



IQ-TREE
<http://www.iqtree.org>

Methods and Practice

Minh Bui
Australian National University

Workshop on Molecular Evolution
Woods Hole, June 2022

IQ-TREE DEVELOPMENT TEAM

Australia



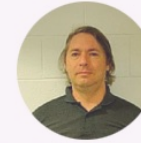
James Barbetti

Contribution: Software engineering for COVID-19 data



Thomas Wong

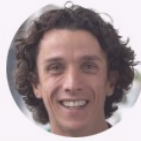
Contribution: ModelFinder 2



Michael Woodhams

Google Scholar

Contribution: Lie Markov models.



Robert Lanfear

Google Scholar

Contribution: Inspiring ideas and advice.



Bui Quang Minh

Google Scholar

Contribution: Team leader, software core, ultrafast bootstrap, model selection.



Nhan Trong Ly

Contribution: sequence simulations.

Austria



Olga Chernomor

Google Scholar

Contribution: Partition models and phylogenomic search.



Arndt von Haeseler

Google Scholar

Contribution: Inspiring ideas and advice.



Dominik Schrempf

Google Scholar

Contribution: Polymorphism-aware models (PoMo).



Heiko A. Schmidt

Google Scholar

Contribution: Integration of **TREE-PUZZLE** features.



Diep Thi Hoang

Contribution: Improving ultrafast bootstrap.

Vietnam

Thanks to plenty of users for feedback and bug reports!

Why IQ-TREE?

Next generation sequencing data represent both a blessing and a curse:

- Blessing: (Phylo)genomic data help to elucidate many phylogenetic questions.
- Curse: Many model assumptions become increasingly distant from the truth due to growing data complexity.

“All models are wrong, but some are useful” (Box, 1976)

With IQ-TREE we aim to:

- Analyze ultra-large data sets.
- Provide many (if not most) “useful” models of sequence evolution.

But still, there are RAxML, PhyML out there, why do I need IQ-TREE?

- We better have at least 2 software independently developed for similar purpose. Only then, the pros and cons (sometimes **bugs**) can be identified. This creates a *friendly* competition, which helps to advance the field!
- Same as having MrBayes, RevBayes, BEAST for Bayesian inference.

Typical phylogenetic analysis under maximum likelihood

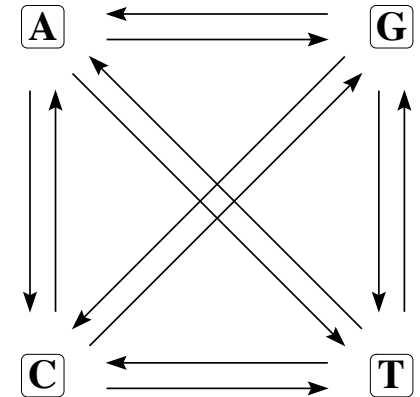
Multiple sequence alignment

```
ACGGGAT--C--C----CATTAC
ACGGGAT--C--C----CACTAC
CCGGGATAGCTTC----CATTAC
ACCCCCTATC--CACTGGATTAC
ACGACATATC--CACTGGATTCC
```

Model selection

ModelFinder (2017)

Substitution model

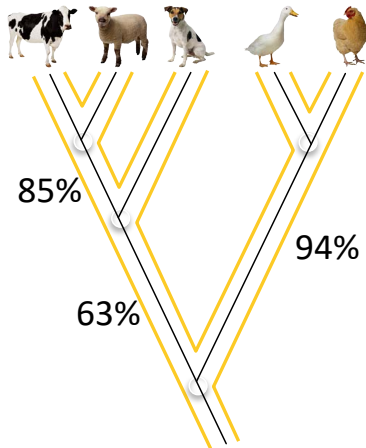


My work focused on improving all three steps for large datasets!

