.052  30.189  63.974  1.00 53.00      1GKY  74
ATOM      9  OG  SER     1       7.294  31.409  63.930  1.00 57.79      1GKY  75
ATOM     10  N   ARG     2      11.360  28.819  62.827  1.00 36.48      1GKY  76
ATOM     11  CA  ARG     2      12.548  28.316  63.532  1.00 30.20      1GKY  77
ATOM     12  C   ARG     2      13.502  29.501  63.500  1.00 25.54      1GKY  78
...
ATOM   1444  CB  LYS   186      13.836  22.263  57.567  1.00 55.06      1GKY1510
ATOM   1445  CG  LYS   186      12.422  22.452  58.180  1.00 53.45      1GKY1511
ATOM   1446  CD  LYS   186      11.531  21.198  58.185  1.00 49.88      1GKY1512
ATOM   1447  CE  LYS   186      11.452  20.402  56.860  1.00 48.15      1GKY1513
ATOM   1448  NZ  LYS   186      10.735  21.104  55.811  1.00 48.41      1GKY1514
ATOM   1449  OXT LYS   186      16.887  23.841  56.647  1.00 62.94      1GKY1515
TER    1450      LYS   186                                              1GKY1516
```



Molecular Biology Information: Whole Genomes

- The Revolution Driving Everything

  Fleischmann, R. D., Adams, M. D., White, O., Clayton, R. A., Kirkness, E. F., Kerlavage, A. R., Bult, C. J., Tomb, J. F., Dougherty, B. A., Merrick, J. M., McKenney, K., Sutton, G., Fitzhugh, W., Fields, C., Gocayne, J. D., Scott, J., Shirley, R., Liu, L. I., Glodek, A., Kelley, J. M., Weidman, J. F., Phillips, C. A., Spriggs, T., Hedblom, E., Cotton, M. D., Utterback, T. R., Hanna, M. C., Nguyen, D. T., Saudek, D. M., Brandon, R. C., Fine, L. D., Fritchman, J. L., Fuhrmann, J. L., Geoghagen, N. S. M., Gnehm, C. L., McDonald, L. A., Small, K. V., Fraser, C. M., Smith, H. O. & Venter, J. C. (1995). "Whole-genome random sequencing and assembly of Haemophilus influenzae rd." *Science* 269: 496-512.

  (Picture adapted from TIGR website, http://www.tigr.org)

- Timeline

  1995, HI (bacteria): 1.6 Mb & 1600 genes done

  1997, yeast: 13 Mb & ~6000 genes for yeast

  1998, worm: ~100Mb with 19 K genes

  1999: >30 completed genomes!

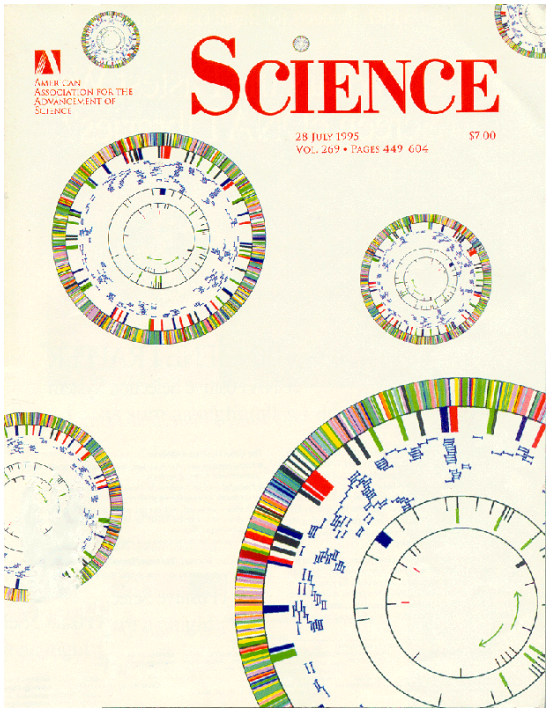
  2000, draft human

  2003, human: 3 Gb & 100 K genes...

  2010, 1000 human genomes!

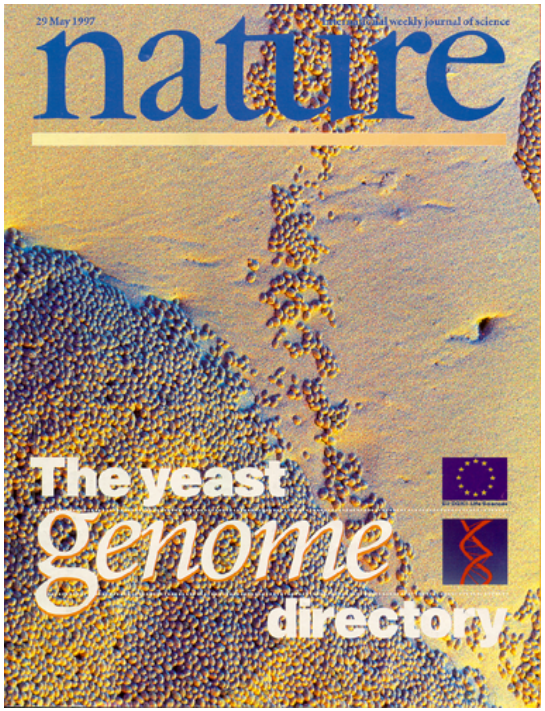
1995

Bacteria,
1.6 Mb,
~1600 genes
[*Science* **269**: 496]



1997

Eukaryote,
13 Mb,
~6K genes
[Nature **387**: 1]



1998

Animal,
~100 Mb,
~20K genes
[*Science* **282**:
1945]

2000?

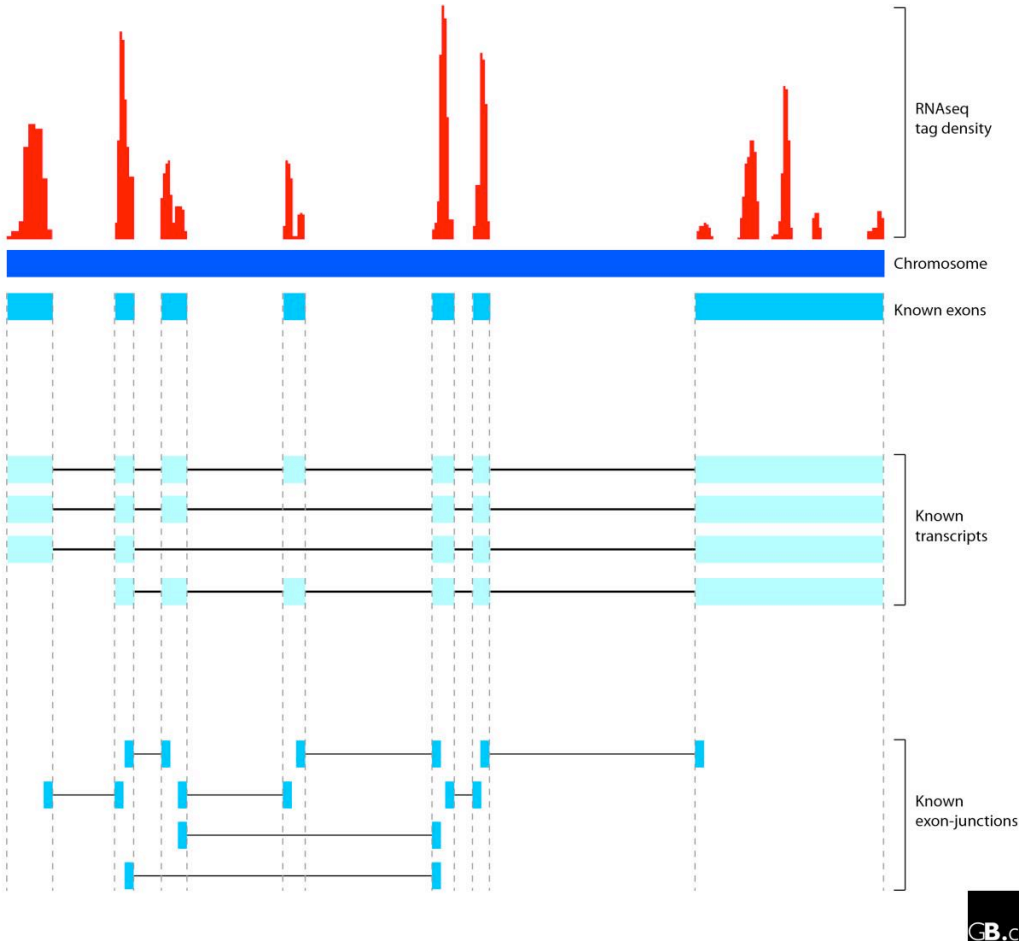
Human,
~3 Gb,
~20K genes

‘98 spoof

A Bioinformatics prediction that came true!

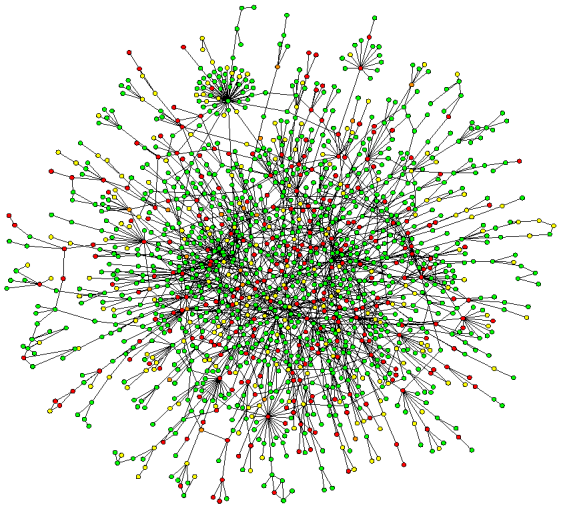
real thing, Apr ‘00

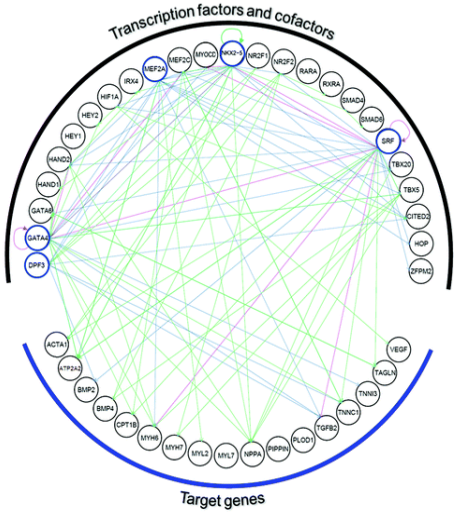
Gene Expression Data: On & Off



- Early experiments yeast
  - ◊ Complexity at 10 time points,
    6000 x 10 = 60K floats
- Then tiling array technology
  - ◊ 50 M data points to tile the human genome at ~50 bp res.
- Now Next-Gen Sequencing (RNAseq)
  - ◊ 10M+ reads on the human genome, counts
- Can only sequence genome once but can do an infinite variety of expression experiments

Molecular Networks: Connectivity





Regulatory Networks

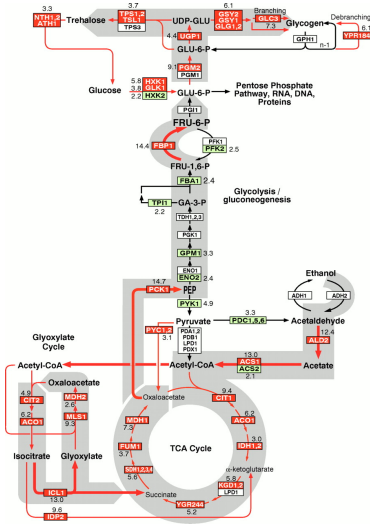
Get readouts of where proteins bind to DNA : Chip-chip then chip-seq

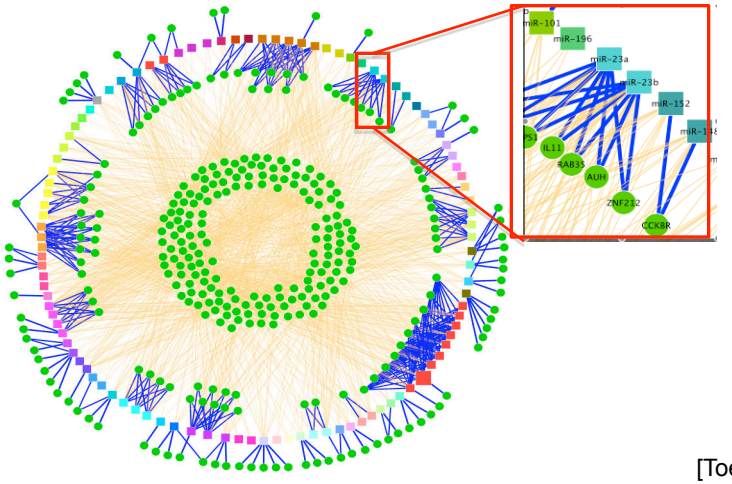
Protein Interaction Networks

For yeast: 6000 x 6000 / 2 ~ 18M possible interactions (maybe ~30K real)

Protein-protein Interaction networks

TF-target-gene Regulatory networks





Metabolic pathway networks

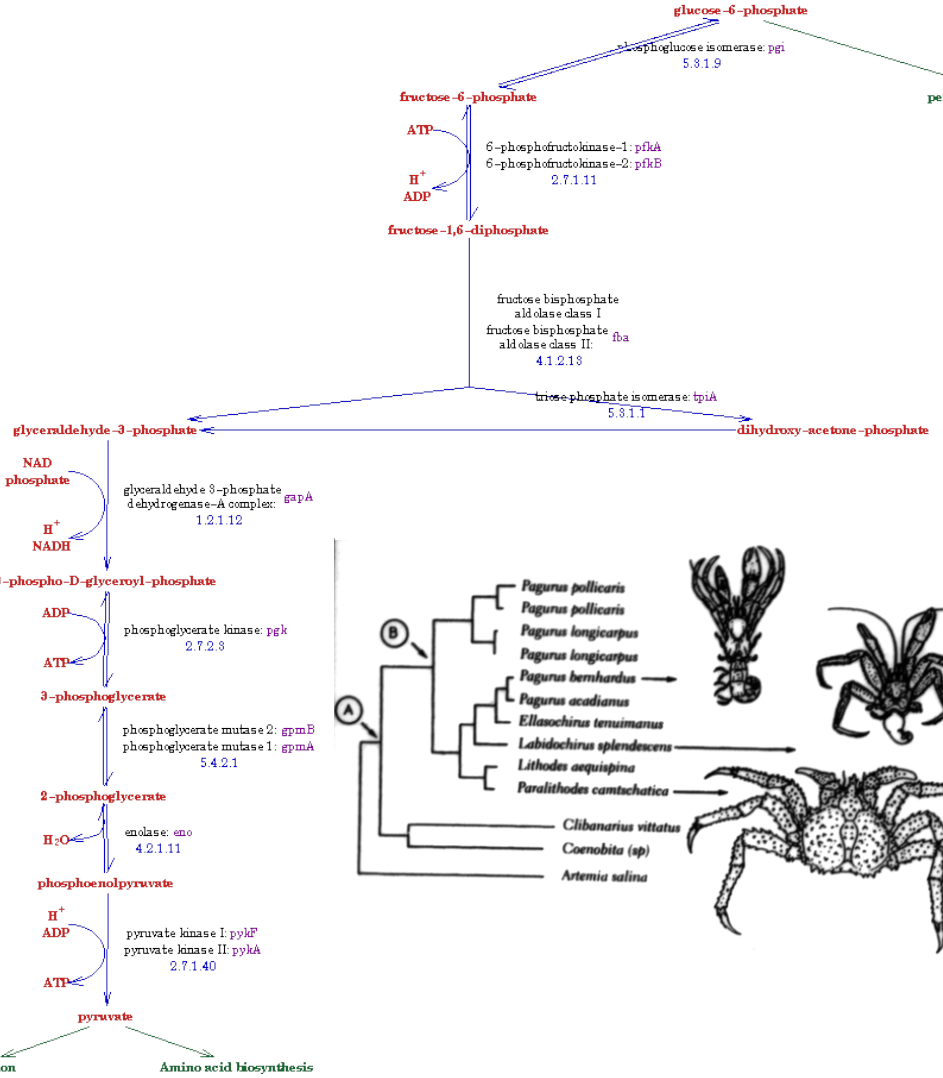
miRNA-target networks

[Toenjes, *et al*, *Mol. BioSyst.* (2008); Jeong *et al*, *Nature* (2001); [Horak, et al, Genes & Development, 16:3017-3033; DeRisi, Iyer, and Brown, Science, 278:680-686]

Molecular Biology Information: Other Integrative Data

- Information to understand genomes
  - ◊ Whole Organisms Phylogeny, traditional zoology
  - ◊ Environments, Habitats, ecology
  - ◊ Phenotype Experiments (large-scale KOs, transposons)
  - ◊ The Literature (MEDLINE)
- The Future....

