

Visualization and Biology: Fertile Ground for Collaboration

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<http://www.cs.ubc.ca/~tmm/talks.html#harvard09>

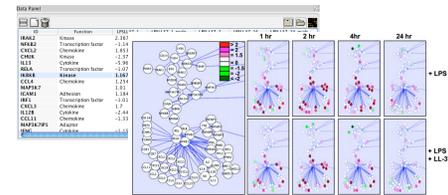
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Outline

- visualization ideas and background
- combining interaction networks, microarray data
 - Cerebral system
- comparing phylogenetic trees
 - TreeJuxtaposer system
- discussion

Why do visualization?

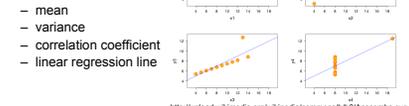
- pictures help us think
 - substitute perception for cognition
 - external memory: free up limited cognitive/memory resources for higher-level problems



When should we bother doing vis?

- need a human in the loop
 - augment, not replace, human cognition
 - for problems that cannot be (completely) automated
- simple summary not adequate
 - statistics may not adequately characterize complexity of dataset distribution

Anscombe's quartet:
same



- mean
- variance
- correlation coefficient
- linear regression line

<http://upload.wikimedia.org/wikipedia/commons/b/b6/Anscombe.svg>

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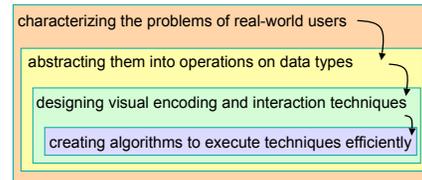
What does visualization allow?

- discovering new things
 - hypothesis discovery, "eureka moment"
- confirming conjectured things
 - hypothesis confirmation
- contradicting conjectured things
 - especially (inevitably?) data cleansing
- novel capabilities
 - tool supports fundamentally new operations
- **speedup**
 - tool accelerates workflow (most common)

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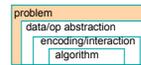
Multiple levels of problem-driven vis

- cascading levels: output above is input below



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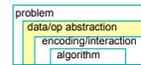
Characterizing problems



- understanding domain concepts and current workflow
- finding gaps, breakdowns, slowdowns
 - where conjecture that vis would help

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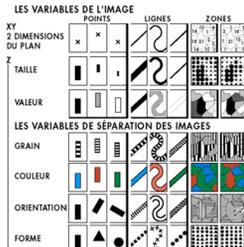
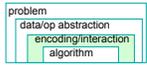
Abstracting into operations on data types



- data types
 - tables of numbers
 - relations: networks/graphs, hierarchies/trees
 - spatial data: geographic, positions in space
- operations
 - sorting, filtering, browsing, comparison, characterizing trends and distributions, finding anomalies and outliers, finding correlation...
 - relations: following path through network...

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Designing encoding and interaction

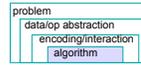


Semiology of Graphics. Jacques Bertin, Gauthier-Villars 1967, EHESS 1999

- visual encoding
 - marks: points, lines, areas
 - attributes: position, color, shape, size, orientation, ...
- interaction
 - selecting, navigating, ordering, ...

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Creating efficient algorithms



- classic computer science problem
 - create algorithm given clear specification

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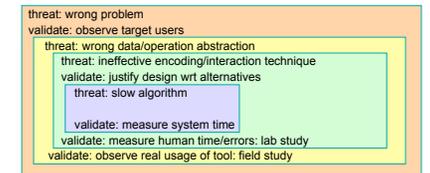
Design decisions

- huge space of design alternatives
 - conflicting tradeoffs
 - iterative refinement often necessary
- many/most choices are ineffective
 - wrong visual encoding can mislead, confuse
 - principled reasons to make choices usually not obvious to untrained people

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Validation: Is problem solved?