IQ-TREE (2015, 2020)

Tree reconstruction

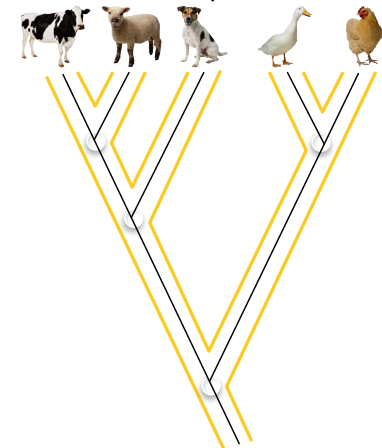
`iqtree2 -s ALN_FILE -B 1000`



Tree with branch supports

Ultrafast bootstrap (2013, 2018)

Assessment of branch supports



Phylogenetic tree

Step 1: Model selection

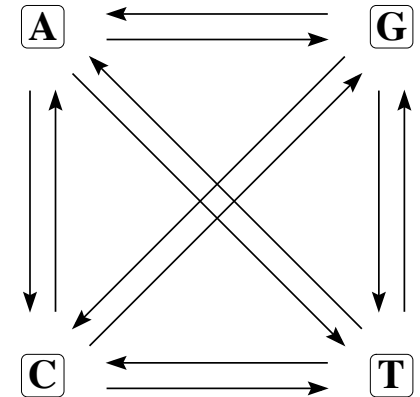
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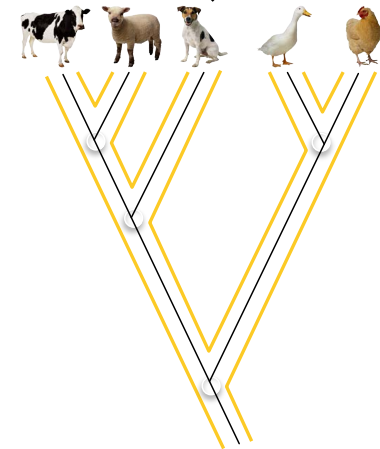
Model selection

ModelFinder (2017)