(Pathway drawing from P Karp's EcoCyc, Phylogeny from S J Gould, Dinosaur in a Haystack)



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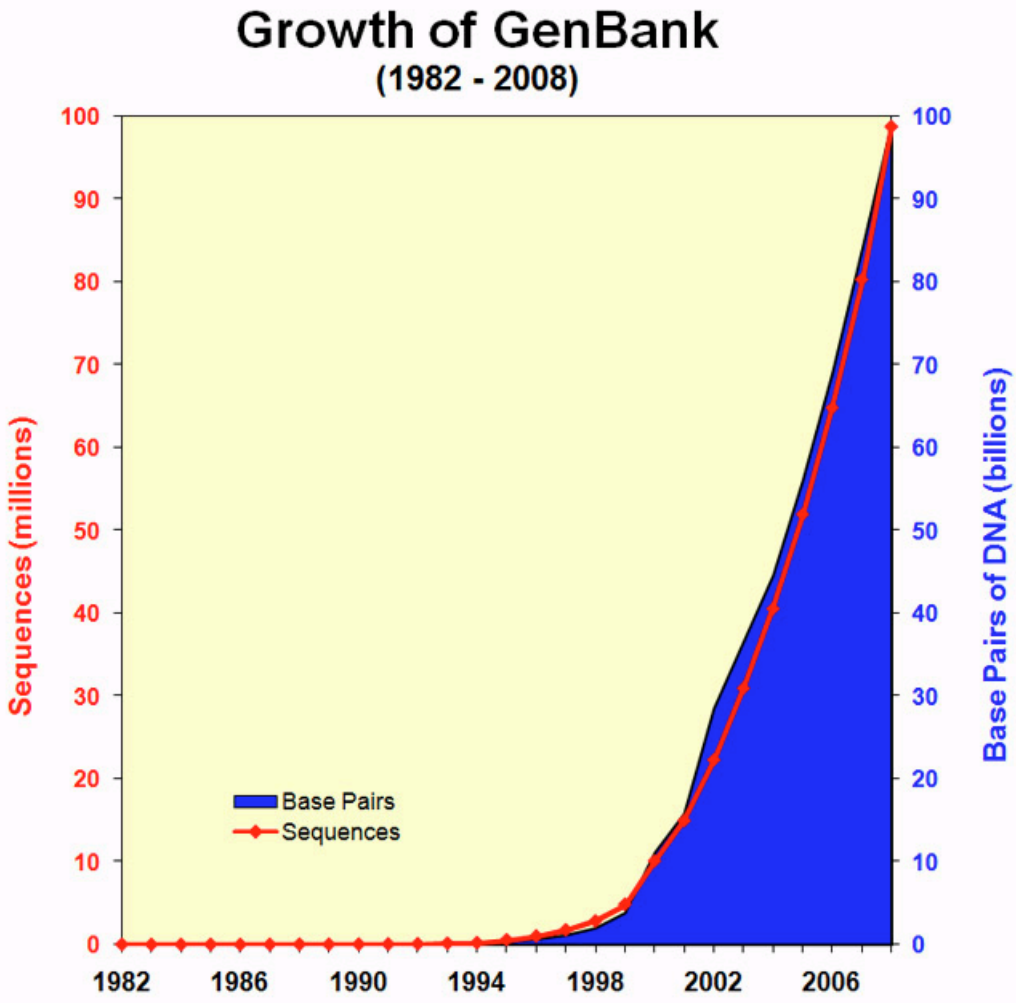
- Bioinformatics is a practical discipline with many **applications**.

[Luscombe et al. ('01). *Methods Inf Med* 40: 346]

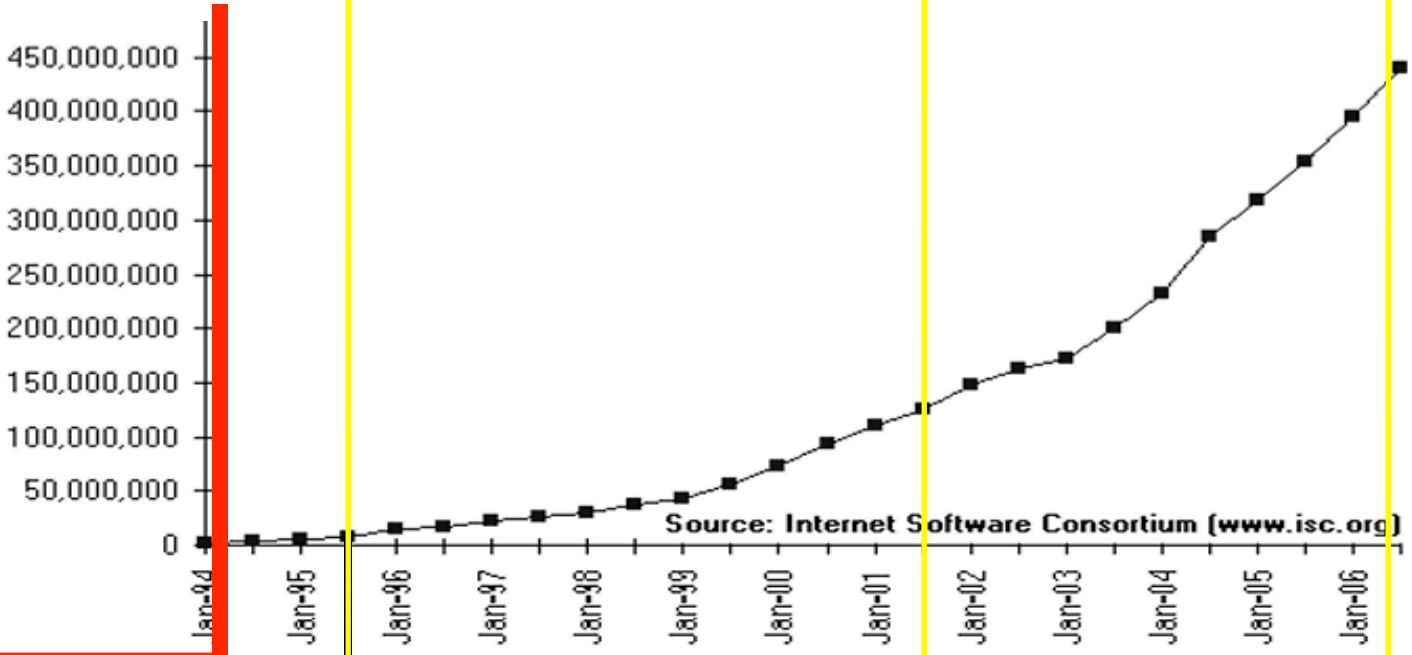
Large-scale Information: Explonential **Scaling** of Data Matched by Development of Computer Technology

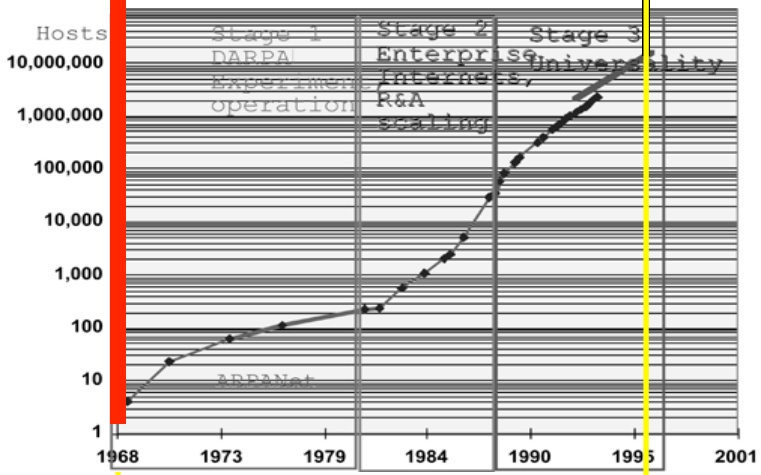
- CPU vs Disk & Net
  - ◊ As important as the increase in computer speed has been, the ability to store large amounts of information on computers is even more crucial
  - ◊ Comparison with **Moore's Law**
- A Driving Force in Bioinformatics



Internet Hosts

(adapted from D Brutlag, Stanford & http://navigators.com/stats.html)



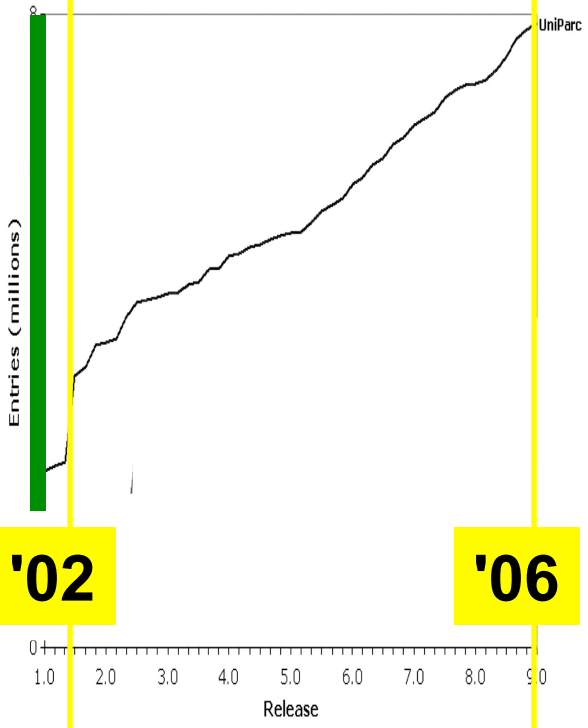


'68

'95

Proteins

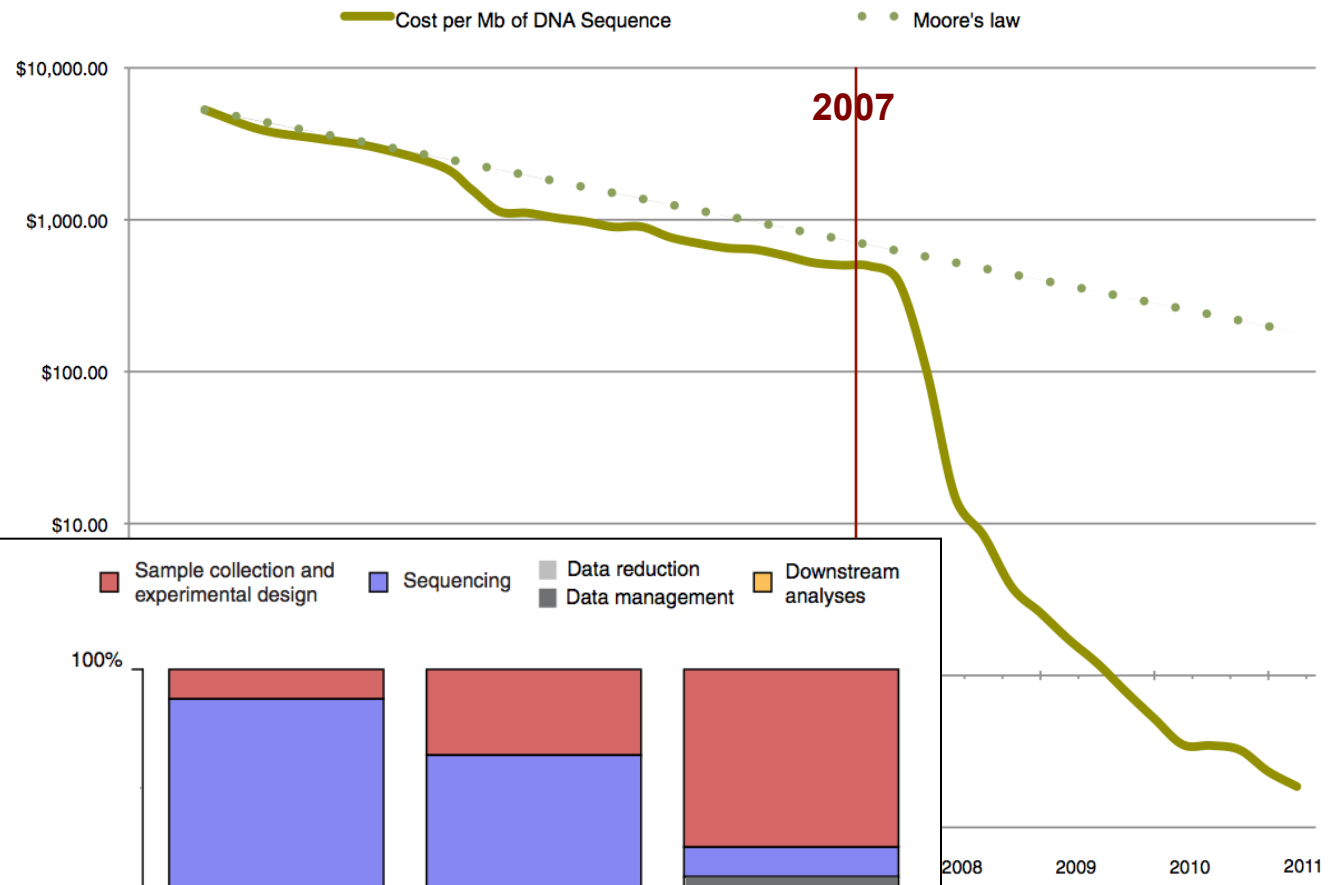
Suzek, B. E. et al. Bioinformatics 2007 23:1282-1288; doi: 10.1093/bioinformatics/btm098

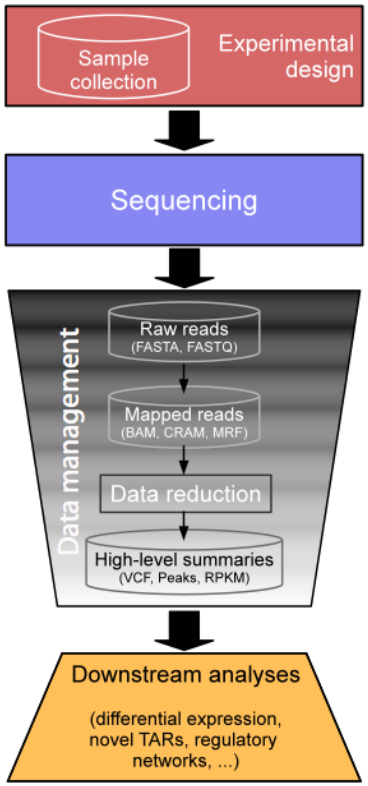


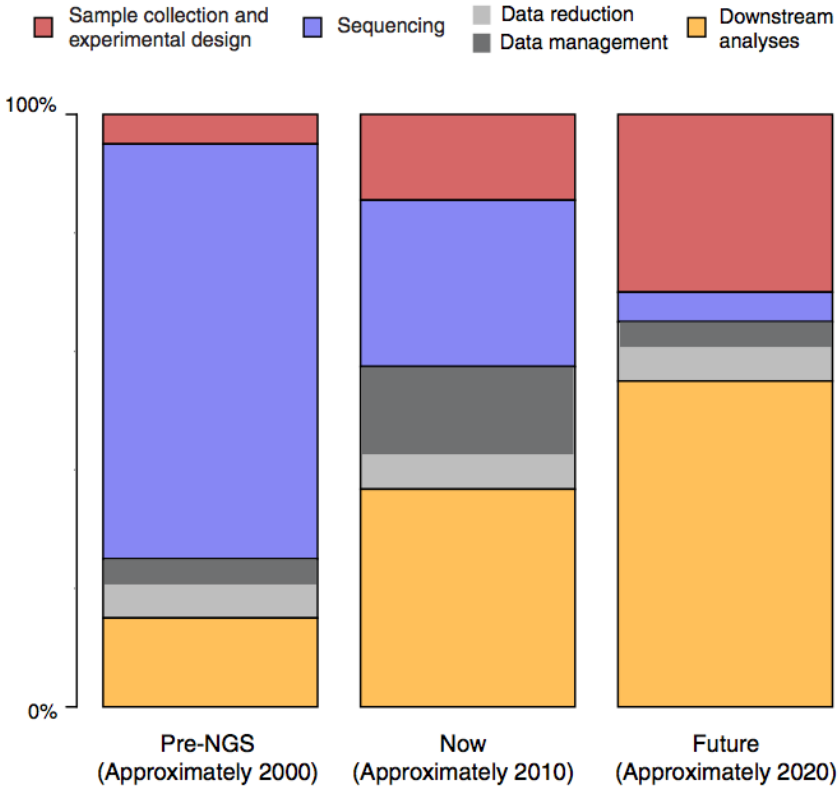
'02

'06

Sequencing Data Explosion: Going to $0/base



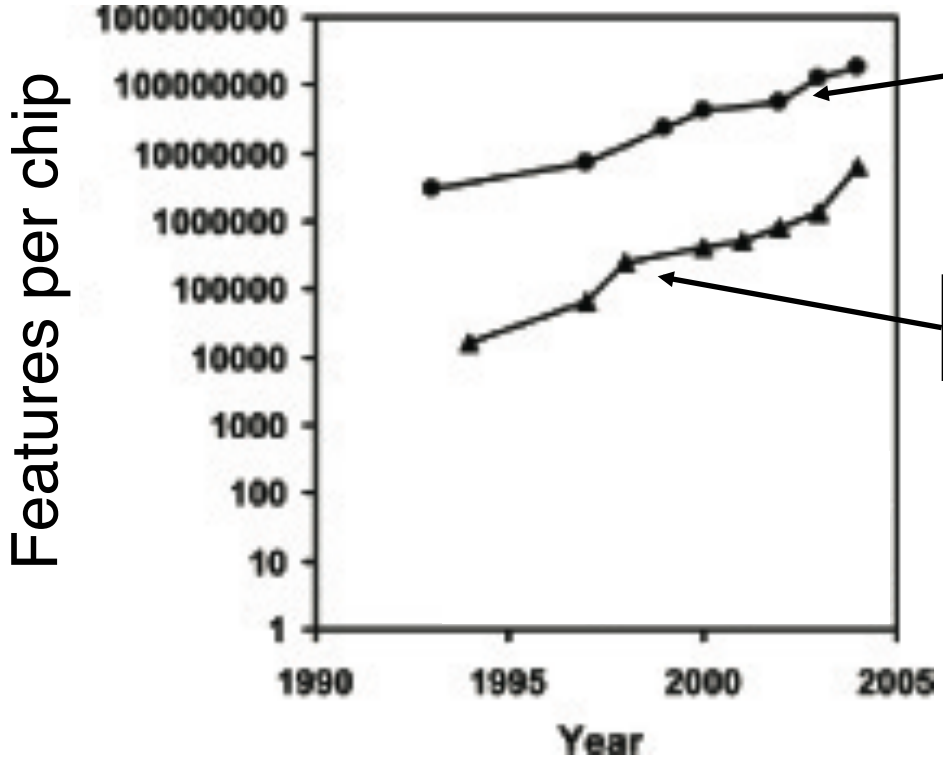




From '00 to ~' 20, cost of DNA sequencing expt. shifts from the actual seq. to sample collection & analysis

[Sboner et al. ('11) GenomeBiology]

