## Phylogenetic Biology Week 7

Biology 1425
Professor: Casey Dunn, dunnlab.org
Brown University
2013

### Front matter...

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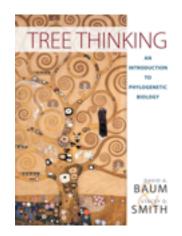


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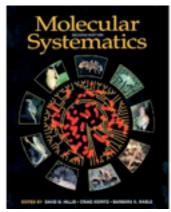
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### Sources

### Some non-original content is drawn from:



Baum, D and S. Smith (2012) Tree Thinking: and Introduction to Phylogenetic Biology. Roberts and Company Publishers. ISBN 9781936221165



INFERRING PHYLOGENIES

Joseph Felsenstein

Swofford, D. L., Olsen, G. J., Waddell, P. J., & Hillis, D. M. (1996). Phylogenetic inference. In: Molecular Systematics, Second Edition. eds: D. M. Hillis, C Moritz, & B. K. Mable. Sinauer Associates. ISBN 9780878932825

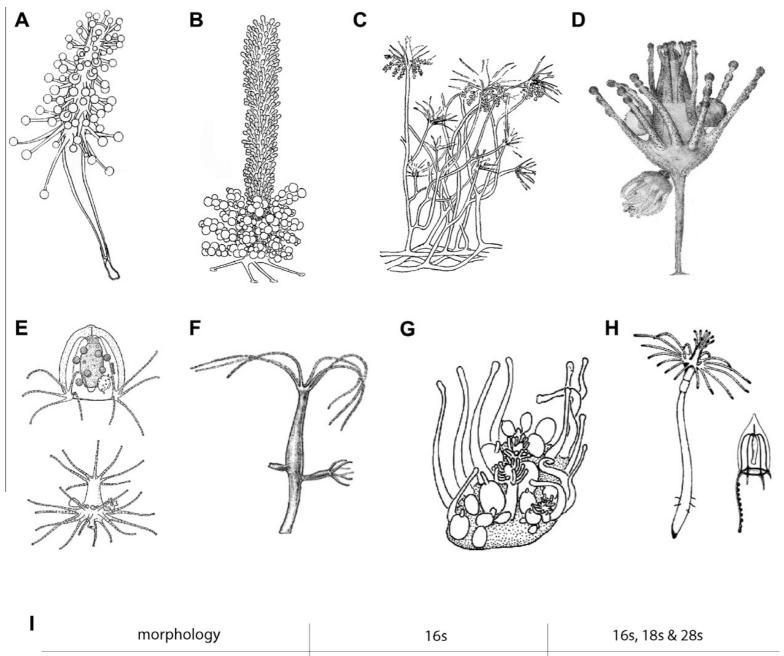
Felsenstein, J. (2003) Inferring Phylogenies. Sinauer Associates. ISBN 978-0878931774

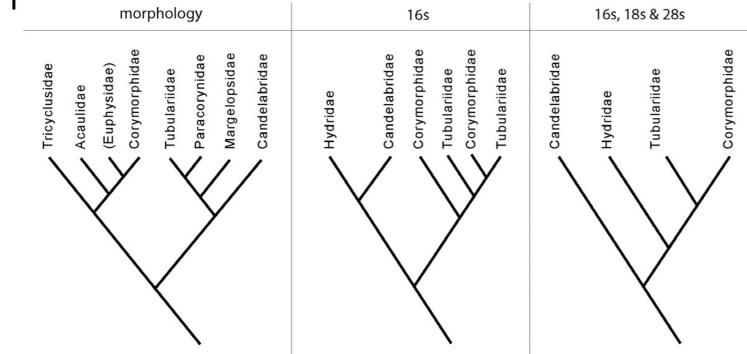
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## Phylogenetics in the wild-Project design and the challenges that arise when executing a study