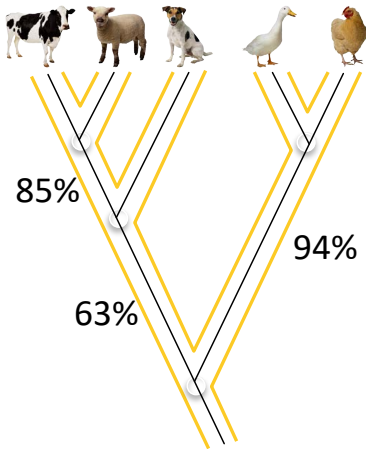
Substitution model



Tree reconstruction



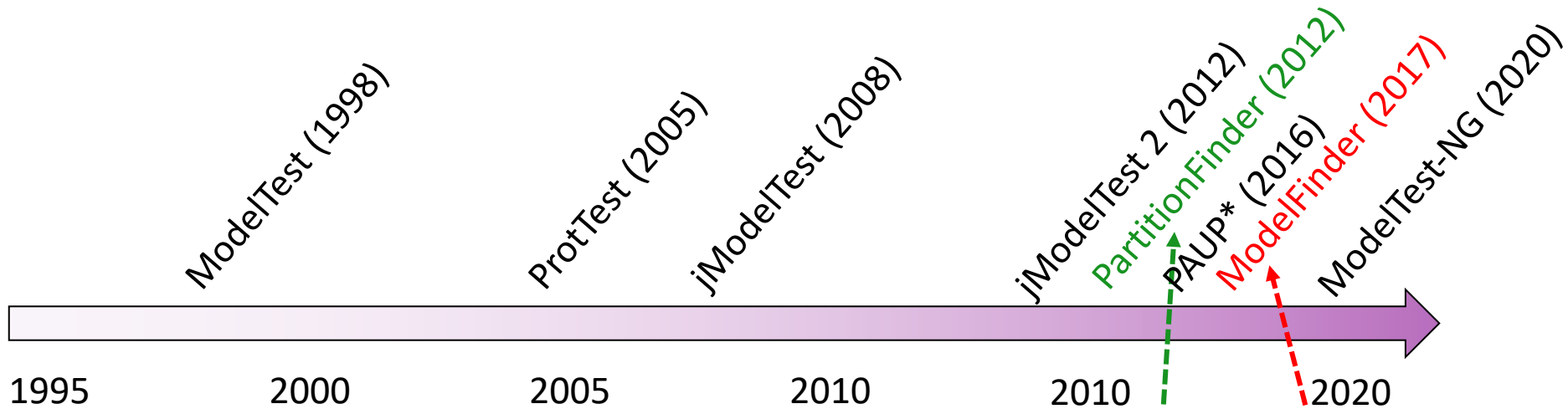
Assessment of branch supports



Tree with branch supports

Phylogenetic tree

Model selection approaches



- (j)Modeltest / ProtTest: slow and limited on models.
- **PartitionFinder**: better models for genomic data but still slow.
- **ModelFinder**: >10x faster and more realistic models.
- **Current work: ModelFinder 2 = ModelFinder + PartitionFinder**



Robert Lanfear
(ANU)

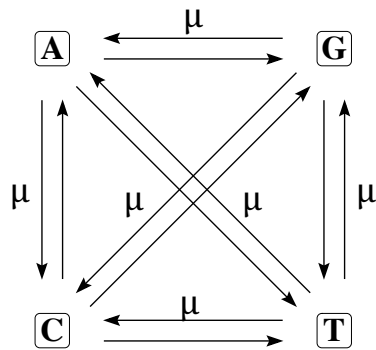


Lars Jermin
(ANU & CSIRO)



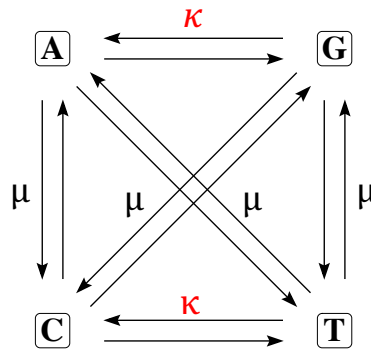
Thomas Wong
(ANU)

Models of sequence evolution



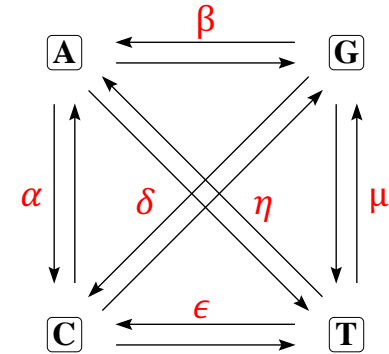
JC

(Jukes & Cantor 1969)



HKY

(Hasegawa, Kishino,
Yano 1985)



GTR

(General Time
Reversible, 1986)

Rate heterogeneity: alignment sites evolved at different rates. Some slow, some fast.

Rate model	Explanation
+I	Some sites are <i>invariable</i> (zero rate), e.g. due to selective force.
+G	Site rates follow a <i>Gamma</i> distribution.
+I+G	Some sites are invariable, the rest follow a Gamma distribution.
+R	Sites fall into several categories from slow to fast rates. No assumption of rate distribution (free-rate model).

A model = substitution model + rate heterogeneity, e.g. “GTR+G”