Features per Slide



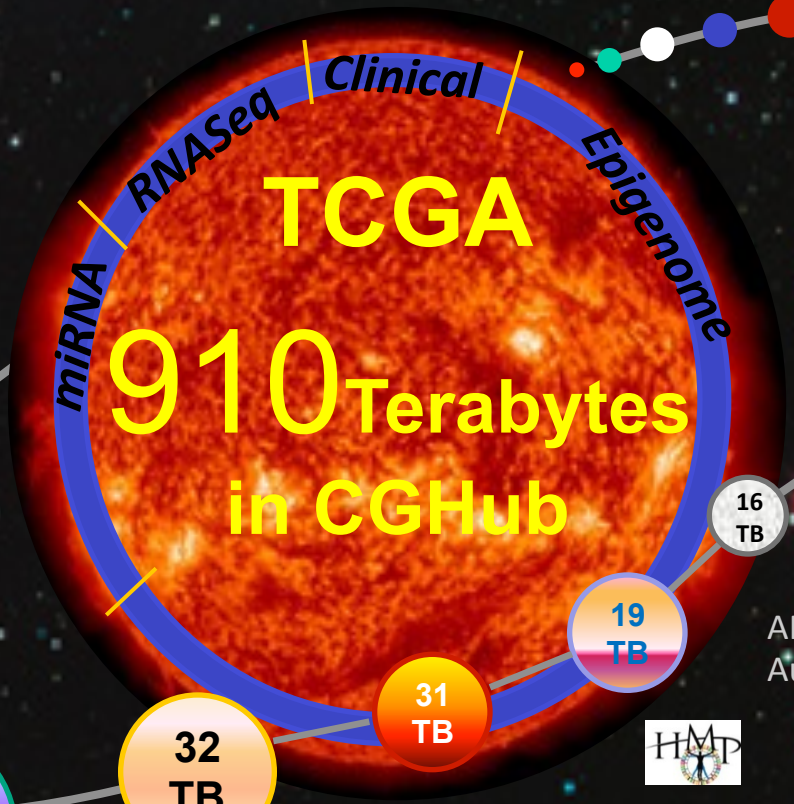
Chip Technology

Seq Universe

[from Heidi Sofia, NHGRI]

SRA >1 petabyte

TCGA endpoint: ~2.5 Petabytes
- **~1.5 PB exome**
- **~1 PB whole genome**



220 TB

46 TB

32 TB

31 TB

19 TB

16 TB

9 TB

ENCODE

ARRA
Autism

HMP

National Human Genome Research Institute

ADSP

NHLBI ESP

NHGRI LSSP

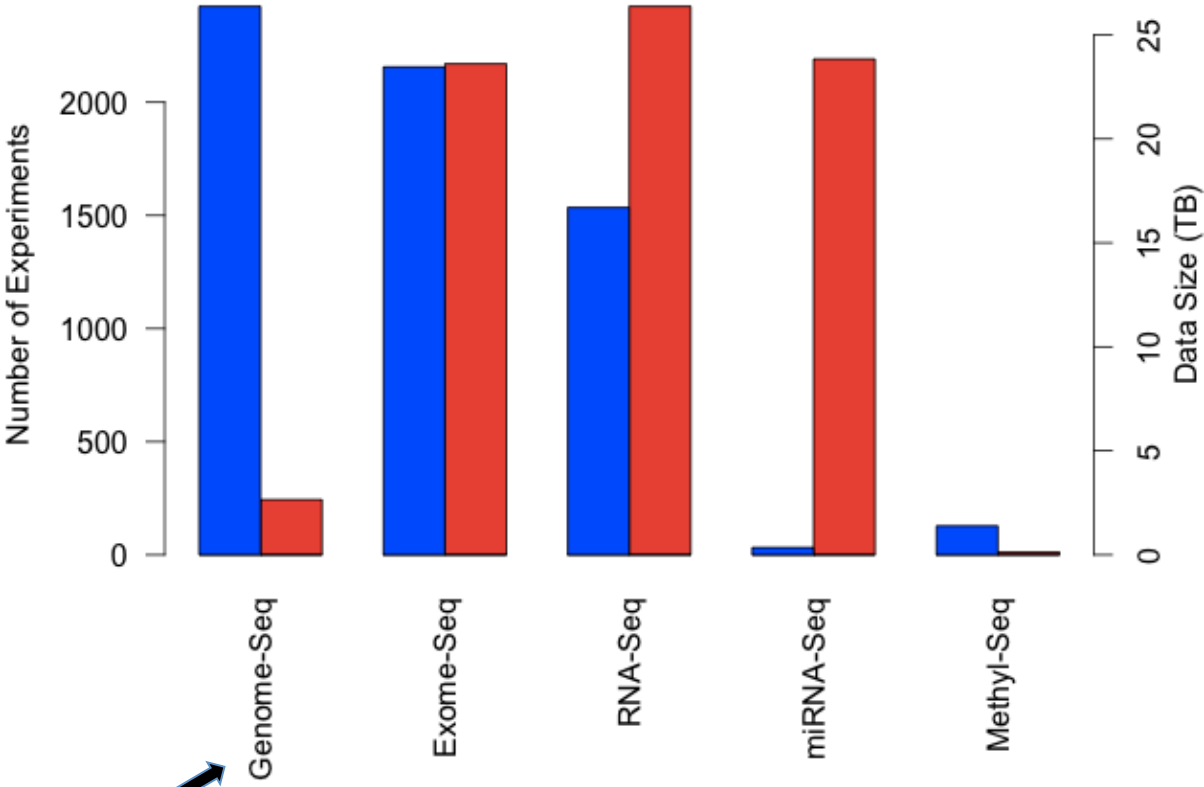
Star formation
100K Genomes England

Sofia, 2-28-14

JESS3

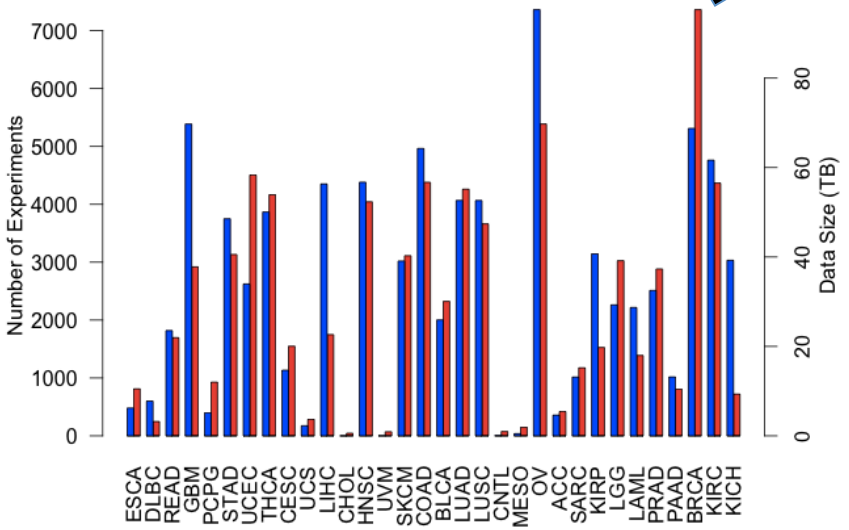
TCGA: What's in a petabyte?

- >73,000 Expt
- 34 Cancer Types
- ~5,000 Patients



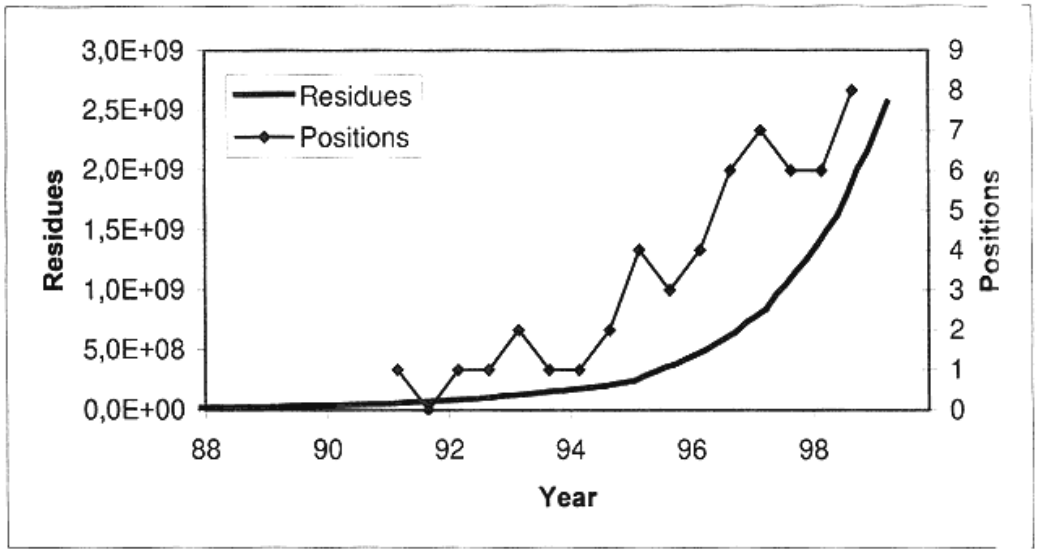
Experiments
Data Size (TB)

TCGA Cancer Types

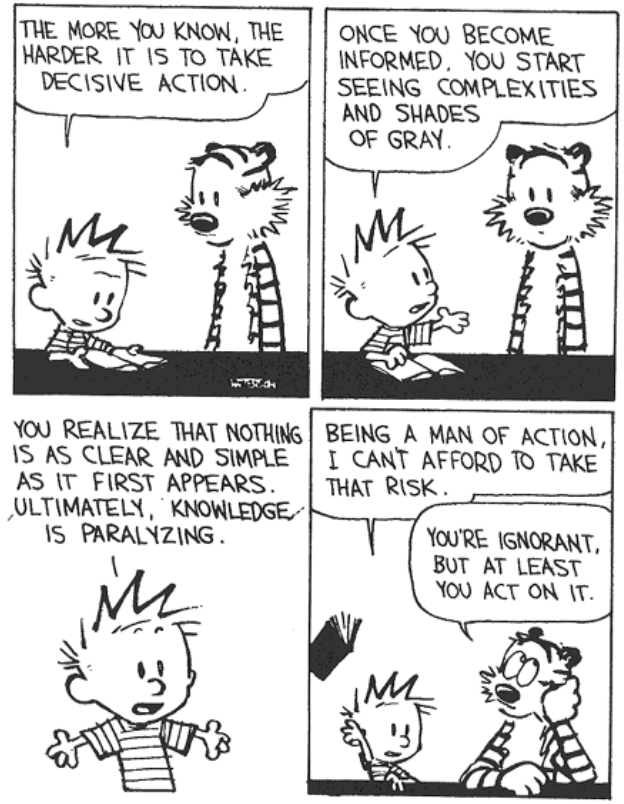


Breast Cancer Expt. Types

Jobs: Bioinformatics is born!



Growth in number of residues in Genbank, a central database for sequence data, compared to the request for people with competence in bioinformatics. The request for scientists is estimated from the number of relevant positions advertised in the first number of Nature in March and September of each year.



B. Watterson, "*There's treasure everywhere*", Andrews and McMeel, 1996.

(courtesy of Finn Drablos)

What is Bioinformatics?

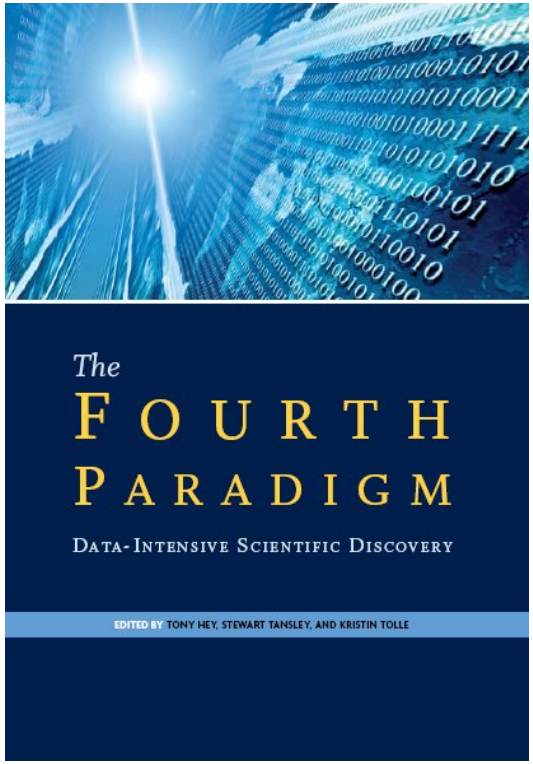
- *(Molecular)* **Bio - `informatics`**
- One idea for a definition?