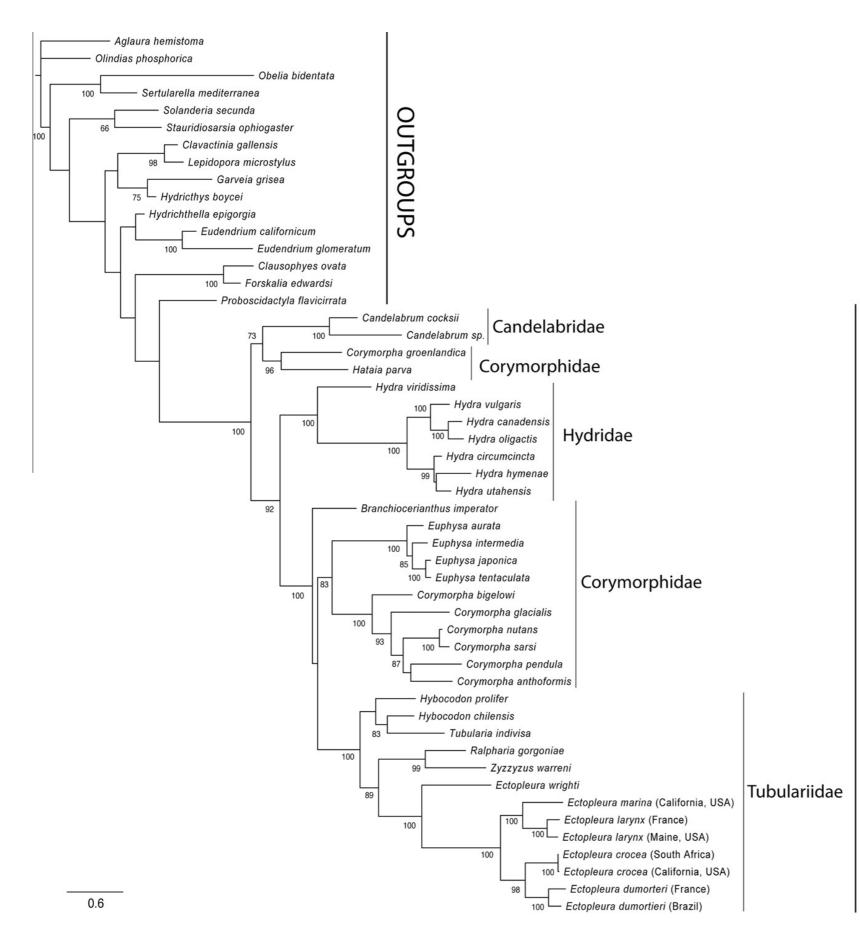
### An example study

Nawrocki, A. M., Collins, A. G., Hirano, Y. M., Schuchert, P. & Cartwright, P. **Phylogenetic placement of Hydra and relationships within Aplanulata (Cnidaria: Hydrozoa)**. Molecular Phylogenetics and Evolution 67, 60–71 (2013). http://dx.doi.org/10.1016/j.ympev.2012.12.016





Nawrocki et al 2013, Figure 1



## Bootstrap, all markers

Nawrocki et al 2013, Figure 3

**Aplanulata** 

## How important is taxon sampling?

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The better your taxon sampling, the more questions you can address.

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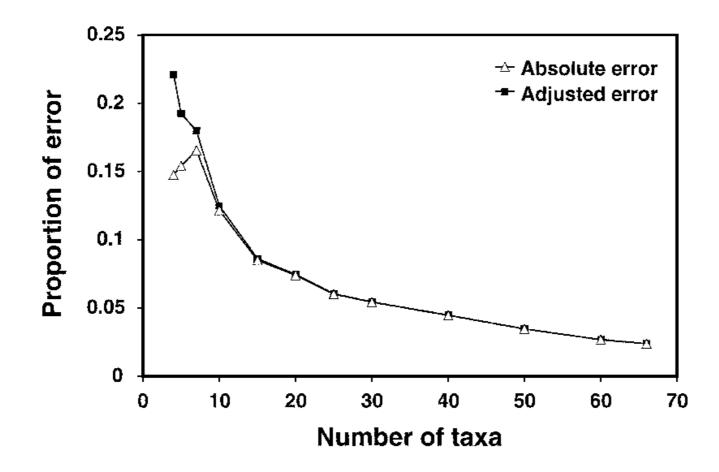
Better taxon sampling can improve accuracy

Syst. Biol. 51(4):588–598, 2002 DOI: 10.1080/10635150290102339

#### Increased Taxon Sampling Greatly Reduces Phylogenetic Error

#### DERRICK J. ZWICKL AND DAVID M. HILLIS

Section of Integrative Biology and Center for Computational Biology and Bioinformatics, University of Texas, Austin, Texas 78712, USA; E-mail: zwickl@mail.utexas.edu and dhillis@mail.utexas.edu