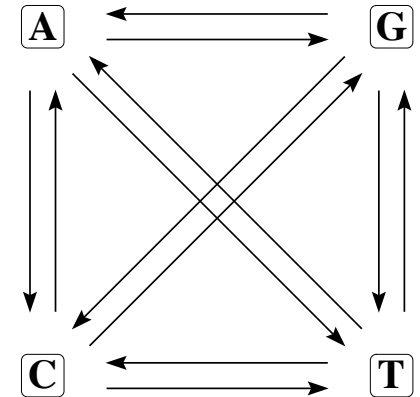
Step 2: Tree reconstruction

Multiple sequence alignment

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```

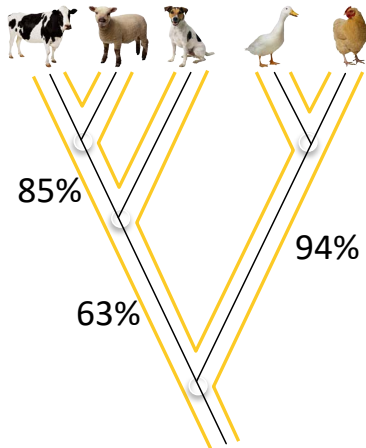
Model selection

Substitution model



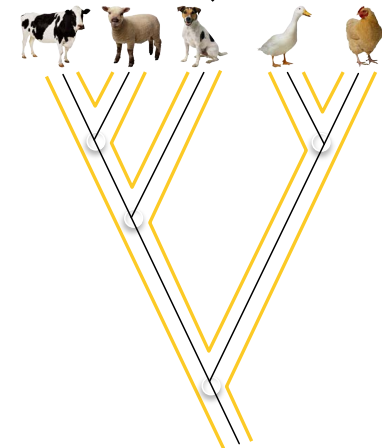
IQ-TREE (2015, 2020)