  Bioinformatics is conceptualizing **biology in terms of molecules** (in the sense of physical-chemistry) and then applying **"informatics" techniques** (derived from disciplines such as applied math, CS, and statistics) to **organize, mine, model & understand the information associated** with these molecules, **on a large-scale.**
- Bioinformatics is a practical discipline with many **applications**.

[Luscombe et al. ('01). *Methods Inf Med* 40: 346]

General Types of "Informatics" techniques in Computational Biology

- Databases
  - Building, Querying
  - Representing Complex data
- Data mining
  - Machine Learning techniques
  - Clustering & Tree construction
  - Rapid Text String Comparison & textmining
  - Detailed statistics of significance & association
- Network Analysis
  - Analysis of Topology (eg Hubs)
  - Predicting Connectivity
- Structure Analysis & Geometry
  - Graphics (Surfaces, Volumes)
  - Comparison & 3D Matching (Vision, recognition, docking)
- Physical Modeling
  - Newtonian Mechanics
  - Electrostatics
  - Numerical Algorithms
  - Simulation
  - Modeling Chemical Reactions & Cellular Processes

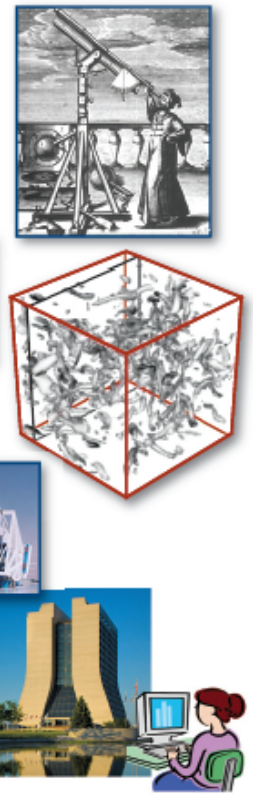
Jim Gray's 4th Paradigm



Science Paradigms

- Thousand years ago:
  science was **empirical**
  describing natural phenomena
- Last few hundred years:
  theoretical branch
  using models, generalizations
- Last few decades:
  a **computational** branch
  simulating complex phenomena
- Today: **data exploration** (eScience)
  unify theory, experiment, and simulation
  - Data captured by instruments or generated by simulator
  - Processed by software
  - Information/knowledge stored in computer
  - Scientist analyzes database/files using data management and statistics

$$\left(\frac{\dot{a}}{a}\right)^2 = \frac{4\pi G p}{3} - K\frac{c^2}{a^2}$$



Jim Gray's 4th Paradigm

#3 - Simulation

Prediction based on physical principles (eg Exact Determination of Rocket Trajectory)

Emphasis on: Supercomputers

#4 - Data Mining

Classifying information & discovering unexpected relationships

Emphasis: networks, "federated" DBs

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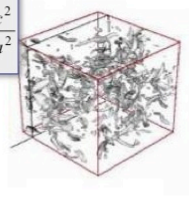
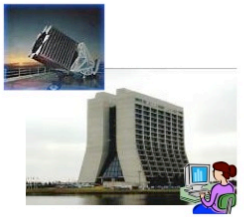
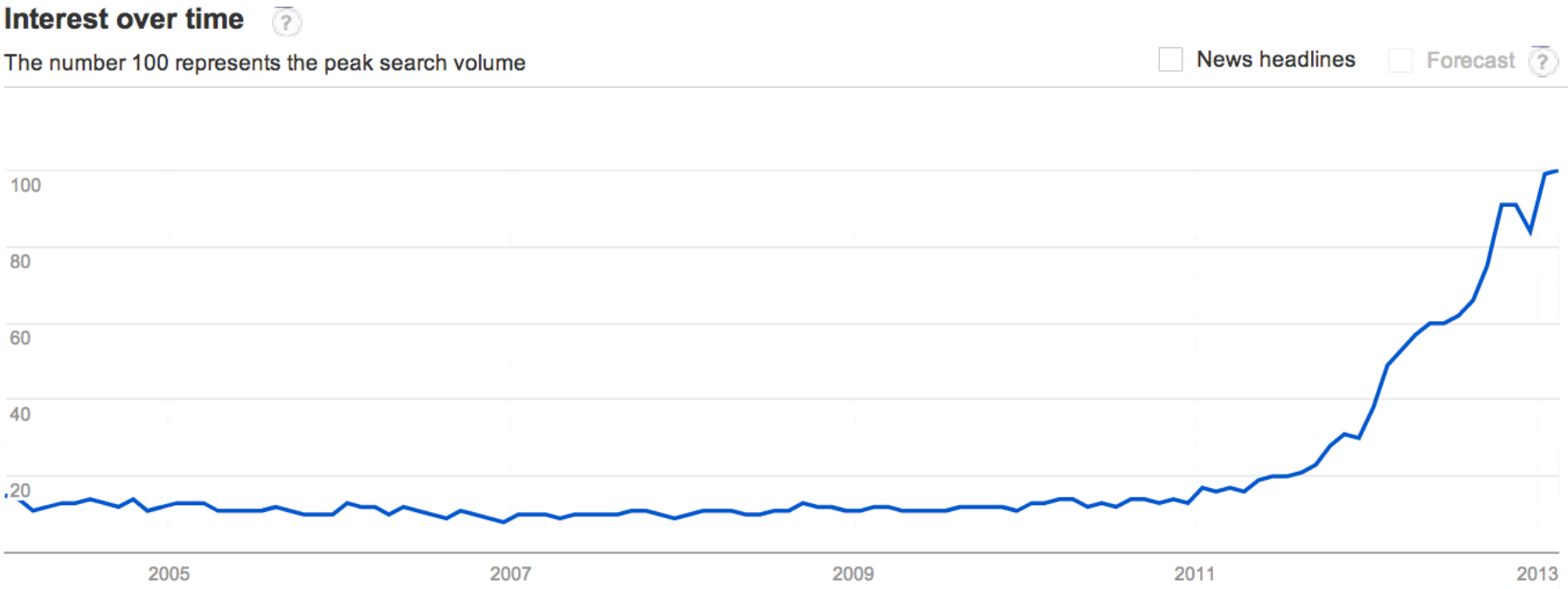




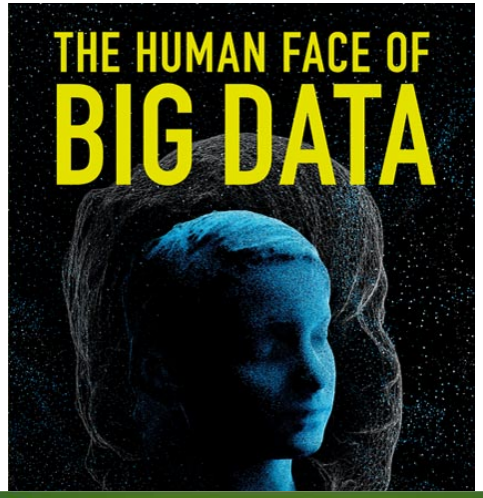
Gray died in '07.
Book about his ideas came out in '09…..

[Slide from : http://research.microsoft.com/en-us/um/people/gray/talks/stanford%2520symbolic%2520systems%2520seminar.ppt]

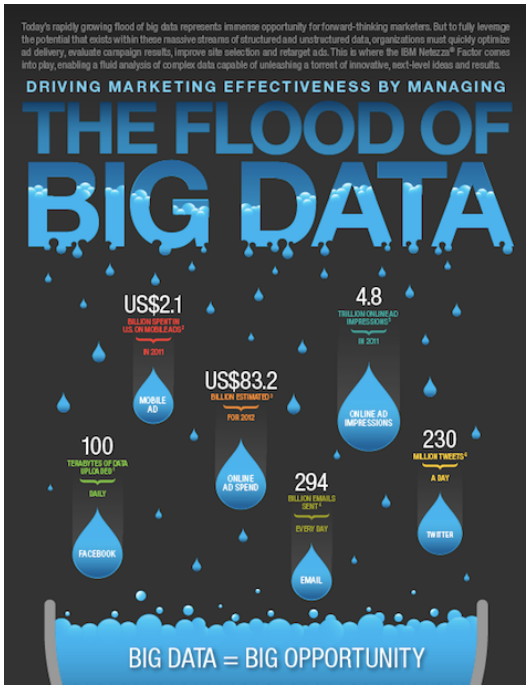
Since ‘07, continued “steady” increase in computation;Even Larger Data Growth in Last Decade, Giving Rise to the Analysis of Many Big Data Sets

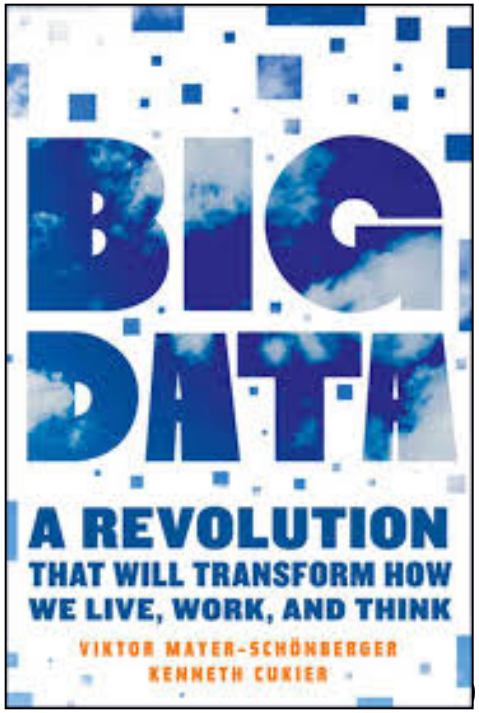


Google Trends



Commercial World Data: Financial & Retail Data





Big Data:

a current buzz-word

Harvard Business Review

Data Scientist: The Sexiest Job of the 21st Century

by Thomas H. Davenport and D.J. Patil

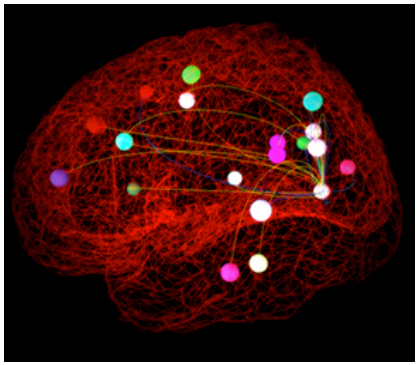
Artwork: **Tamar Cohen, *Andrew J Buboltz,*** 2011, silk screen on a page from a high

When Jonathan Goldman arrived for work in June 2006 at LinkedIn, the business ne
up. The company had just under 8 million accounts, and the number was growing qu
friends and colleagues to join. But users weren't seeking out connections with the pe
rate executives had expected. Something was apparently missing in the social expe

[Oct. '12 issue]

Big data is transforming science

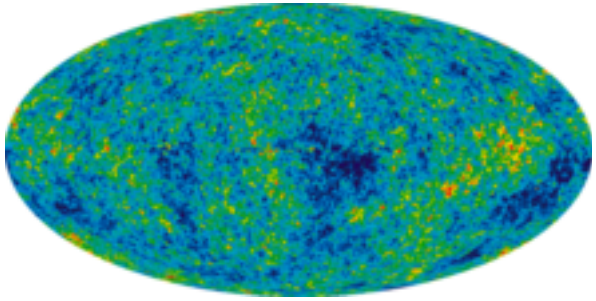
High energy physics -
Large Hadron Collider



Neuroscience -
The Human Connectome Project

Ecology - Fluxnet

Genomics
DNA sequencer



Astronomy -
Sloan Digital Sky survey

ISI Web of KNOWLEDGE
Transforming Research

Knowledge of knowledge
Meta-data of scientific documents

Computational social science
Online communities

What do people do with Big Data ?