Zwickl and Hillis 2002, http://dx.doi.org/10.1080/10635150290102339

# How important are outgroups?

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Poor outgroup sampling is one of the biggest rookie mistakes in phylogenetic analyses

# How important is character sampling?

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Better character sampling can improve accuracy

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Better character sampling can improve accuracy

Better character sampling can provide more rigorous evaluation of hypotheses

# How does missing data impact phylogenetic analyses?

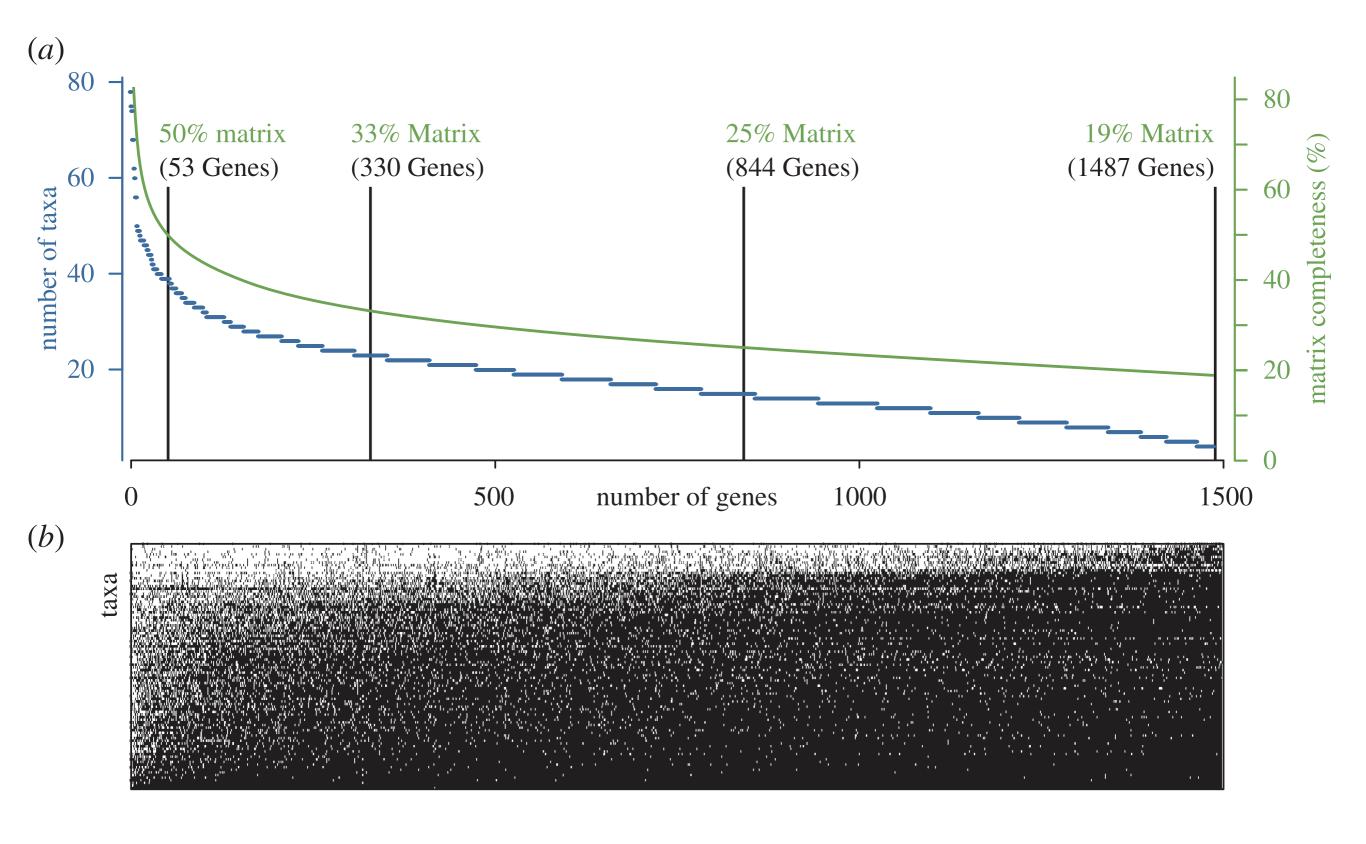
## Missing data leads to loss of signal.