Tree reconstruction



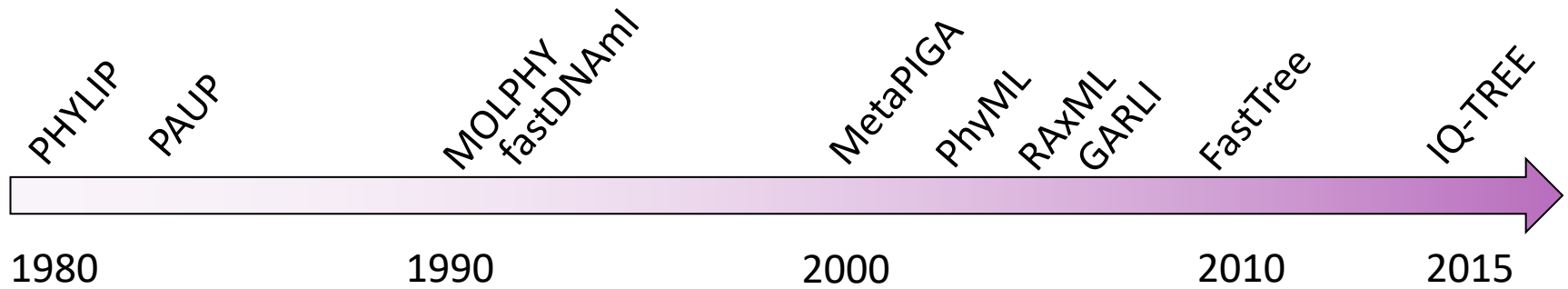
Tree with branch supports

Assessment of branch supports

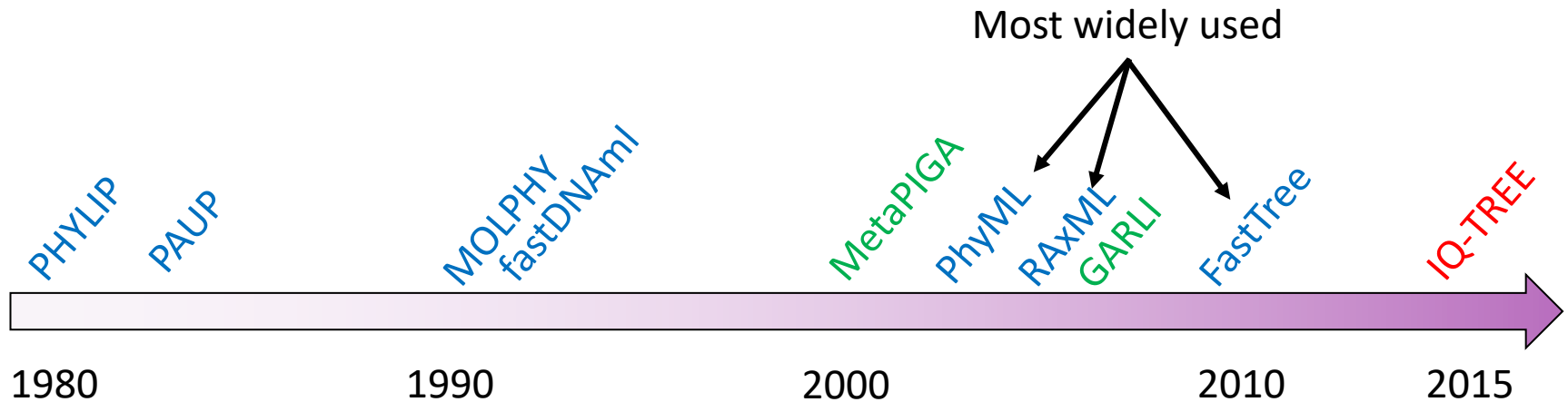


Phylogenetic tree

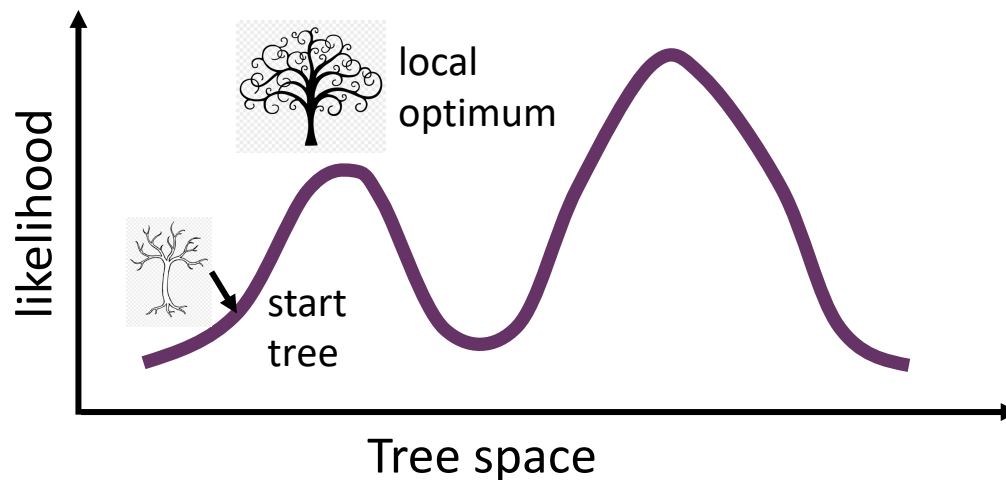
Search heuristics for finding maximum likelihood trees