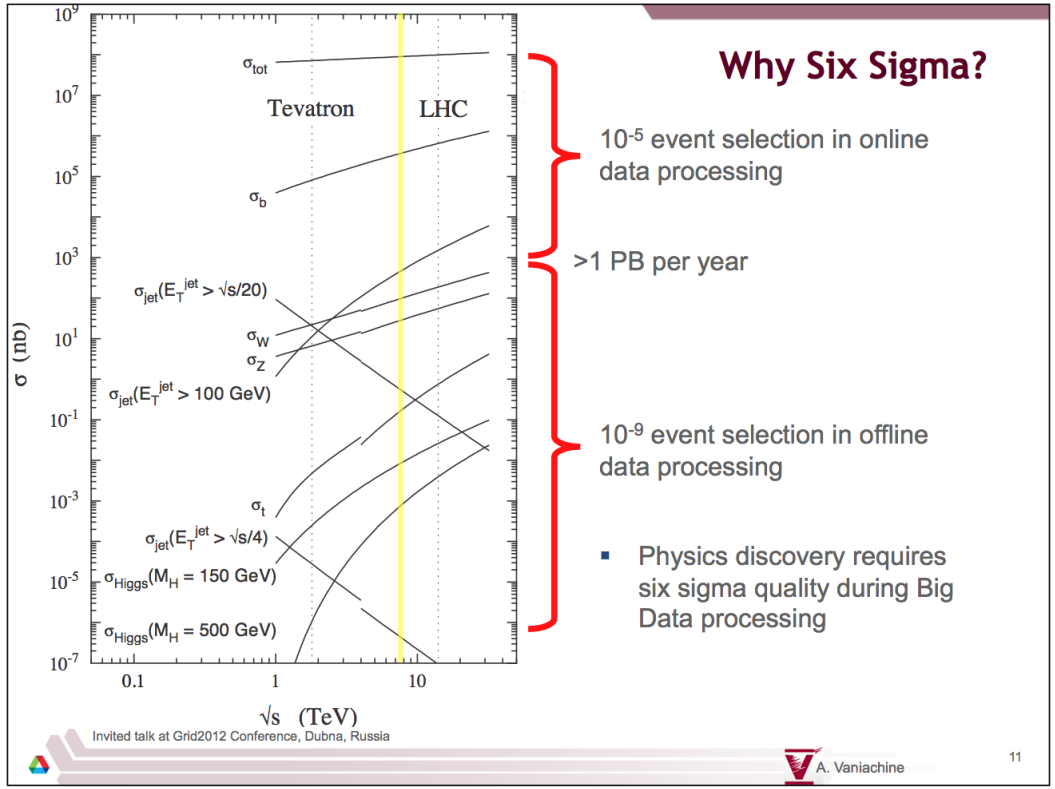
[*Nature* **489:** 208]

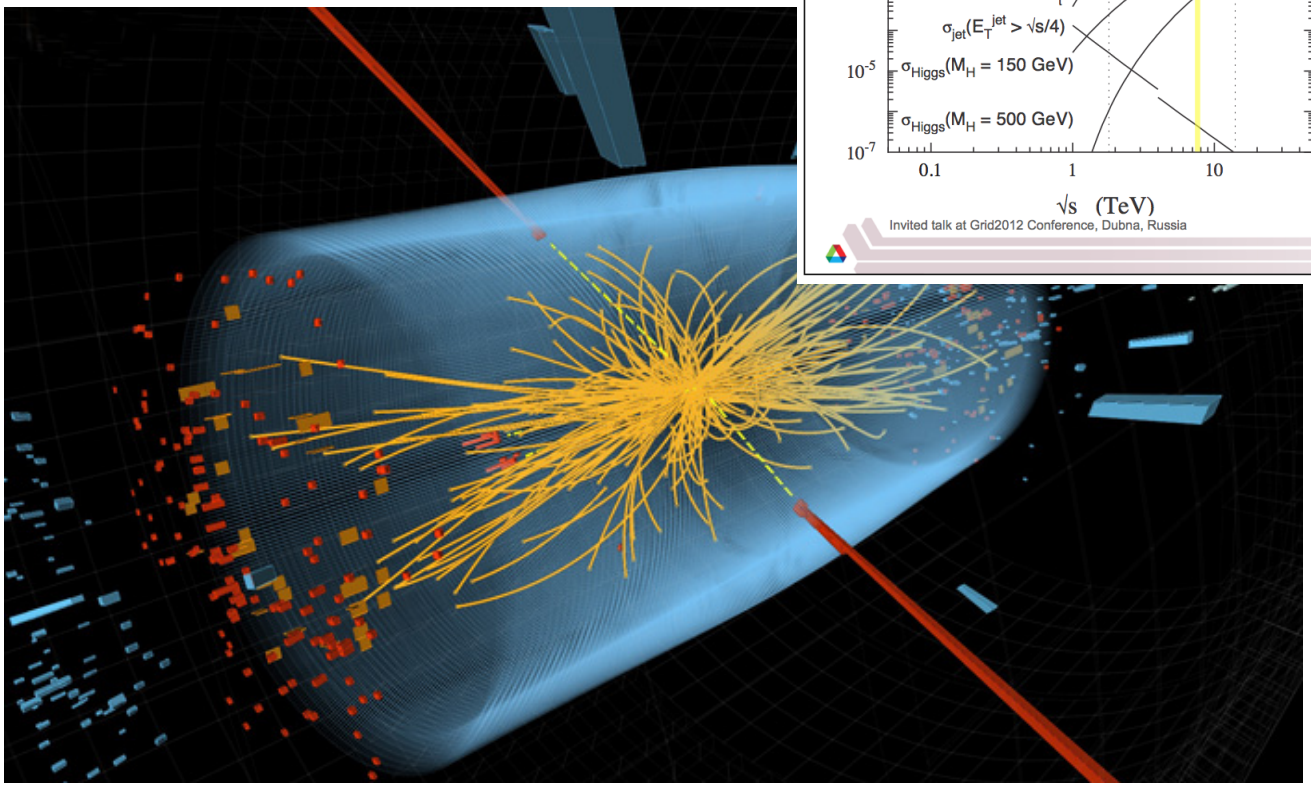
- Fundamental goal is general understanding & answering specific Qs: modeling & making predictions
- **Explicit Description of Data not Important --** Fast query, hiding underlying structure (e.g. Google **Search**)
- **Explicit Description of Data Important –** Organization highlighting underlying structure (e.g. Google **Maps**)

http://www.theatlantic.com/technology/archive/2012/01/to-know-but-not-understand-david-weinberger-on-science-and-big-data/250820/

Higgs Boson: Searching Through Many Events for a Few Needles

http://images.iop.org/objects/phw/news/16/7/4/higgs-cms.jpg



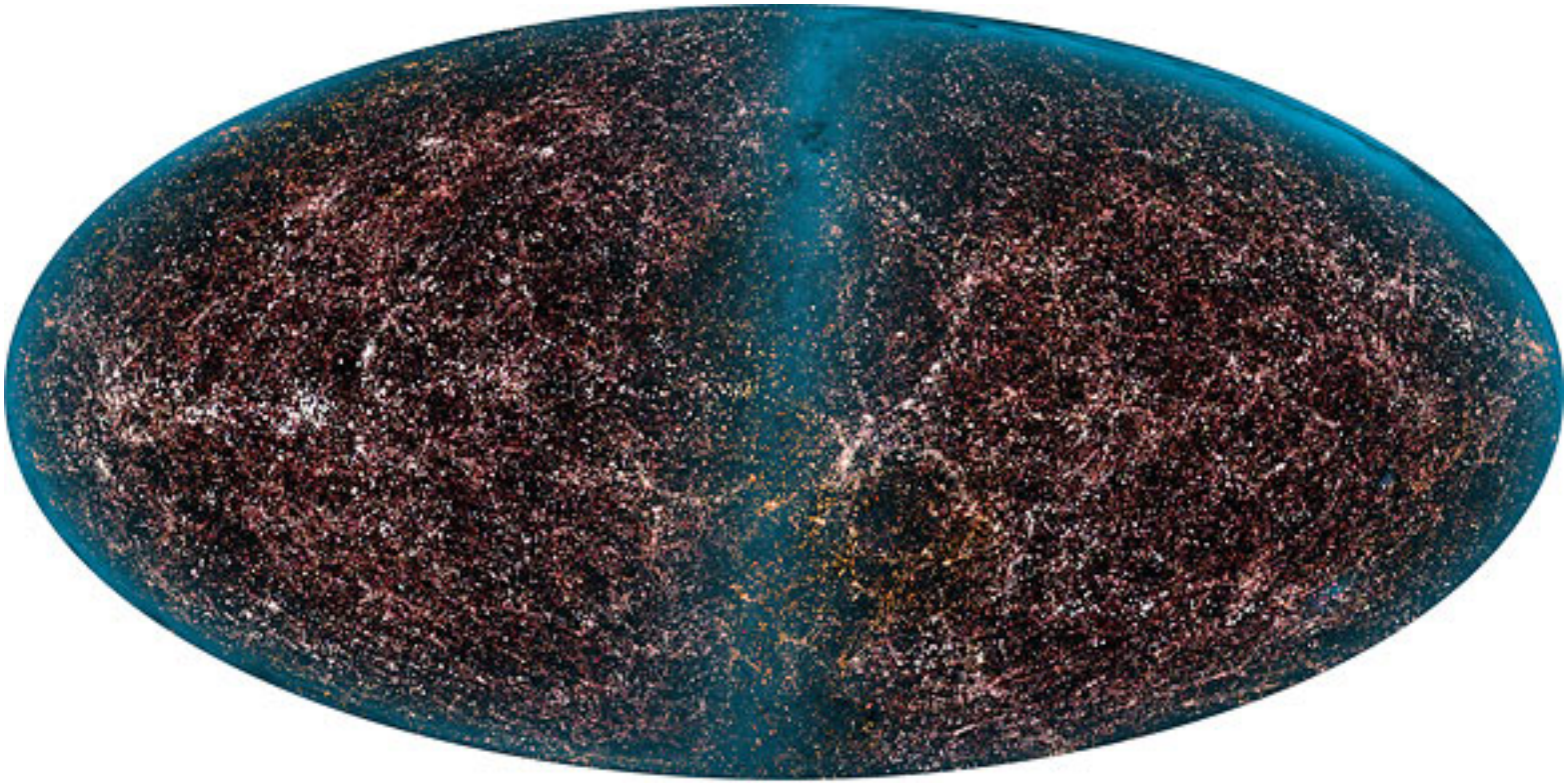


"Golden" Events

One H → 4 l / Billion

http://cdsweb.cern.ch/record/1462234/files/ATL-SOFT-SLIDE-2012-441.pdf

Making Intuitive Maps, Highlighting Large-scale Structure of Stars & the Earth

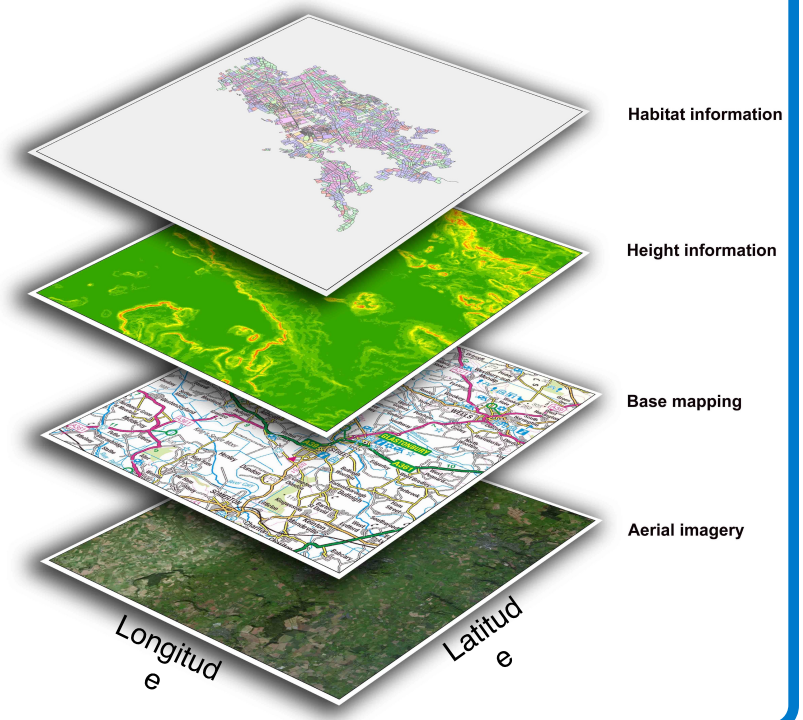


[SDSS.org]

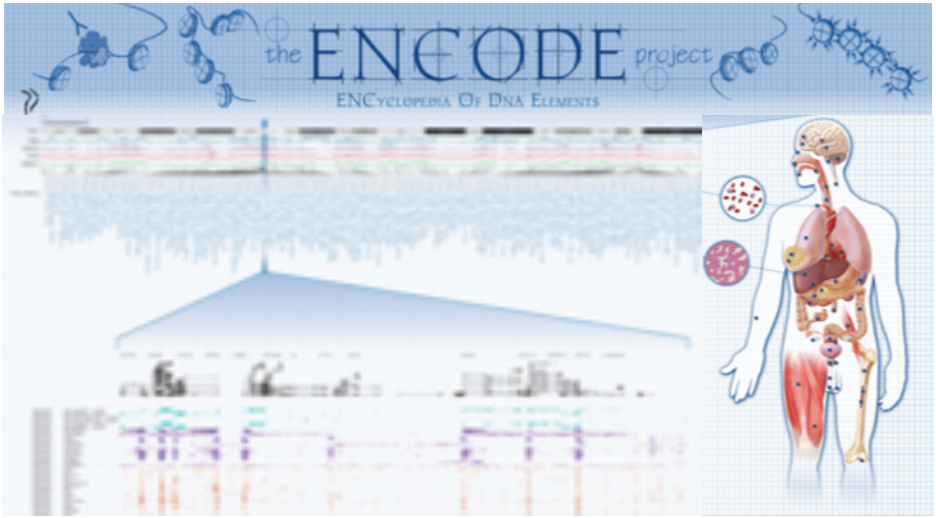
http://www.nrc-cnrc.gc.ca/eng/education/astronomy/tapping/2011/2011-03-02.html

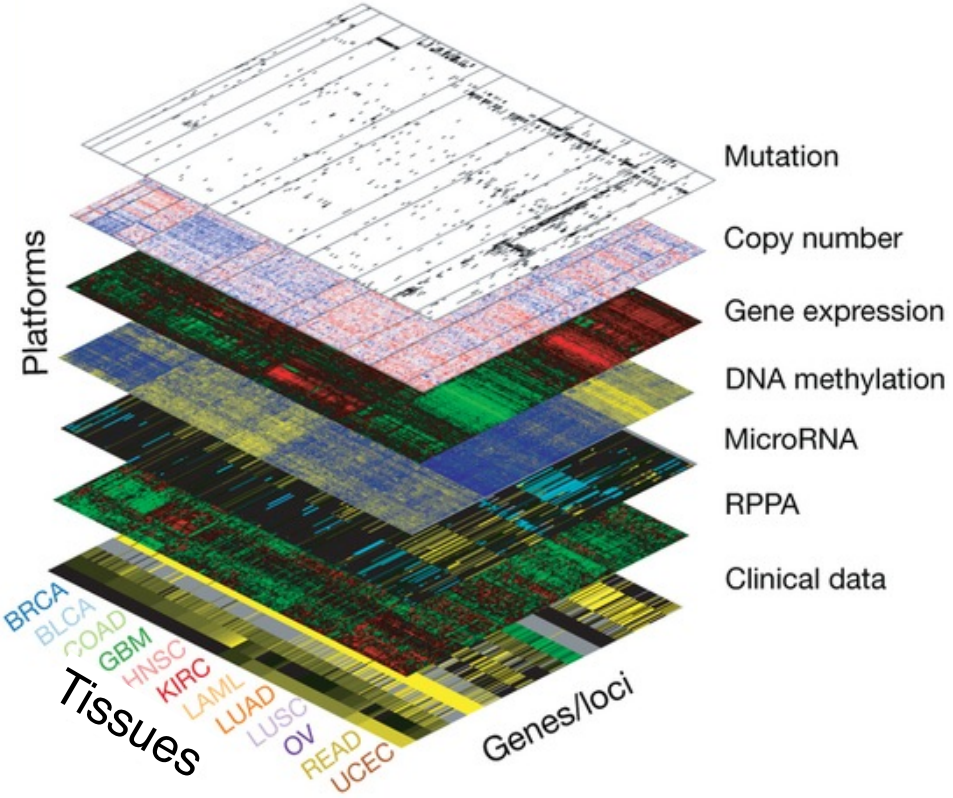
Human genome annotation — a non-intuitive map

geographical information



genomic information

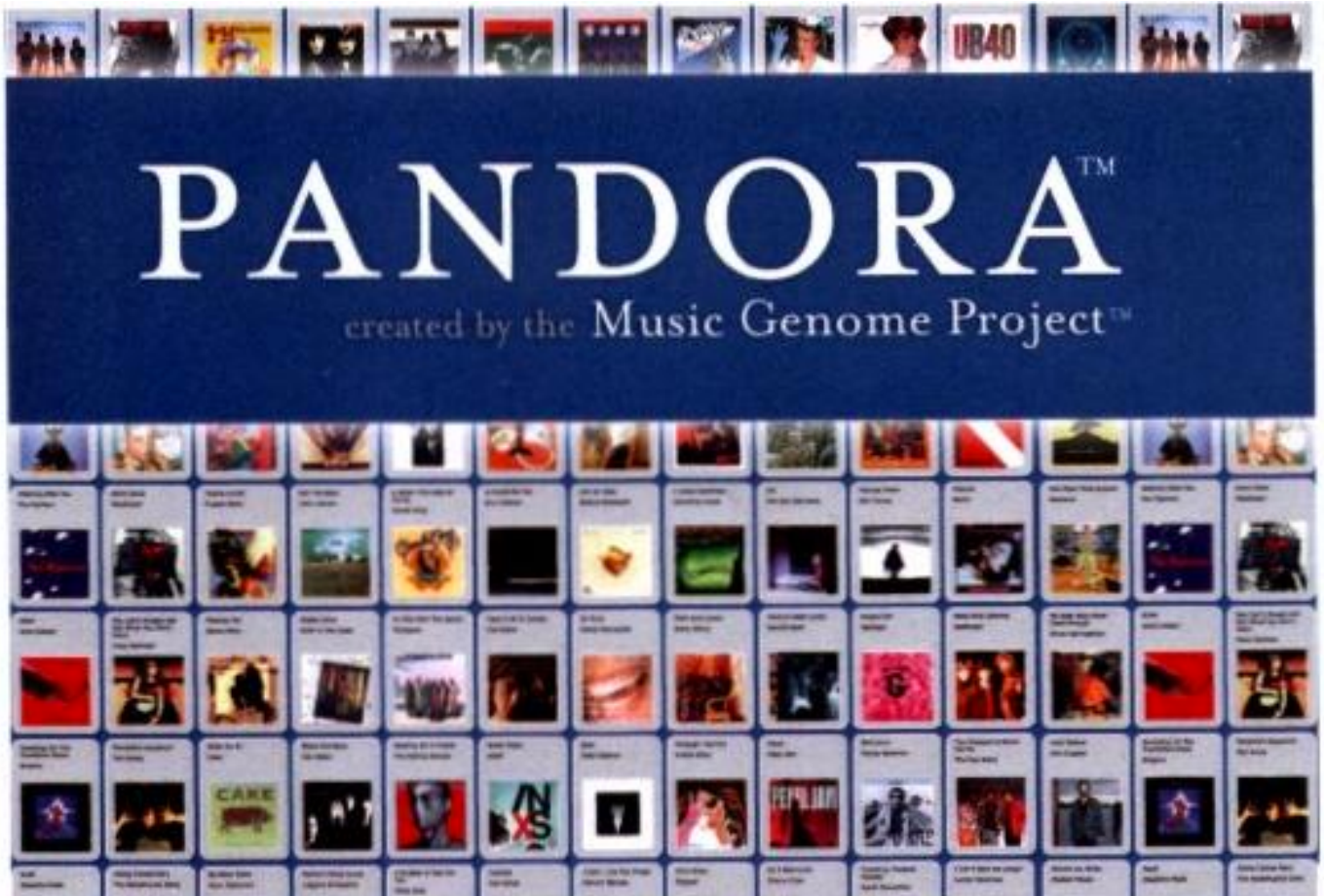




- Large-scale organisation providing an overview of the genome
- Integration of heterogeneous data

Genomics: as an exemplar Data Science sub-discipline

- Developing ways of organizing & mining genomic information on a large scale
  - Very fundamental & early form of "Big Data"
- Perhaps we can learn from other disciplines &, in turn, teach them how to do this?