## Missing data leads to loss of signal.

## Can it also lead to inconsistency?

# Missing data in phylogenetic analyses are rarely distributed randomly.

Some genes are better sampled than others, some taxa are better sampled than others.



Hejnol et al. 2009, http://dx.doi.org/10.1098/rspb.2009.0896

Should you add poorly sampled genes and taxa to an analysis, or have a smaller but more complete analysis?

## Amount of missing data: less more

Eliminate poorly sampled taxa

Eliminate poorly sampled genes

Additional expense

Broader taxon sampling

Broader gene sampling

Reduced expense

Could be impacted by inconsistency



#### Available online at www.sciencedirect.com



Biomedical Informatics

Journal of Biomedical Informatics 39 (2006) 34-42

www.elsevier.com/locate/yjbin

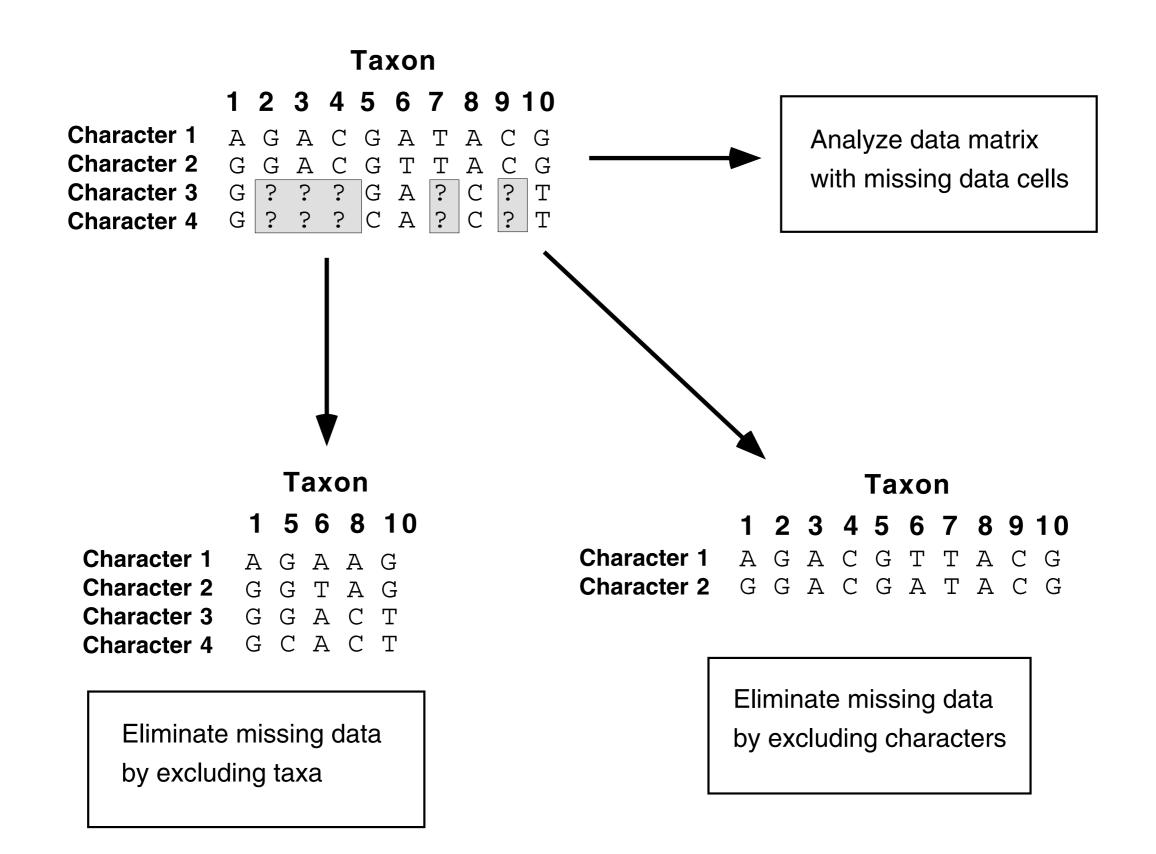
Methodological Review

#### Missing data and the design of phylogenetic analyses

John J. Wiens \*

Department of Ecology and Evolution, Stony Brook University, Stony Brook, NY 11794-5245, USA

Received 5 March 2005 Available online 6 May 2005



"Recent simulations suggest that highly incomplete taxa can be accurately placed in phylogenies, as long as many characters have been sampled overall. Furthermore, adding incomplete taxa can dramatically improve results in some cases by subdividing misleading long branches. Adding characters with missing data can also improve accuracy, although there is a risk of long-branch attraction in some cases."