Search heuristics for finding maximum likelihood trees



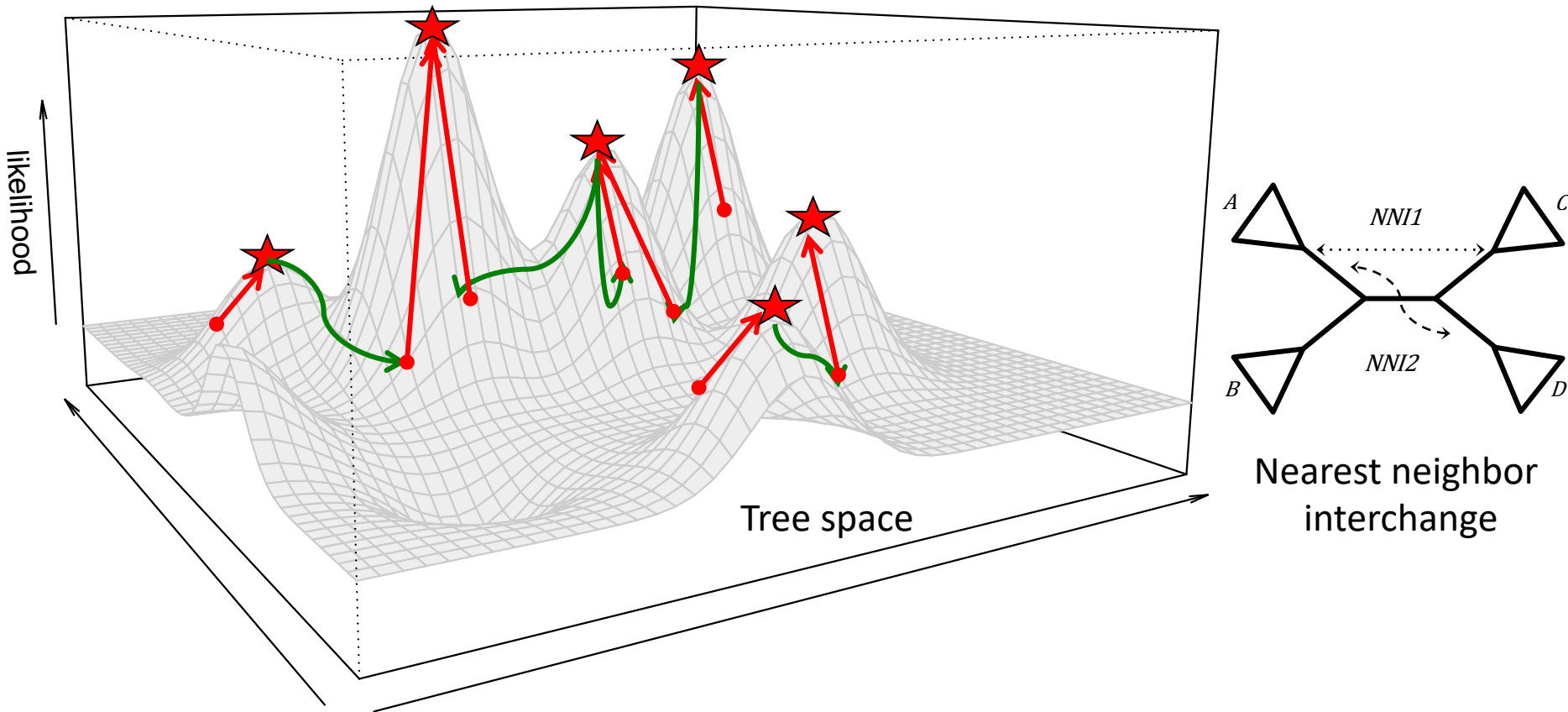
1. Hill-climbing / greedy algorithms: Fast but local optimum
2. Genetic algorithm: Slow but escaping local optima
3. IQ-TREE: Fast and escaping local optima



Tree search algorithms in RAxML and IQ-TREE

Feature	RAxML	IQ-TREE
Starting tree	Parsimony: Stepwise addition + subtree pruning and regrafting (SPR)	99 parsimony trees (like RAxML) and 1 Neighbor-joining tree
Tree search heuristics	Hill-climbing SPR	Stochastic: Hill-climbing Nearest Neighbor Interchange (NNI) and downhill NNI