- humans in the loop for outer three levels



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Collaboration: Complementary expertise

- vis researchers
 - vis design alternatives
 - human perceptual capabilities
 - scalable graphics algorithms
 - validation methodology
- domain scientists
 - deep knowledge of driving problems, data
- both benefit from new tools
 - scientist: you get something helpful
 - vis researcher: we get to watch you use it
 - see if problem actually solved
 - feed new knowledge back into our design principles

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Good driving problems for vis research

- big data
- reasonably clear questions
- need for humans in the loop
- many areas of science are a great match
 - biology particularly appealing

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Cerebral

collaboration with researchers at UBC Hancock Lab studying innate immunity

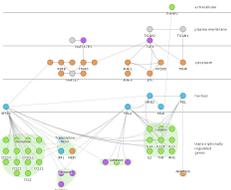
Cerebral: Visualizing Multiple Experimental Conditions on a Graph with Biological Context
Aaron Barsky, Computer Science, UBC
Tamara Munzner, Computer Science, UBC
Jennifer Gardy, Microbiology and Immunology, UBC
Robert Kincaid, Agilent Technologies
IEEE Transactions on Visualization and Computer Graphics (Proc. InfoVis 2008) 14(6) (Nov-Dec) 2008, p 1255-1260
<http://www.cs.ubc.ca/labs/imager/tr/2008/cerebral/>
<http://www.cs.ubc.ca/labs/imager/tr/2008/BarskyMscThesis/>

open-source software download (Cytoscape plugin)
<http://www.pathogenomics.ca/cerebral/>
deployed in InnateDB (mammalian innate immunity database)
<http://www.innatedb.ca>

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Systems biology model

- graph $G = \{V, E\}$
 - V: proteins, genes, DNA, RNA, tRNA, etc.
 - E: interacting molecules



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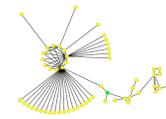
Model - Experiment cycle

- conduct experiments on cells
 - interpret results in current graph model
 - propose modifications to refine model
- vis tool to accelerate workflow?

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Goal: Integrate model with measurements

- system model
 - interaction graph $G = \{V, E\}$
 - meta-data for each v in V
 - labels, biological attributes
- experimental measurements
 - multiple floats for each v in V
 - microarray data

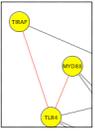


Label	Protein	LPS1L7_1	LPS1L7_1_1	LPS1L7_2	LPS1L7_24	LPS1L7_24_1
MAP2	Microtubule	2.867	0.251	1.537	-1.513	0.803
MAP2K2	Transcription factor	-1.14	0.672	-1.03	1.393	0.803
CELSF2	Chaperone	3.653	0.176	4.113	1.819	0.763
CYR61	Chaperone	-1.176	0.177	2.212	1.196	0.807
SLC4	Chaperone	-1.265	0.166	2.188	2.128	0.802
MAP2K1	Chaperone	2.214	0.166	-1.096	1.071	0.76
MAPK8	Kinase	3.847	0.28	3.482	3.887	0.286
MAPK9	Kinase	0.67	0.166	-1.096	1.071	0.76
MAPK1	Adhesion	1.184	0.166	1.137	1.102	0.811
MAPK10	Transcription factor	-1.613	0.166	1.146	1.081	0.802
CELSF3	Chaperone	1.1	0.166	1.192	1.188	0.521
SLC18	Chaperone	-2.648	0.166	-1.673	-2.109	0.8
CELSF1	Chaperone	-1.138	0.166	-1.995	-1.793	0.229
MAP2K10P1	Chaperone	-1.138	0.166	-1.995	-1.793	0.229

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Model summarizes extensive lab work

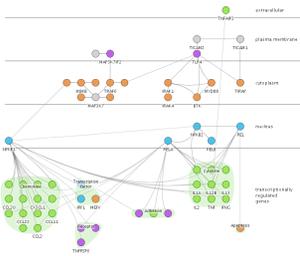
- graphs come from hand-curated databases
 - dynamic, change with each new publication
- each edge has provenance from experimental evidence
 - TIRAP: an adapter molecule in the Toll signaling pathway. Hong T, Barton GM, Medzhitov R.
 - Mai (MyD88 adapter-like) is required for Toll-like receptor-4 signal transduction. Fitzgerald KA, Palsson-McDermott EM, Bowie AG, Jefferson CA, Manless AS, Brady G, Brint E, Dunne A, Gray P, Harte MT, McMurray D, Smith DE, Sims JE, Bird TA, O'Neill LA.
- choose scope to manage complexity



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TLR4 biomolecule: E=74, V=54

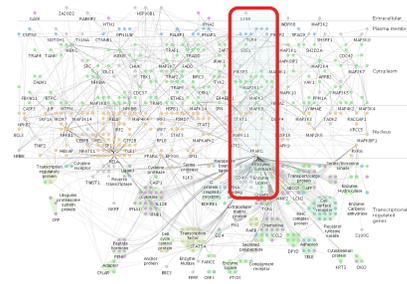
- very local view



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Immune system: E=1263, V=760