Syst. Biol. 58(1):130–145, 2009 Copyright © Society of Systematic Biologists DOI:10.1093/sysbio/syp017 Advance Access publication on May 21, 2009

### The Effect of Ambiguous Data on Phylogenetic Estimates Obtained by Maximum Likelihood and Bayesian Inference

Alan R. Lemmon<sup>1,2,3,\*</sup>, Jeremy M. Brown<sup>1</sup>, Kathrin Stanger-Hall<sup>4</sup>, and Emily Moriarty Lemmon<sup>1,3</sup>

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<sup>3</sup>Present address: Department of Biological Science, Florida State University, Tallahassee, FL 32306, USA;

<sup>4</sup>Plant Biology Department, University of Georgia, 403 Biosciences Building, Athens, GA 30602, USA;

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FL 32306-4120, USA; E-mail: alemmon@evotutor.org.

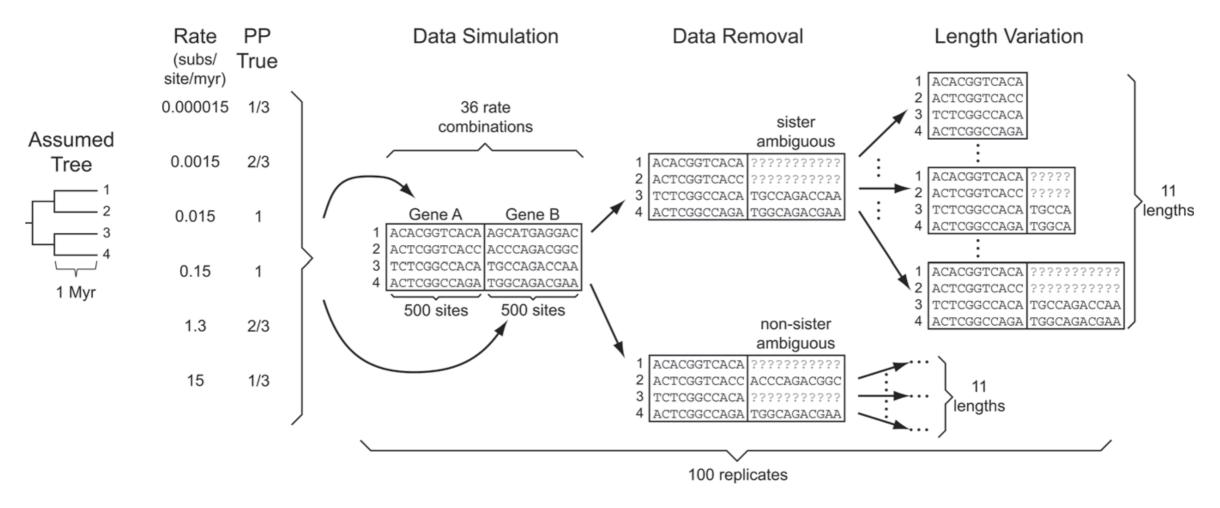


FIGURE 1. Simulation design. Among-site rate variation was simulated using 6 rates of evolution (chosen to produce the desired PP for the true tree with 500 sites) combined across 2 genes to form 36 rate combinations. Gene A contained unambiguous sites, whereas Gene B contained ambiguous sites. Ambiguous characters were present for either sister or nonsister taxa. Although Gene A always contained 500 sites, the length of Gene B varied from 0 to 500 sites. Note that Gene B contained no topological information, regardless of the rate of evolution. PP = posterior probabilities.

### Lemmon et al. 2009, http://dx.doi.org/10.1093/sysbio/syp017

"We find that in both ML and Bayesian frameworks, among-site rate variation can interact with ambiguous data to produce misleading estimates of topology and branch lengths."