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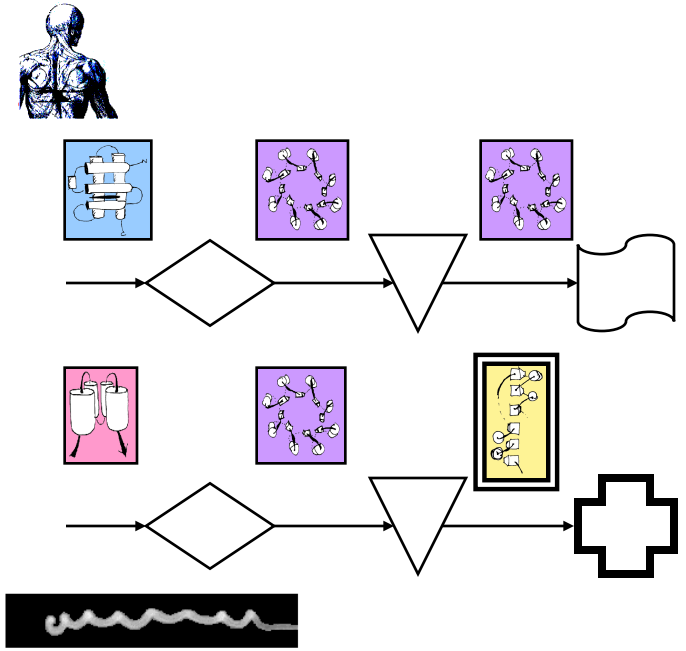
[Luscombe et al. ('01). *Methods Inf Med* 40: 346]

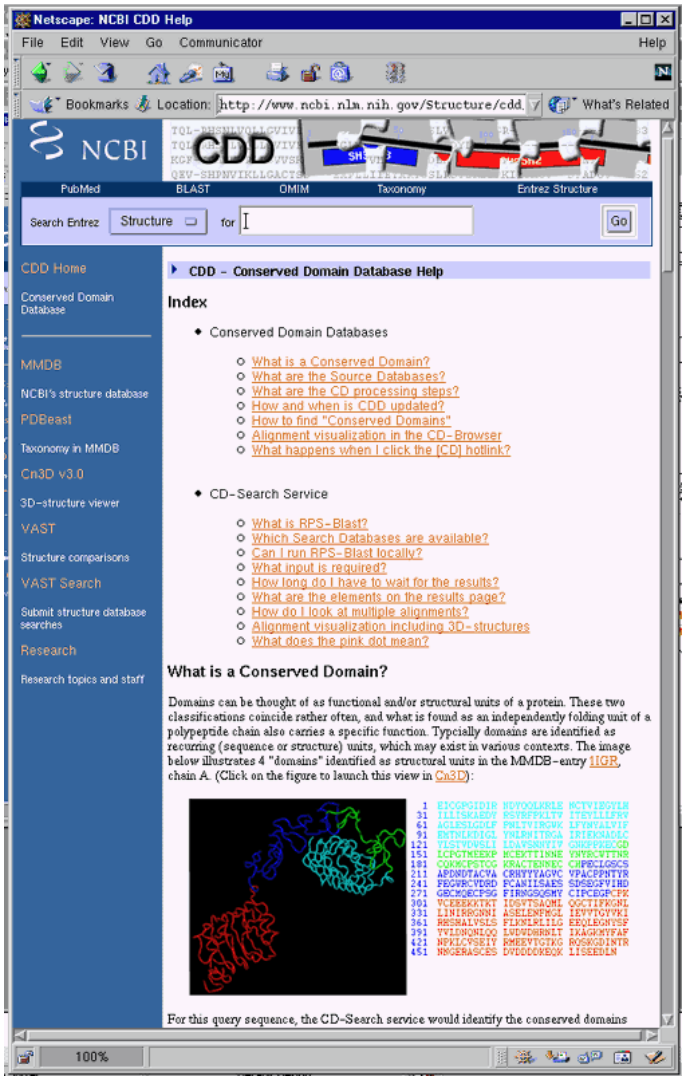
Organizing Molecular Biology Information: Redundancy and Multiplicity

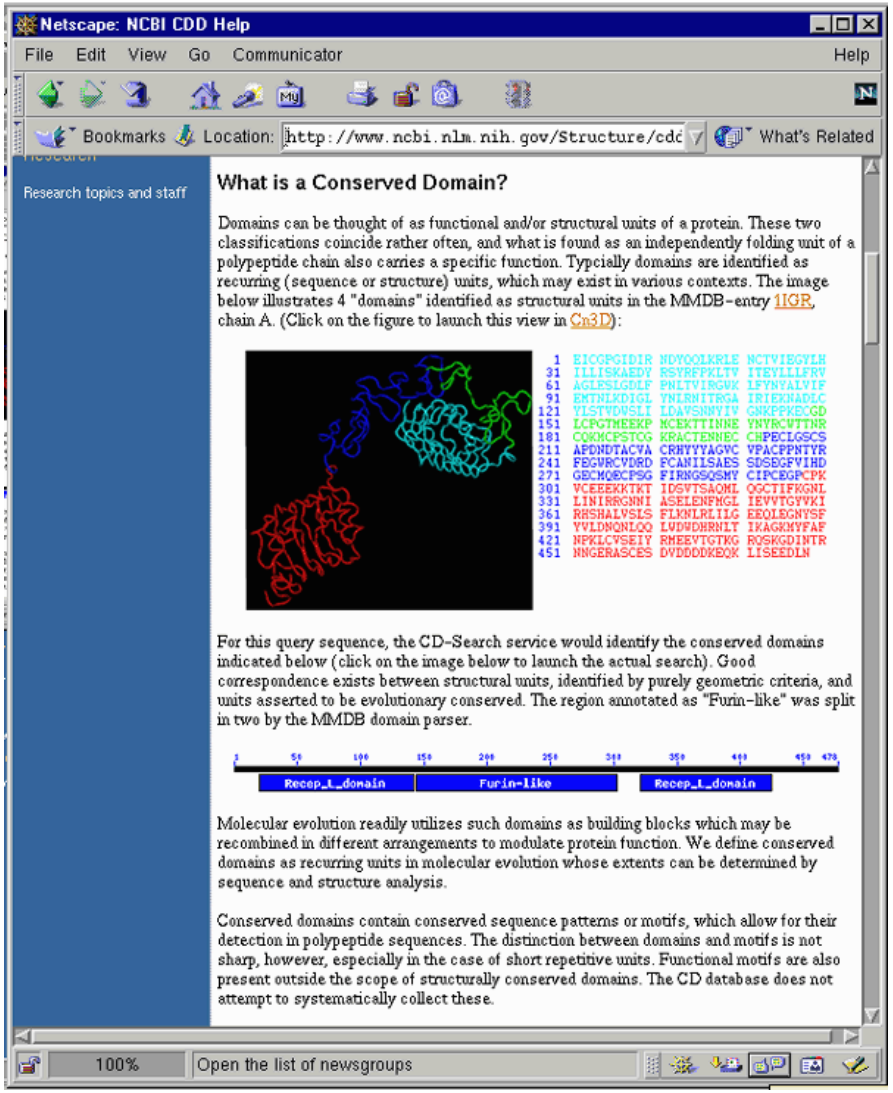


- Different Sequences Have the Same Structure
- Organism has many similar genes
- Single Gene May Have Multiple Functions
- Genes are grouped into Pathway & Networks
- Genomic Sequence Redundancy due to the Genetic Code
- **How do we find the similarities?.....**

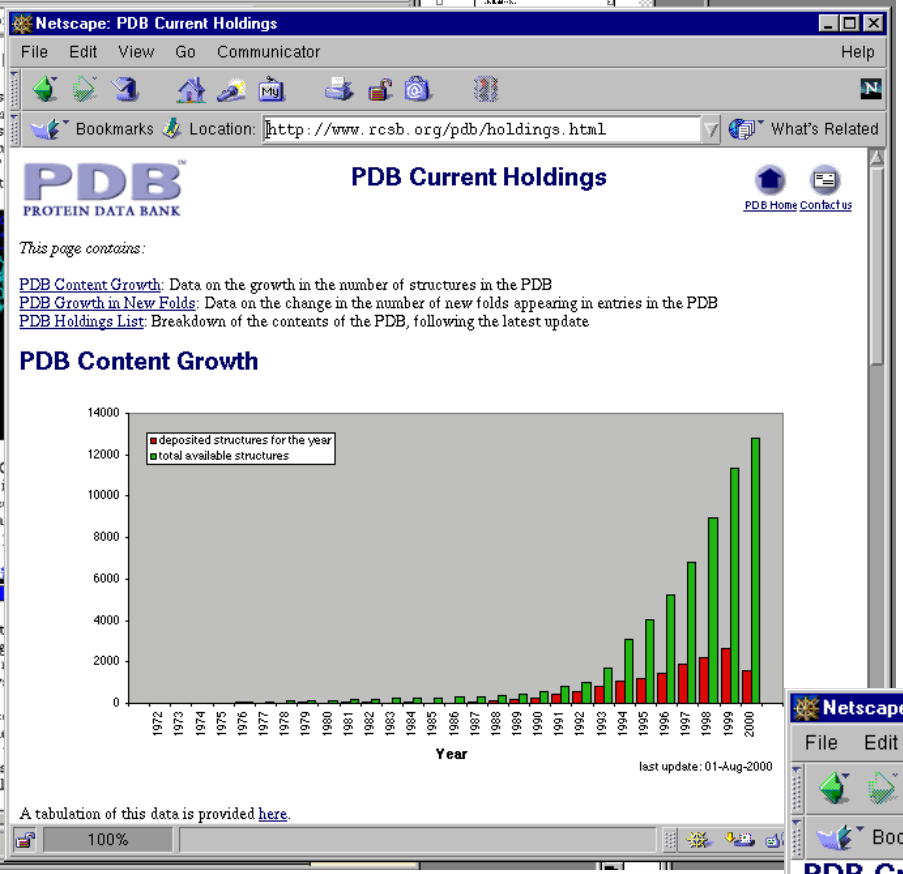
Integrative Genomics - genes ↔ structures ↔ **functions** ↔ **pathways** ↔ expression levels ↔ regulatory systems ↔ ….

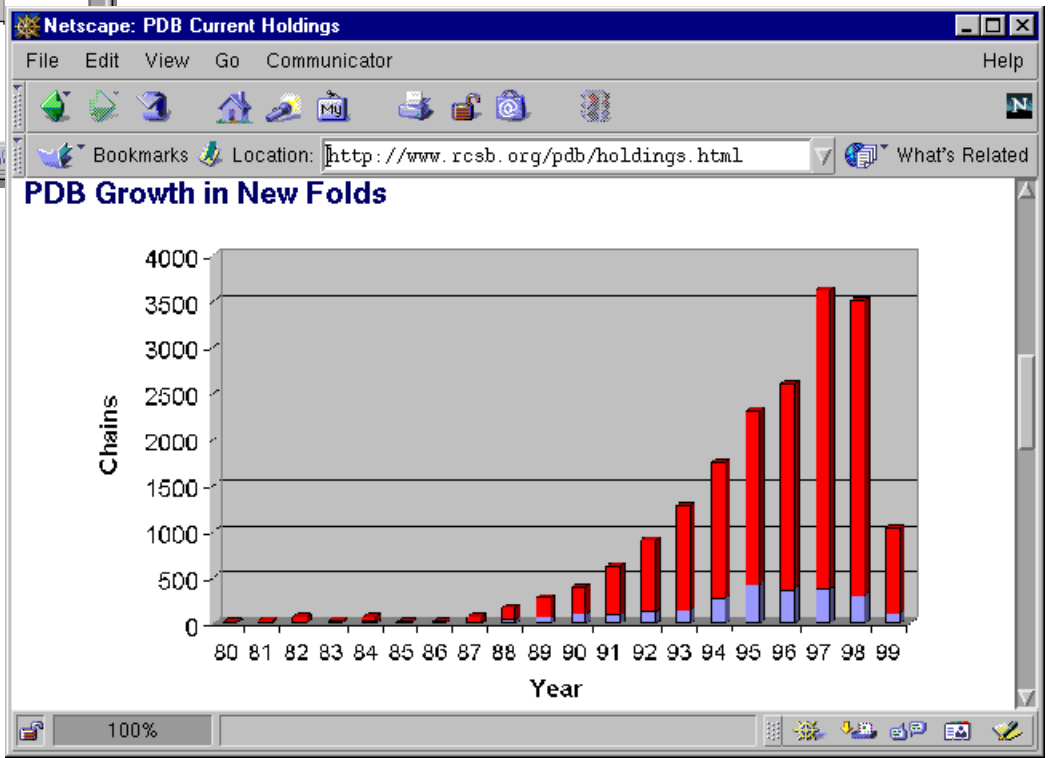
Molecular Parts = Conserved Domains, Folds, &c





Vast Growth in (Structural) Data...