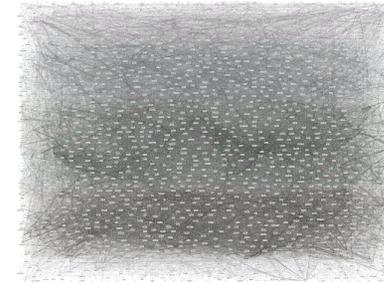
- bigger picture, target size for Cerebral



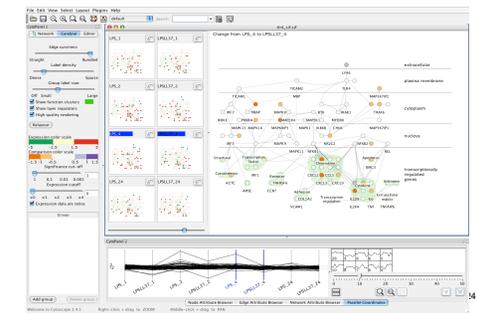
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Human interactome: E~50,000, V~10,000

- too complex, beyond scope of tool



Cerebral video



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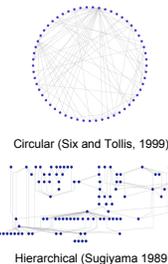
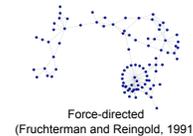
Encoding and interaction design decisions

- create custom graph layout
 - guided by biological metadata
- use small multiple views
 - one view per experimental condition
- show measured data in graph context
 - not in isolation

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Traditional graph layout

- given graph $G=\{V,E\}$
- create layout in 2D plane
- heavily studied
 - hundreds of papers
 - annual Graph Drawing conf



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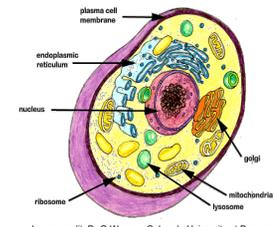
Existing layouts did not suit immunologists

- graph drawing goals
 - visualize graph structure
- biologist goals
 - visualize biological knowledge
 - some relationships happen to form a graph
 - cell location also relevant

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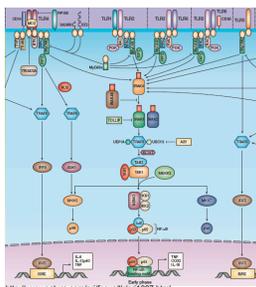
Biological cells divided by membranes

- interactions generally occur within a compartment
- crossing membranes is interesting



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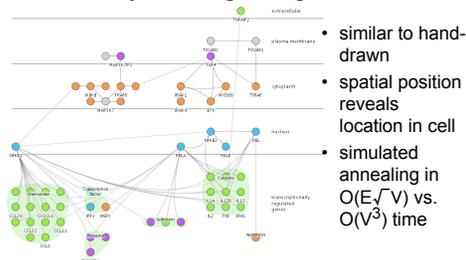
Hand-drawn diagrams



- cellular location encoded spatially
- infeasible to create by hand in era of big data

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Cerebral layout using biological metadata

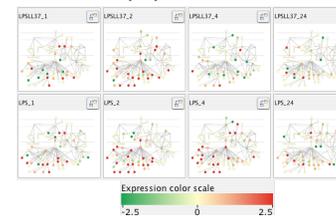


- similar to hand-drawn
- spatial position reveals location in cell
- simulated annealing in $O(E\sqrt{V})$ vs. $O(V^3)$ time

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Use small multiple views

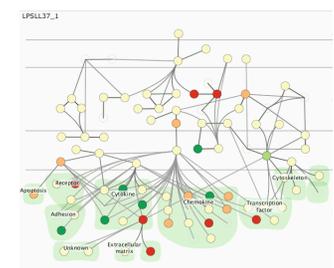
- one graph instance per experimental condition
 - same spatial layout
 - color differently, by condition



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Why not animation?

- global comparison difficult



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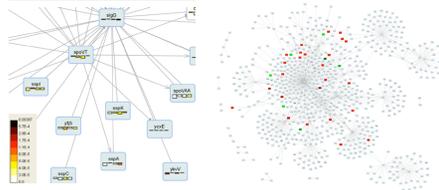
Why not animation?