### **BMC Evolutionary Biology**



Research article



### Using ESTs for phylogenomics: Can one accurately infer a phylogenetic tree from a gappy alignment?

Stefanie Hartmann<sup>1,2</sup> and Todd J Vision\*<sup>1</sup>

Address: <sup>1</sup>Department of Biology, University of North Carolina, Chapel Hill, NC 27599, USA and <sup>2</sup>Institute for Biochemistry and Biology, Karl-Liebknecht-Strasse 24-25, University of Potsdam, 14476 Potsdam, Germany

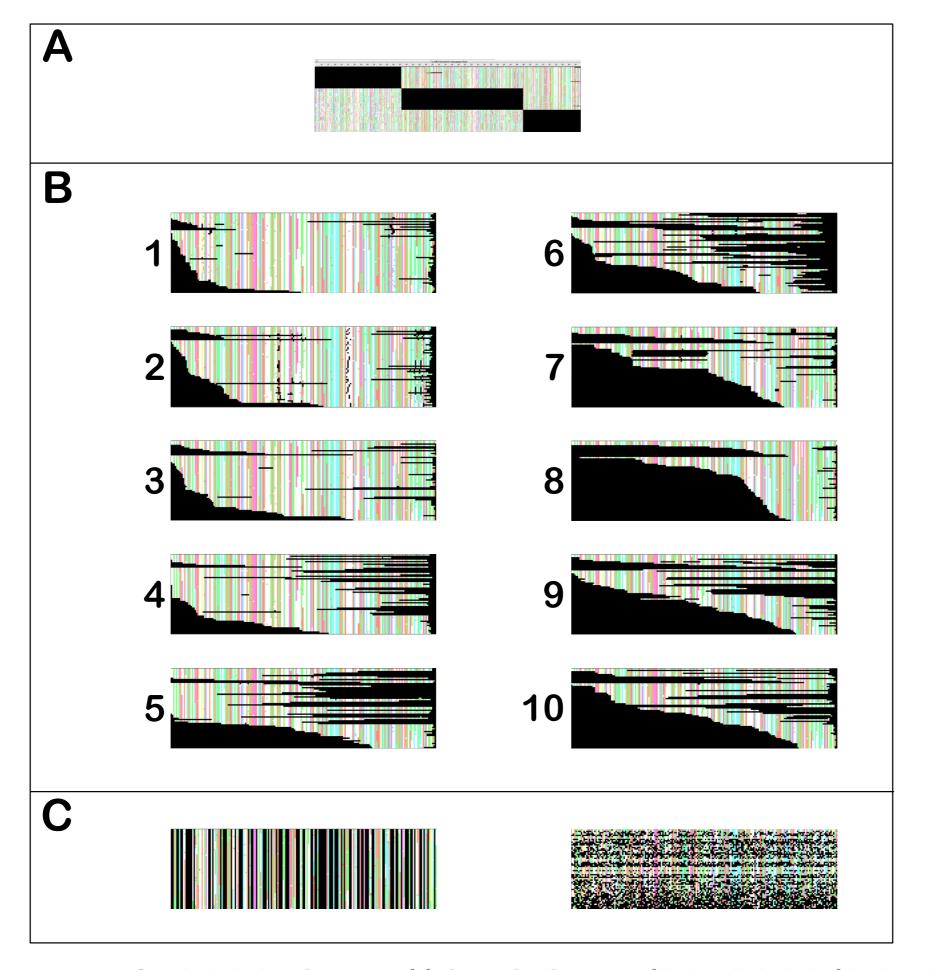
Email: Stefanie Hartmann - stefanie.hartmann@uni-potsdam.de; Todd J Vision\* - tjv@bio.unc.edu