IQ-TREE: A new stochastic algorithm

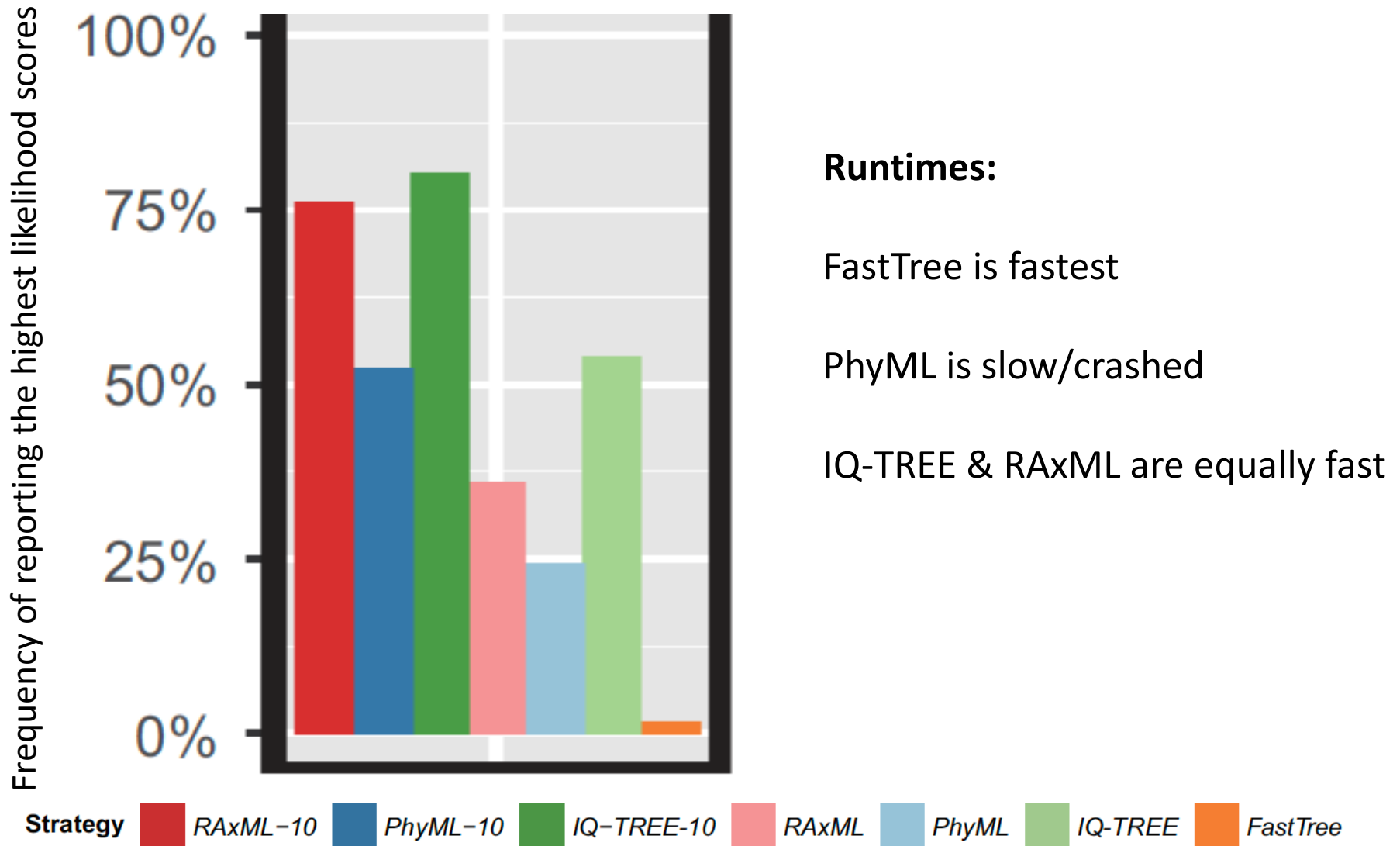


Metaheuristics:
*Random restart, Iterated local search,
Evolution strategy*

Lam-Tung Nguyen Heiko Schmidt Arndt von Haeseler



An independent benchmark by Zhou et al. (2018)