- limits of human visual memory
 - compared to side by side visual comparison
- Matthew Plumlee and Colin Ware. Zooming versus multiple window interfaces: Cognitive costs of visual comparisons. *ACM Trans. Computer-Human Interaction (TOCHI)*, 13(2):179-209, 2006.
- Barbara Tversky, Julie Bauer Morrison, and Mireille Bejrancourt. Animation: can it facilitate? *International Journal of Human-Computer Studies*, 57(4):247-262, 2002.

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Why not glyphs?

- embed multiple conditions as a chart inside node
- clearly visible when zoomed in
- but cannot see from global view
 - only one value shown in overview

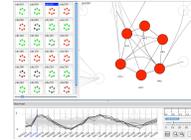


[M. A. Weisenberg, S. A. F. T. van Hün, O. P. Kuipers, J. B. T. M. Roerdink. Visualizing Genome Expression and Regulatory 34 Network Dynamics in Genomic and Metabolic Context. *Computer Graphics Forum*, 27(3):887-894, 2008.]

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Show measured data in graph context

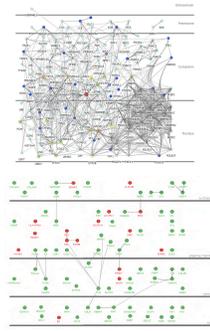
- data driven hypothesis
 - clusters indicate similar function?
 - same pattern of gene expression → same role in cell?
- clusters are often untrustworthy artifacts!
 - noisy data: different clustering alg. → different results
 - measured data alone potentially misleading
 - show in context of graph model



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Adoption by biologists

- Matthew D Dyer, T. M Murali, and Bruno W Sobral. The landscape of human proteins interacting with viruses and other pathogens. *PLoS Pathogens*, 4(2):e32, 2008.
- Liqui He et al. The glomerular transcriptome and a predicted protein-protein interaction network. *Journal of the American Society of Nephrology*, 19(2):260-268, 2008.



InnateDB links to Cerebral

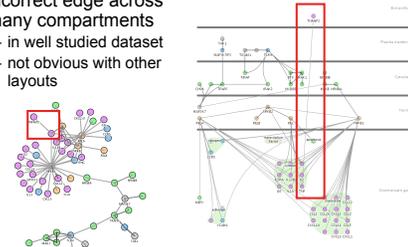
- InnateDB: facilitating systems-level analyses of the mammalian innate immune response
 - David J Lynn, Geoffrey L Winsor, Calvin Chan, Nicolas Richard, Matthew R Laird, Aaron Barsky, Jennifer L Gardy, Fiona M Roche, Timothy H W Chan, Naisha Shah, Raymond Lo, Misbah Naseer, Jiammie Que, Melissa Yau, Michael Acab, Dan Tulpan, Matthew D Whiteside, Avinash Chikamaria, Bernadette Mah, Tamara Munzner, Karsten Hokamp, Robert E W Hancock, Fiona S L Brinkman. *Molecular Systems Biology* 2008; 4:218
 - <http://innatedb.ca>

Gene	Accession	Species	Pathway	Function	Location	Expression	Interactions
CD14	U05389	Human	CD14	CD14	CD14	CD14	CD14
CD14	U05389	Human	CD14	CD14	CD14	CD14	CD14
CD14	U05389	Human	CD14	CD14	CD14	CD14	CD14

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Data cleansing example

- incorrect edge across many compartments
 - in well studied dataset
 - not obvious with other layouts



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Cerebral summary

- supports interactive exploration of multiple experimental conditions in graph context
- provides familiar representation by using biological metadata to guide graph layout

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 - Cerebral system
- comparing phylogenetic trees
 - TreeJuxtaposer system
- discussion

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TreeJuxtaposer

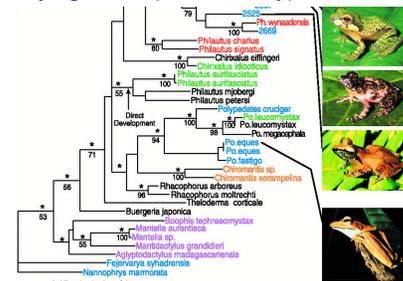
collaboration with biologists at UT-Austin Hillis Lab