\* Corresponding author

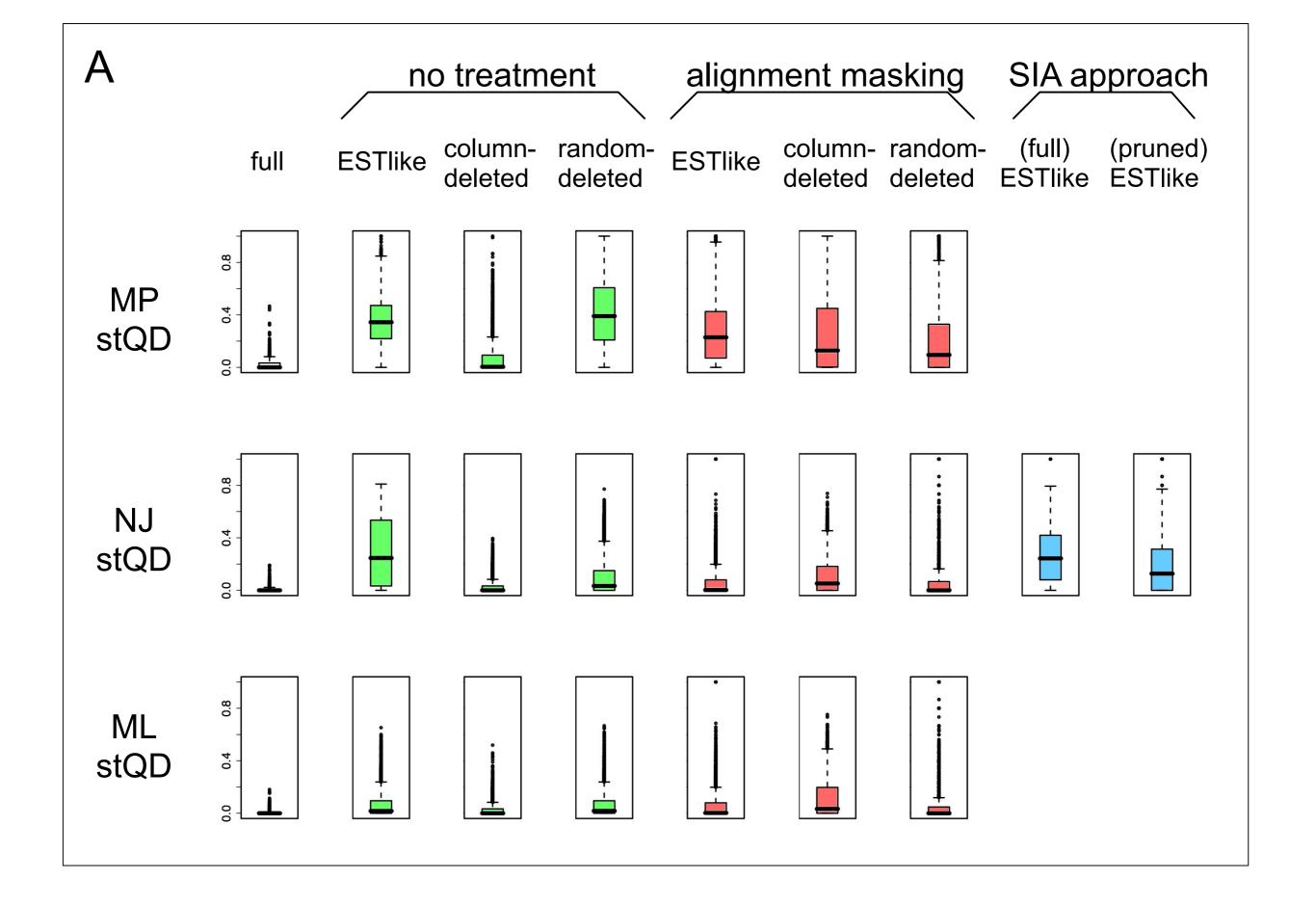
Published: 26 March 2008

BMC Evolutionary Biology 2008, 8:95 doi:10.1186/1471-2148-8-95

Received: 27 October 2007 Accepted: 26 March 2008



Hartmann et al. 2008, http://dx.doi.org/10.1186/1471-2148-8-95



Hartmann et al. 2008, http://dx.doi.org/10.1186/1471-2148-8-95

## How to collect your molecular data?

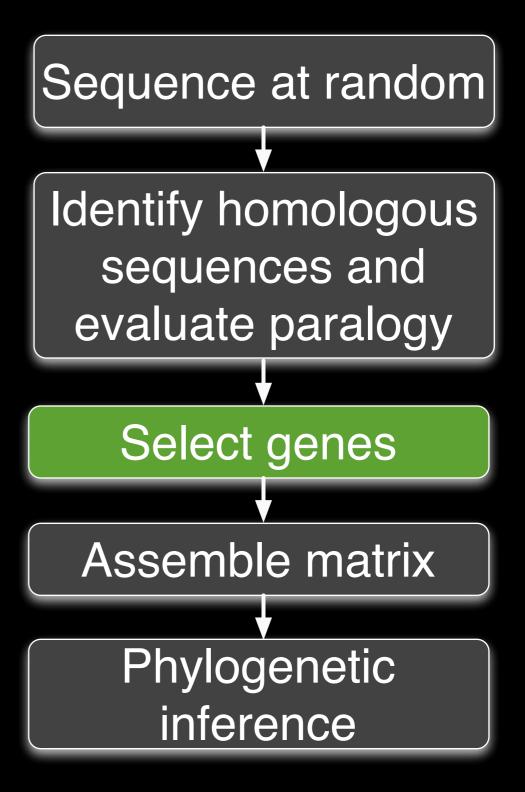
Directed sequencing - pick genes first, then sequence just them

Phylogenomic analyses - use high throughput transcriptome and genome data to broadly sample many genes

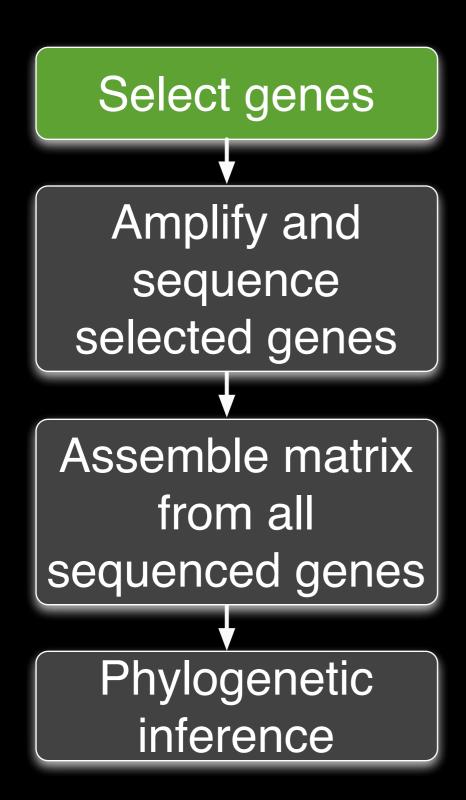