but number of Fundamentally New (Fold) Parts Not Increasing that Fast

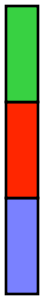




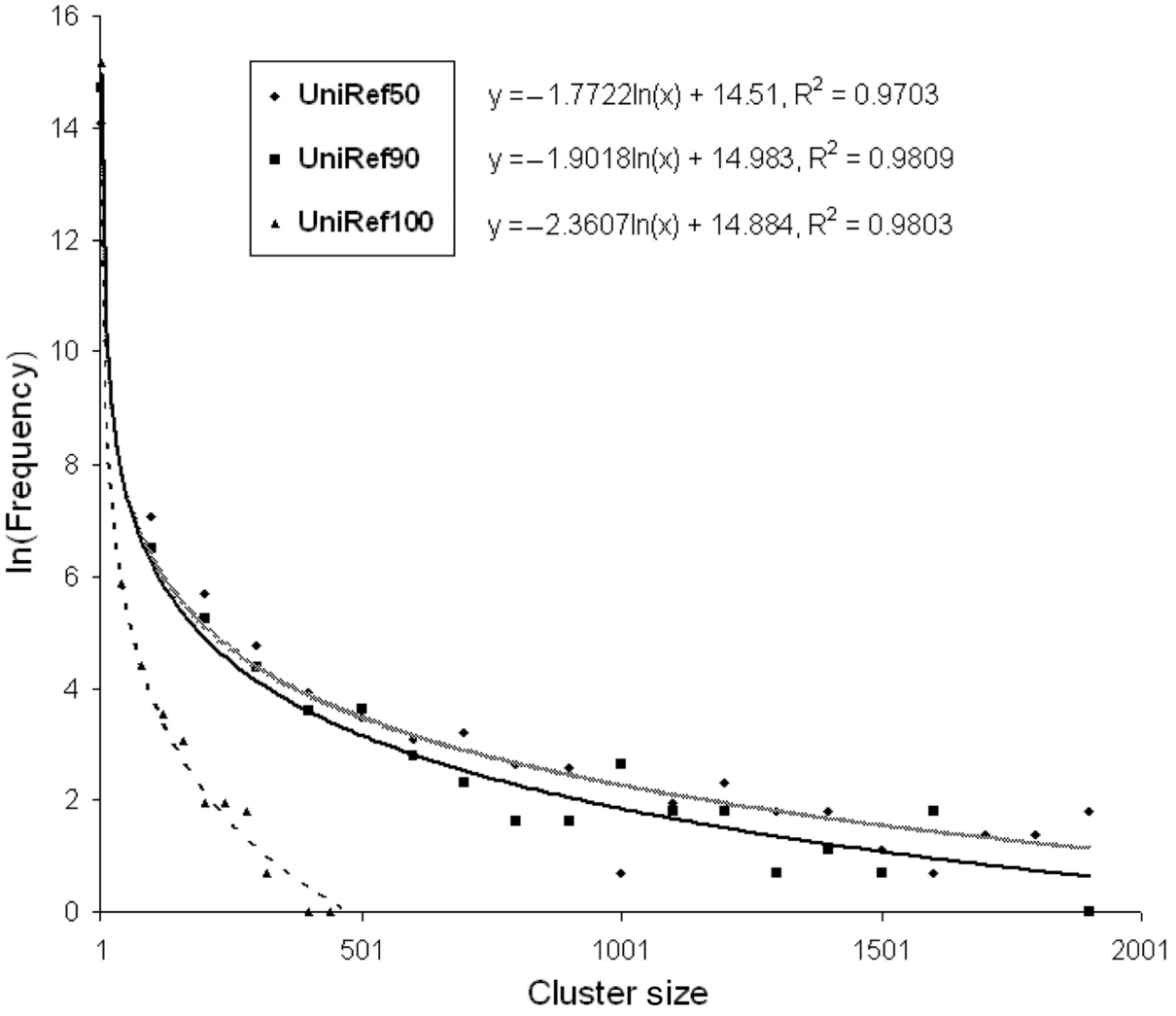
Total in Databank

New Submissions

New Folds



Power-law Size to Protein Families



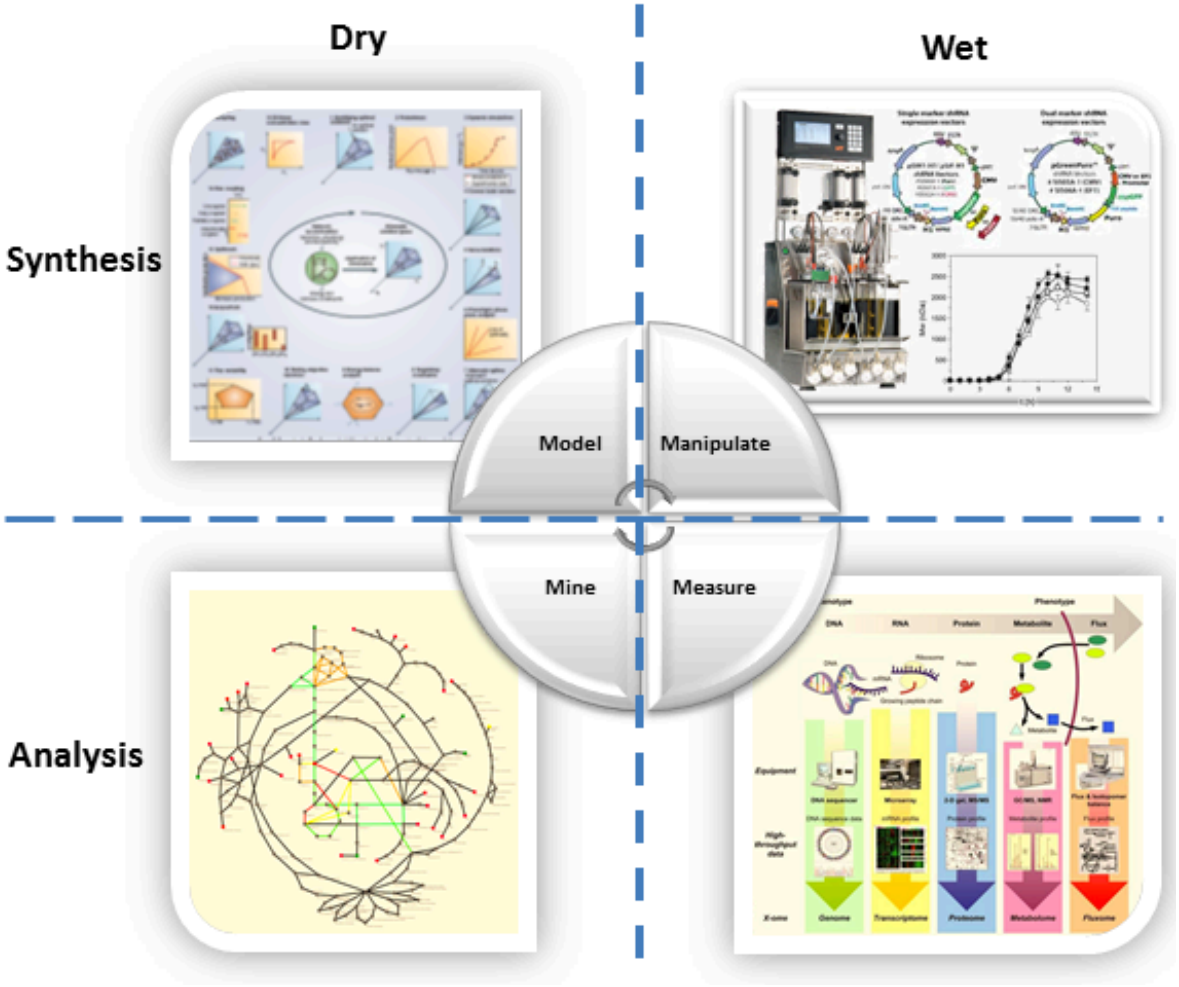
Suzek, B. E. et al. Bioinformatics 2007 23:1282-1288; doi:10.1093/bioinformatics/btm098; See also Luscombe et al., 2002, JMB.

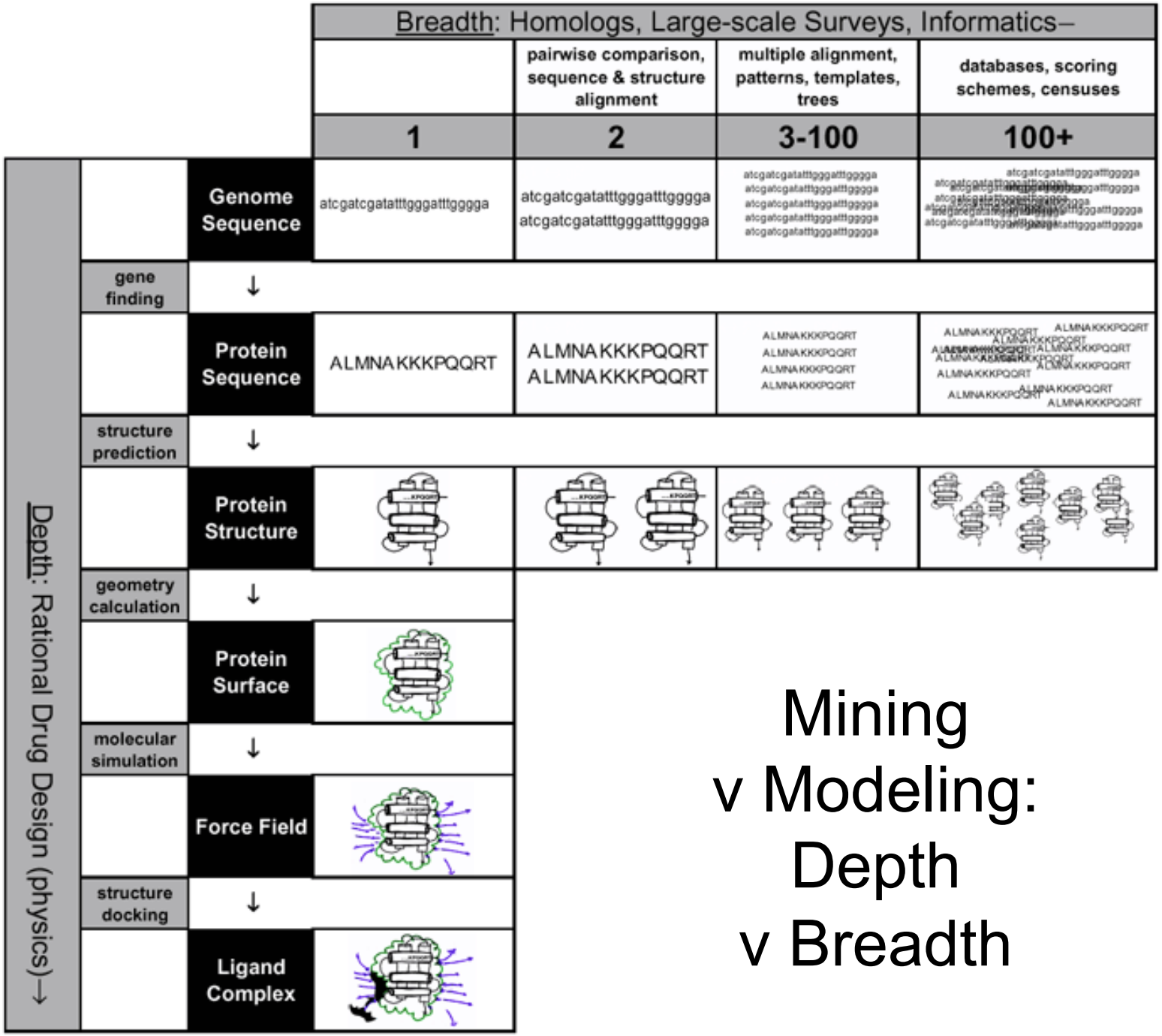
4Ms:

Measurement,
Mining,
Modeling
& Manipulation

TREY IDEKER, L. RAIMOND WINSLOW & A. DOUGLAS LAUFFENBURGER ('06). "Bioengineering and Systems Biology," Annals of Biomedical Engineering DOI: 10.1007/s10439-005-9047-7

Image from http://web.aibn.uq.edu.au/cssb/ResearchProjects.html





Mining
v Modeling:
Depth
v Breadth

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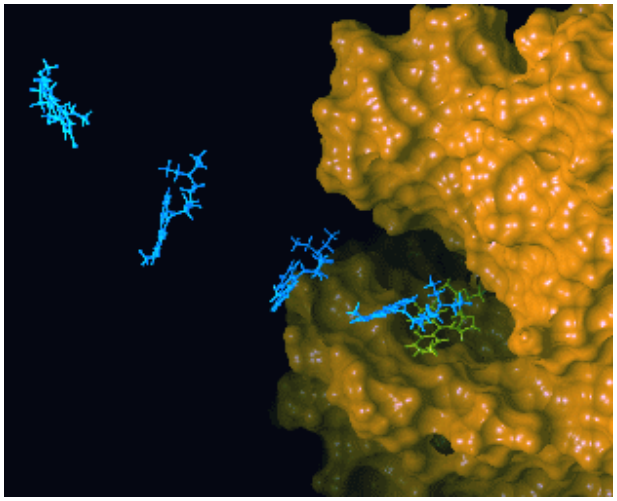
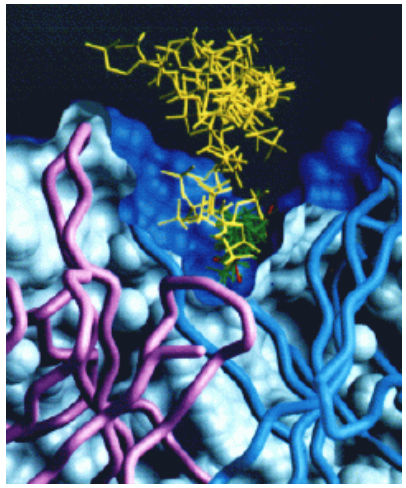
[Luscombe et al. ('01). *Methods Inf Med* 40: 346]

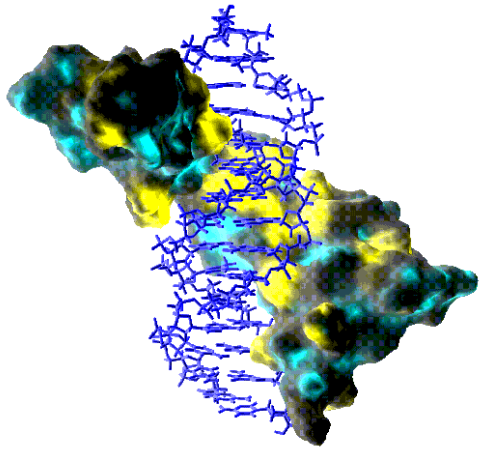
Major Application I: Designing Drugs

- Understanding How Structures Bind Other Molecules (Function)
- Designing Inhibitors
- Docking, Structure Modeling

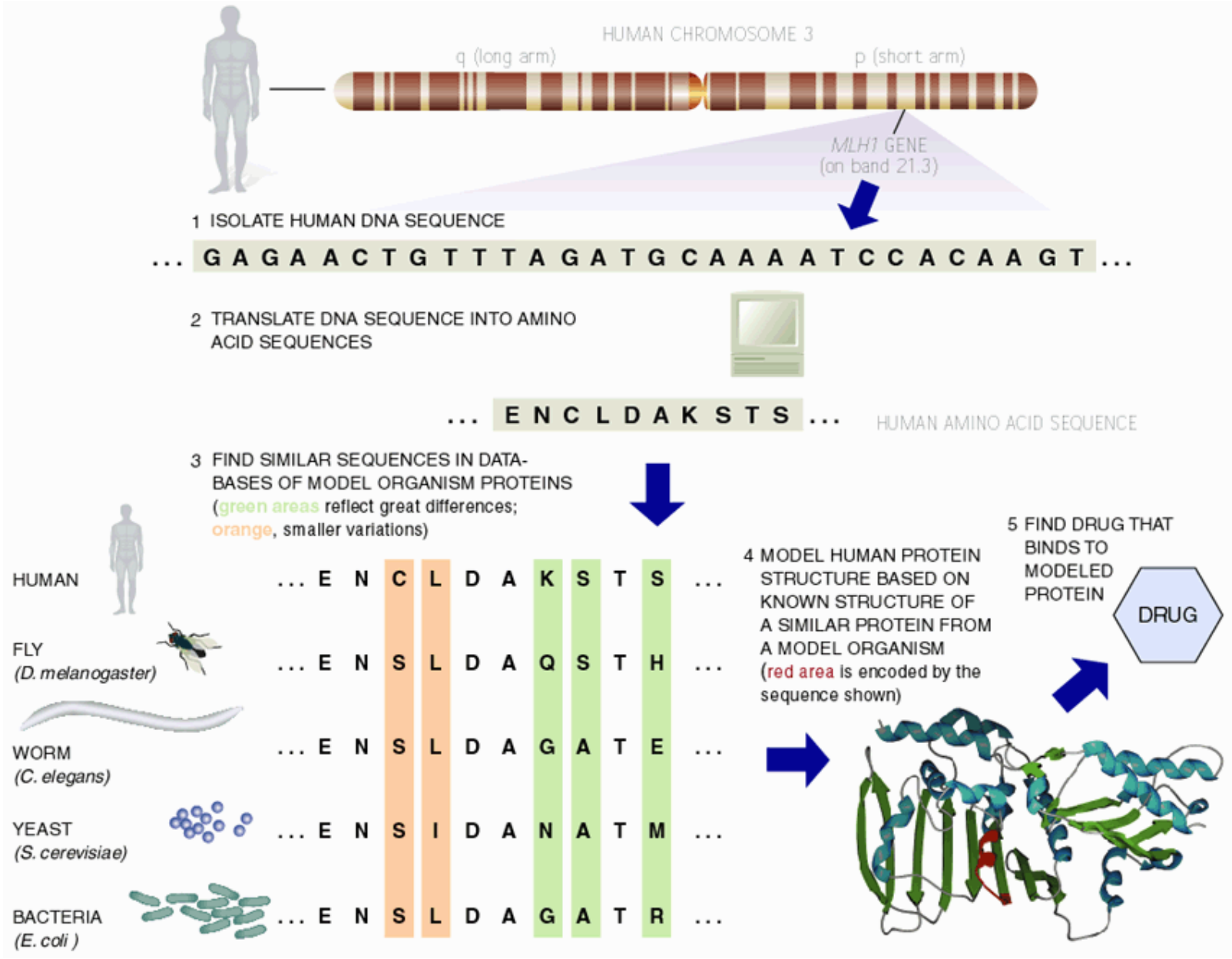
(From left to right, figures adapted from Olsen Group Docking Page at Scripps, Dyson NMR Group Web page at Scripps, and from Computational Chemistry Page at Cornell Theory Center).