TreeJuxtaposer: Scalable Tree Comparison using Focus+Context with Guaranteed Visibility.
 Tamara Munzner, François Guimbertière, Serdar Tasiran, Li Zhang, Yunhong Zhou.
ACM Trans. Graphics 22(3): 453-462, 2003 (Proc. SIGGRAPH 2003).
<http://www.cs.ubc.ca/labs/imager/tr/2003/tj>

open-source software download
<http://olduvai.sourceforge.net/tj>

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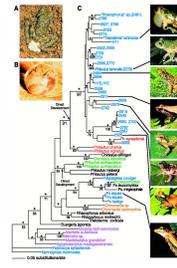
Phylogenetic (evolutionary) tree



M Meegaskumbura et al., *Science* 298:379 (2002)

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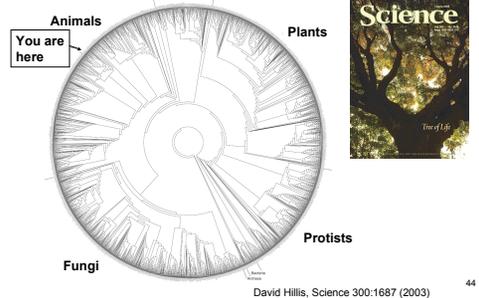
Common dataset size today



M Meegaskumbura et al., *Science* 298:379 (2002)

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Future goal: Full Tree of Life, ~10M nodes



David Hillis, *Science* 300:1687 (2003)

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Operation: Comparing multiple trees

- presentation: single tree shown as final result
- exploration: determine true tree from many possibilities
 - different biological conjectures or data
 - different phylogenetic reconstruction algorithms
 - multiple alternatives from same reconstruction algorithm
- most previous work on browsing
 - necessary but not sufficient for comparison

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Limitations of paper: Scale and speed

- literal: actual paper
- figurative: interfaces with same semantics as paper



need to focus on details

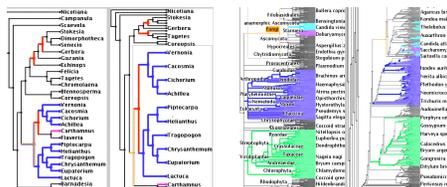


yet maintain context

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TreeJuxtaposer video

- stretch and squish navigation
- linked side by side comparison



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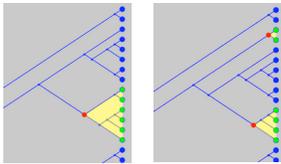
Encoding and interaction design decisions

- linking tree views through node correspondences
 - showing structural differences
- guaranteed visibility of small marks
 - scaling up to millions of nodes

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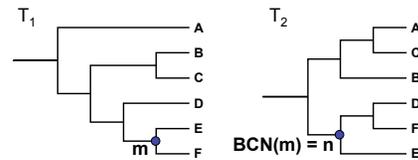
Showing differences via contiguous groups

- **clade**: ancestor + all descendants
- biological questions to support
 - is a clade in one tree also a clade in other?
 - is some group a clade?



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Best corresponding node between trees

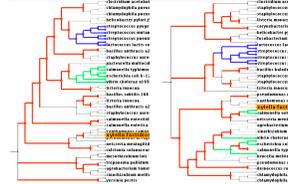


- used for
 - comparing clades between trees
 - linked highlighting on mouseover
 - structural difference highlighting
- algorithm scalability challenge
 - computable in $O(n \log^2 n)$, vs. naive $O(n^2)$

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Guaranteed visibility

- marks are always visible
 - structural differences, search results, user selections
- easy with small datasets
 - regions of interest shown with color highlights



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Guaranteed visibility challenges

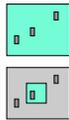
- hard with larger datasets
- reasons a mark could be invisible



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Guaranteed visibility challenges

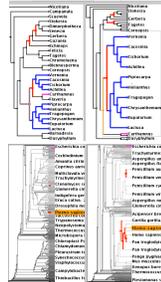
- hard with larger datasets
- reasons a mark could be invisible
 - mark outside the window
 - solution: constrained navigation



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Constrained navigation for visibility

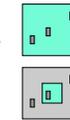
- stretch and squish navigation
 - stretch out part of surface, the rest squishes
 - borders nailed down
 - integrated focus and context
- items never fall outside camera
 - but squished regions can have many items per pixel



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Guaranteed visibility challenges

- hard with larger datasets
- reasons a mark could be invisible
 - mark outside the window
 - solution: constrained navigation
 - mark underneath other marks
 - solution: use 2D not 3D layout