## Gene selection as part of project design

### Select genes Amplify and sequence selected genes Assemble matrix from all sequenced genes Phylogenetic inference

### Gene selection as part of analysis



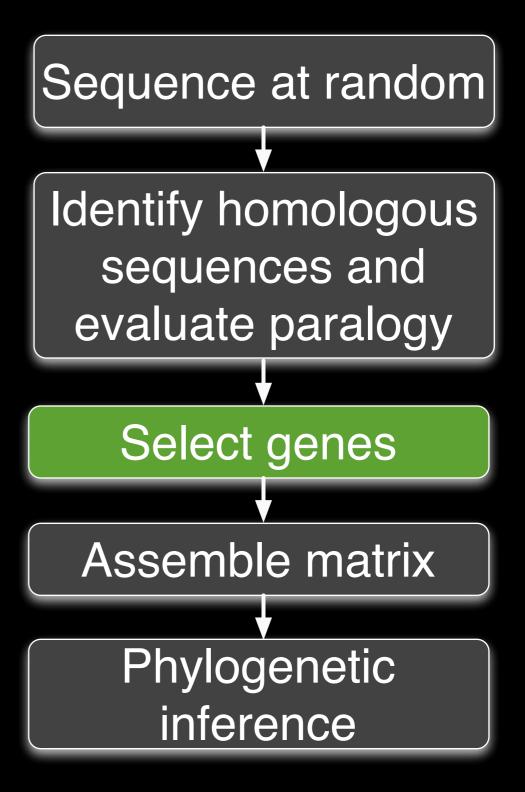
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### Select genes Amplify and sequence selected genes Assemble matrix from all sequenced genes Phylogenetic inference

### Gene selection as part of analysis



# Should you include morphological data in your analysis?

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If you are analyzing fossils, it is essential.

## Common tools for inferring trees:

Parsimony

PAUP\*

Poy

**TNT** 

Likelihood

PAUP\*

RAxML

**GARLI** 

MrBayes

Bayesian

MrBayes

**PhyloBayes**