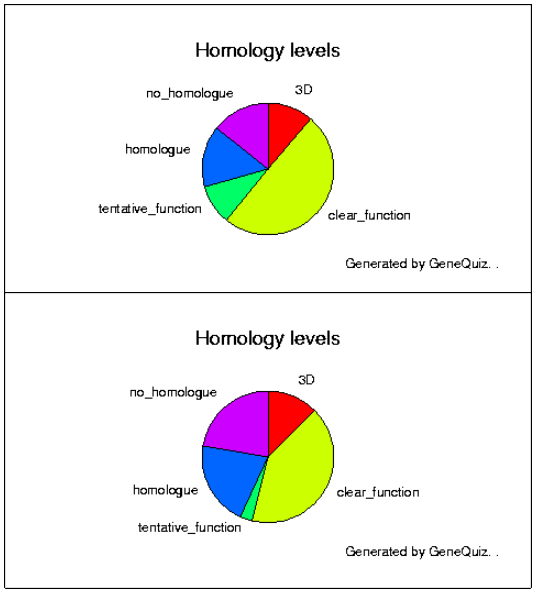


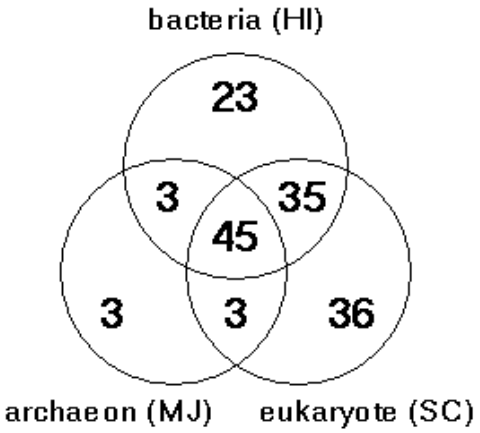
Major Application II: Finding Homologs



Major Application III: Overall Genome Characterization

- Overall Occurrence of a Certain Feature in the Genome
  - ◊ e.g. how many kinases in Yeast
- Compare Organisms and Tissues
  - ◊ Expression levels in Cancerous vs Normal Tissues
- Databases, Statistics
- Using this for **picking drug targets**





(Clock figures, yeast v. Synechocystis,
adapted from GeneQuiz Web Page, Sander Group, EBI)

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Bioinformatics & / or / vs Computational Biology

Biocomputing
Systems Biology
Qbio

Defining the Boundaries of the Field

(Determining the "Support Vectors")

Are They or Aren't They Comp. Bio.? (#1)

- (Digital Libraries & Medical Record Analysis
  - ◊ Automated Bibliographic Search and Textual Comparison
  - ◊ Knowledge bases for biological literature
- (Motif Discovery Using Gibb's Sampling
- (Methods for Structure Determination
  - ◊ Computational Crystallography
    - Refinement
  - ◊ NMR Structure Determination
    - (Distance Geometry
- (Metabolic Pathway Simulation
- (The DNA Computer

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- **(NO)** The DNA Computer

Are They or Aren't They Comp. Bio.? (#2)

- (Gene identification by sequence characteristics
  - ◊ Prediction of splice sites
- (DNA methods in forensics
- (Modeling of Populations of Organisms
  - ◊ Ecological Modeling (predator & prey)
- (Modeling the nervous system
  - ◊ Computational neuroscience
  - ◊ Understanding how brains think & using this to make a better computer
- (Molecular phenotype discovery – looking for gene expression signatures of cancer
  - ◊ What if it included non-molecular data such as age ?

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Are They or Aren't They Comp. Bio.? (#3)

- (RNA structure prediction
- (Radiological Image Processing
  - ◊ Computational Representations for Human Anatomy (visible human)
- (Artificial Life Simulations
  - ◊ Artificial Immunology / Computer Security
  - ◊ (Genetic Algorithms in molecular biology
- (Homology Modeling & Drug Docking
- (Char. drugs & other small molecules (QSAR)
- (Computerized Diagnosis based on Pedigrees
- (Processing of NextGen sequencing image files
- (Module finding in protein networks

Are They or Aren't They Comp. Bio.? (#3, Answers)

- **(YES)** RNA structure prediction
- **(NO)** Radiological Image Processing
  - ◊ Computational Representations for Human Anatomy (visible human)
- **(NO)** Artificial Life Simulations
  - ◊ Artificial Immunology / Computer Security
  - ◊ **(NO?)** Genetic Algorithms in molecular biology
- **(YES)** Homology Modeling & Drug Docking
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Bioinformatics: Practical Application of Simulation and Data Mining

Practical Stuff

http://www.GersteinLab.org/courses/**452**

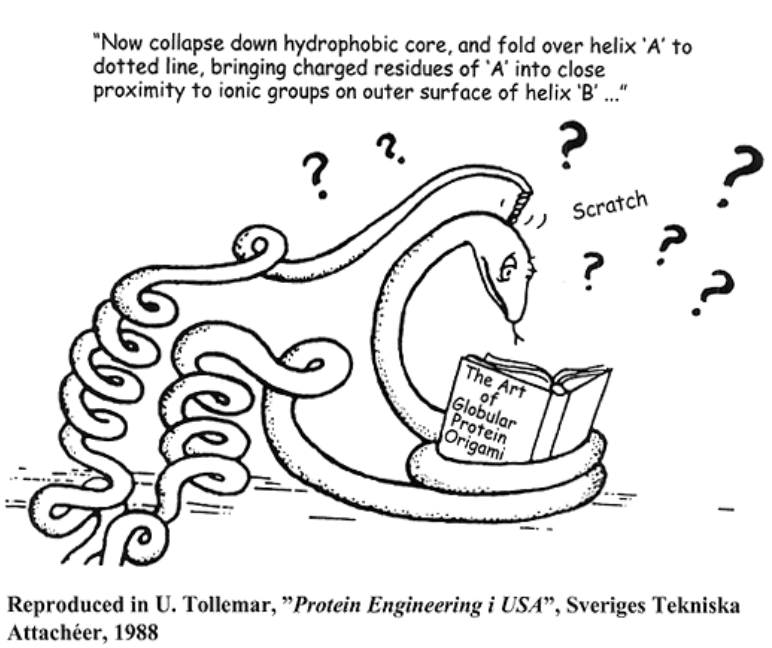
Bioinformatics Topics -- Genome Sequence

- Finding Genes in Genomic DNA
  - ◊ introns
  - ◊ exons
  - ◊ promotors
- Characterizing Repeats in Genomic DNA
  - ◊ Statistics
  - ◊ Patterns
- Duplications in the Genome
  - ◊ Large scale genomic alignment
- Whole-Genome Comparisons
- Finding Structural RNAs

Bioinformatics Topics -- Protein Sequence

- Sequence Alignment
  - ◊ non-exact string matching, gaps
  - ◊ How to align two strings optimally via Dynamic Programming
  - ◊ Local vs Global Alignment
  - ◊ Suboptimal Alignment
  - ◊ Hashing to increase speed (BLAST, FASTA)
  - ◊ Amino acid substitution scoring matrices
- Multiple Alignment and Consensus Patterns
  - ◊ How to align more than one sequence and then fuse the result in a consensus representation
  - ◊ Transitive Comparisons
  - ◊ HMMs, Profiles
  - ◊ Motifs
- Scoring schemes and Matching statistics
  - ◊ How to tell if a given alignment or match is statistically significant
  - ◊ A P-value (or an e-value)?
  - ◊ Score Distributions (extreme val. dist.)
  - ◊ Low Complexity Sequences
- Evolutionary Issues
  - ◊ Rates of mutation and change

Bioinformatics Topics -- Sequence / Structure



Reproduced in U. Tollemar, "*Protein Engineering i USA*", Sveriges Tekniska Attachéer, 1988

- Secondary Structure "Prediction"
  - ◊ via Propensities
  - ◊ \TM-helix finding
  - ◊ Assessing Secondary Structure Prediction
- Structure Prediction: Protein v RNA
- Tertiary Structure Prediction
  - ◊ Fold Recognition
  - ◊ Threading
  - ◊ Ab initio
  - ◊ (Quaternary structure prediction)
- Direct Function Prediction
  - ◊ Active site identification
- Relation of Sequence Similarity to Structural Similarity

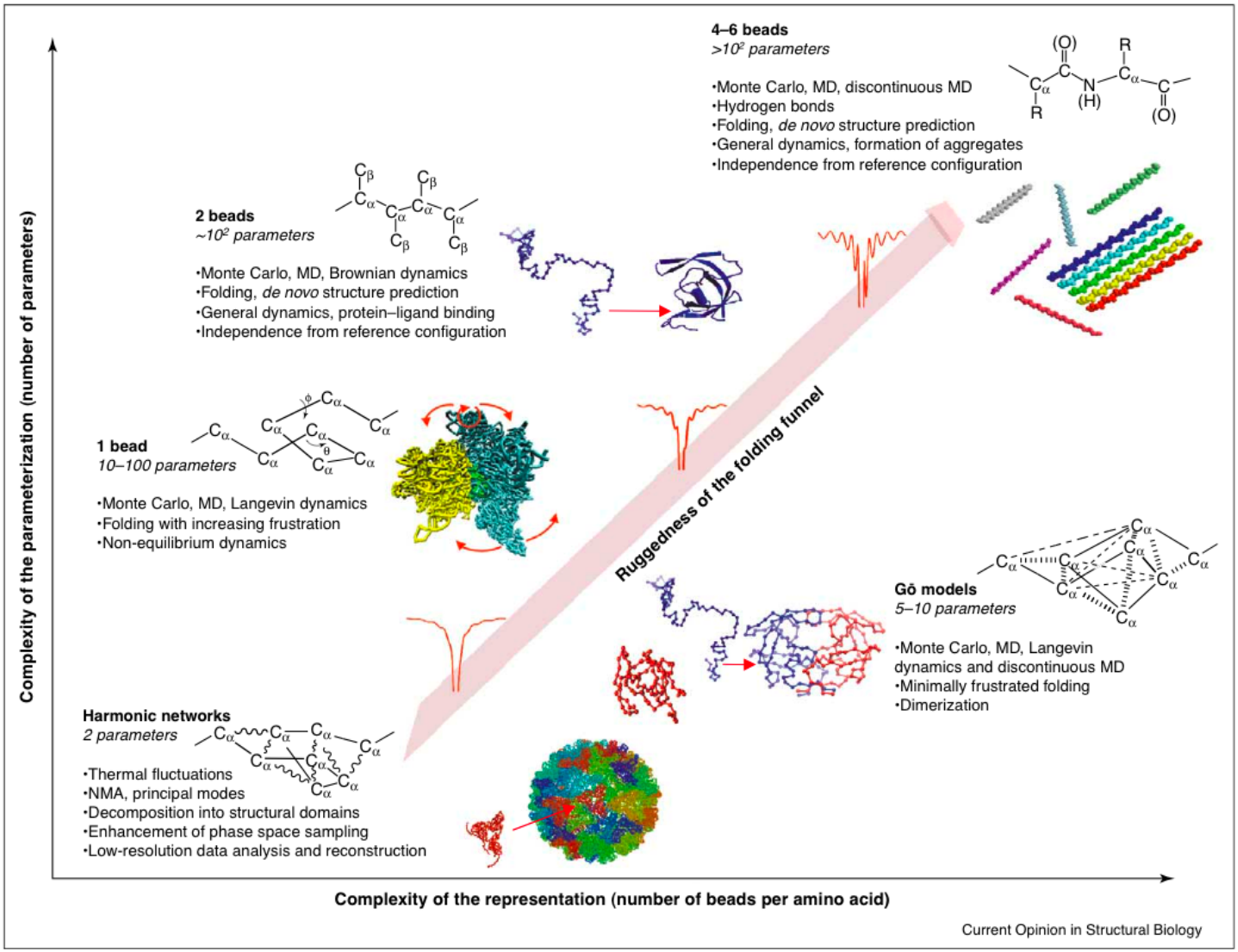
Topics -- Structures

- Structure Comparison
  - ◊ Basic Protein Geometry and Least-Squares Fitting
- Distances, Angles, Axes, Rotations
  - ◊ Calculating a helix axis in 3D via fitting a line
  - ◊ LSQ fit of 2 structures
  - ◊ Molecular Graphics
- Calculation of Volume and Surface
  - ◊ Hinge prediction
  - ◊ Packing Measurement
- Structural Alignment
  - ◊ Aligning sequences on the basis of 3D structure.
  - ◊ DP does not converge, unlike sequences, what to do?
  - ◊ Other Approaches: Distance Matrices, Hashing
- Fold Library
- Docking and Drug Design as Surface Matching

Topics -- Simulation

- Molecular Simulation
  - ◊ Geometry –> Energy –> Forces
  - ◊ Basic interactions, potential energy functions
  - ◊ Electrostatics
  - ◊ VDW Forces
  - ◊ Bonds as Springs
  - ◊ How structure changes over time?
    - How to measure the change in a vector (gradient)
  - ◊ Molecular Dynamics & MC
  - ◊ Energy Minimization
- Parameter Sets
- Number Density
- Simplifications
  - ◊ Poisson-Boltzman Equation
  - ◊ Lattice Models and Simplification

Figure 1



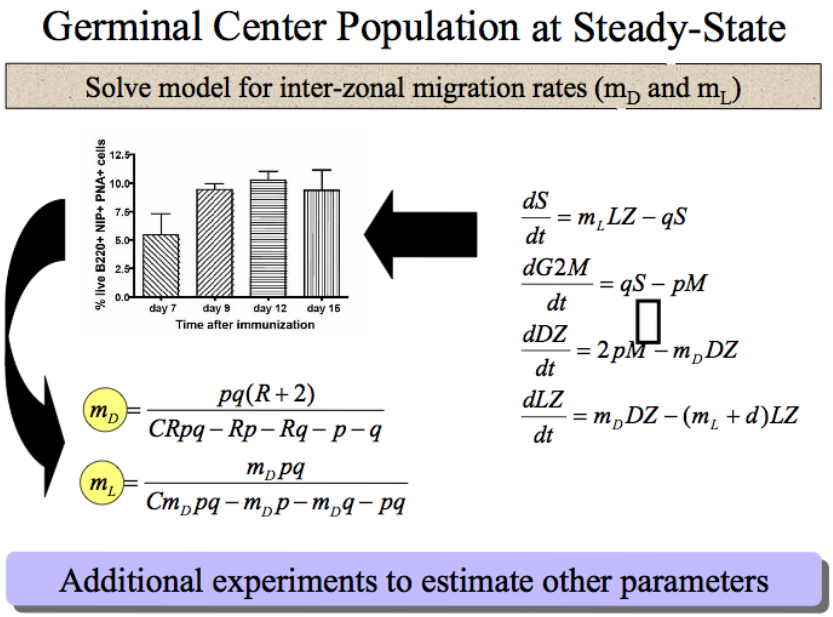
Pictorial representation of the features of bead models. For each class of model, the following aspects are reported: schematic representation of the model, indicative number of parameters, methods of solution, main characteristics and applications. Sample applications are also illustrated with representative pictures (prepared using crystallographic coordinates from the PDB [codes 1hhp, 1cwp, 1mwr, 486d]) intended to show the size of system that can be studied and the kind of study that can be done. The location of the models in the *x-y* plane is intended to qualitatively illustrate their complexity, which increases following the direction of the arrows.

Journal Articles

1. J. D. Honeycutt and D. Thirumalai, “The nature of folded states of globular proteins,” *Biopolymers* **32** (1992) 695.
2. W. C. Swope and J. W. Pitera, “Describing protein folding kinetics By molecular dynamics simulations. 1. Theory,” *J. Phys. Chem. B* **108** (2004) 6571.
3. D. Bratko, T. Cellmer, J. M. Prausnitz, and H. W. Blanch, “Molecular Simulation of protein aggregation,” *Biotechnology and Bioengineering* **96** (2007) 1.

Cell and Immunology Simulations
Prof. Steven Kleinstein

- Modeling cell mutation, division and death
- Population dynamics using ODEs
- Viral dynamics and immunological response
- Optimization and matching experimental data



Next Generation Sequencing & Big Data

- Seq. Tech
- Assembly
- RNA-seq
- ChIP-seq
- Metagenomics

Topics – (Func) Genomics

- Expression Analysis
  - Time Courses clustering
  - Measuring differences
  - Identifying Regulatory Regions
  - Normalization and scoring of arrays
- Function Classification and Orthologs
- Genome Comparisons
  - Large-scale censuses
  - Frequent Words Analysis
  - Genome Annotation
  - Identification of interacting proteins
- Structural Genomics
  - Folds in Genomes, shared & common folds
  - Bulk Structure Prediction

Topics – DBs/Surveys

- Relational Database Concepts and how they interface with Biological Information

- DB interoperation
- What are the Units ?
  - What are the units of biological information for organization?

- Clustering & Trees
  - Basic clustering
  - Evolutionary implications
- Visualization of Large Amounts of Information

Mining

- Information integration and fusion
  - Dealing with heterogeneous data
- Dimensionality Reduction (PCA etc)
- Networks
  - Topology Analysis
  - Prediction of linkages
  - Global structure and local motifs