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Guaranteed visibility challenges

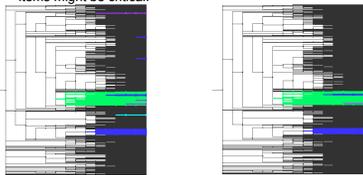
- hard with larger datasets
- reasons a mark could be invisible
 - mark outside the window
 - solution: constrained navigation
 - mark underneath other marks
 - solution: use 2D not 3D layout
 - mark smaller than a pixel
 - solution: smart culling



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Smart culling for small item visibility

- naive culling does not draw all marked items
 - graphics cards optimized for realism: small items far away and thus not important
 - rendering infrastructure for visualization semantics: small items might be critical!



guaranteed mark visibility

no guaranteed visibility

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Guaranteed visibility benefits

- with GV
 - no mark is visible means no need to explore area further
 - without GV
 - risk of false negative conclusions, or
 - user must do tedious exhaustive search to ensure nothing missed
 - algorithm scalability challenge
 - rendering complexity based on number of onscreen pixels
 - not total number of items in dataset
- Partitioned Rendering Infrastructure for Scalable Accordion Drawing (Extended Version). James Slack, Kristian Hildebrand, and Tamara Munzner. Information Visualization, 5(2), p. 137-151, 2006
- Composite Rectilinear Deformation for Stretch and Squish Navigation. James Slack and Tamara Munzner. Proc. Visualization 2006, published as Transactions on Visualization and Computer Graphics 12(9), September 2006.

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TJ summary

- first interactive tree comparison system
 - automatic structural difference computation
 - guaranteed visibility of small marks
- scalable to large datasets
 - 250K to 500K total nodes: original
 - up to 4M nodes: later, with PRISAD
 - subquadratic preprocessing
 - sublinear realtime rendering
 - depends on number of pixels, not number of nodes

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Outline

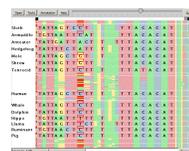
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Many other bio/vis research areas

- multiple sequence alignment
 - SequenceJuxtaposer

open-source software download
<http://olduvai.sourceforge.net/sj/>



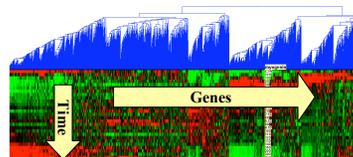
SequenceJuxtaposer: Fluid Navigation For Large-Scale Sequence Comparison In Context. James Slack, Kristian Hildebrand, Tamara Munzner, and Katherine St. John. German Conference on Bioinformatics 2004, pp 37-42

<http://www.cs.ubc.ca/labs/imager/tr/2004/sj/>

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Many other bio/vis research areas

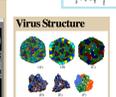
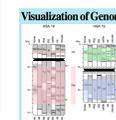
- microarray data
 - Hierarchical Clustering Explorer
<http://www.cs.umd.edu/hcil/hce/>
 Seo and Shneiderman, U Maryland



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Many more, bio/health + others...

- NIH/NSF Visualization Research Challenges Report Johnson, Moorhead, Munzner, Pfister, Rheingans, and Yoo (eds.), IEEE Press
http://vgtc.org/wpmu/techcom/?page_id=11



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Crosscutting themes

- workflow speedups
 - inspecting microarray data with graph
 - minutes vs. hours/days
 - comparing clades between trees
 - seconds vs. hours/days
- contributions from biologist collaborators
 - driving problems and data
 - tool use during iterative refinement

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Vast opportunities

- young field, still much to be done
- think about your current workflow
 - what could you speed up by swapping in perception for cognition?
 - exploit the familiar, yet consider breadth of design alternatives
- finding some friendly neighborhood vis collaborators
 - IEEE VisWeek 2009 (Vis, InfoVis, VAST)
Oct 11-16, Atlantic City
<http://vis.computer.org/VisWeek2009>
 - EuroVis 2009: Jun 10-12, Berlin
<http://www.zib.de/eurovis09>

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More information

- this talk
<http://www.cs.ubc.ca/~tmm/talks.html#harvard09>
- papers, videos
<http://www.cs.ubc.ca/~tmm>
- software
<http://olduvai.sourceforge.net/tj>
<http://www.pathogenomics.ca/cerebral>
<http://www.innatedb.ca>

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