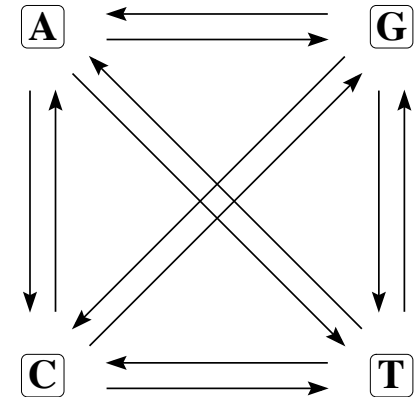
Step 2: Summary

Multiple sequence alignment

```
ACGGGAT--C--C----CATTAC
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```

Model selection

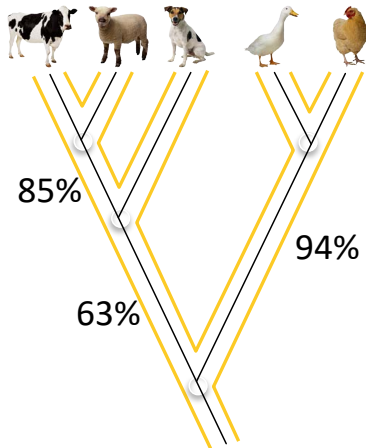
Substitution model



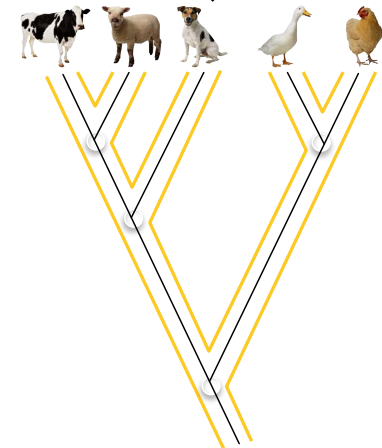
- IQ-TREE algorithm efficiently explores tree space

IQ-TREE (2015, 2020)

Tree reconstruction



Assessment of branch supports



Tree with branch supports

Phylogenetic tree

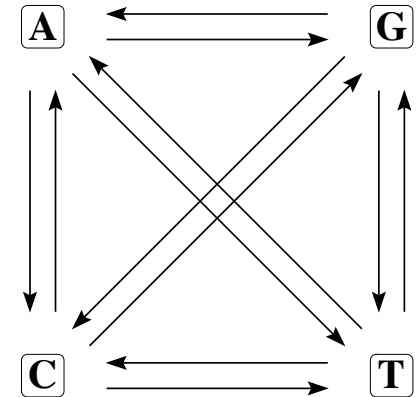
Step 3: Ultrafast bootstrap

Multiple sequence alignment

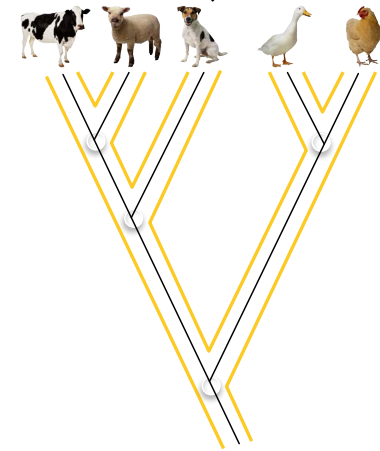
```
ACGGGAT--C--C----CATTAC
ACGGGAT--C--C----CACTAC
CCGGGATAGCTTC----CATTAC
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ACGACATATC--CACTGGATTCC
```

Model selection

Substitution model



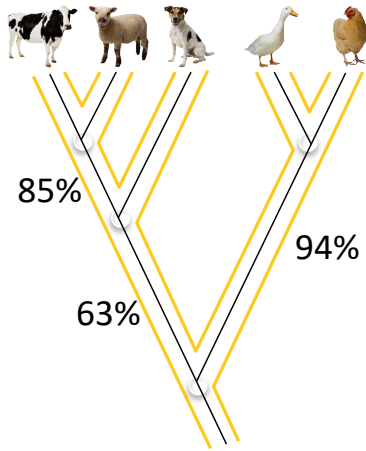
Tree reconstruction



Phylogenetic tree

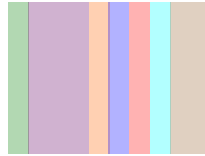
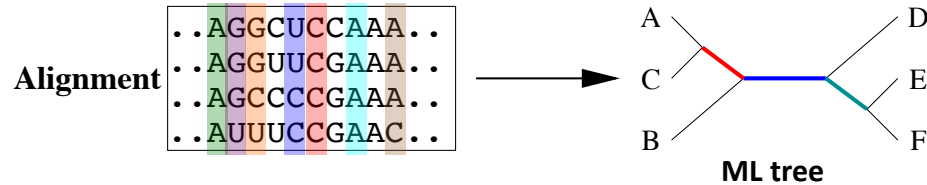
Ultrafast bootstrap (2013, 2018)

Assessment of branch supports



Tree with branch supports

Bootstrap: How reliable are branches of the tree?



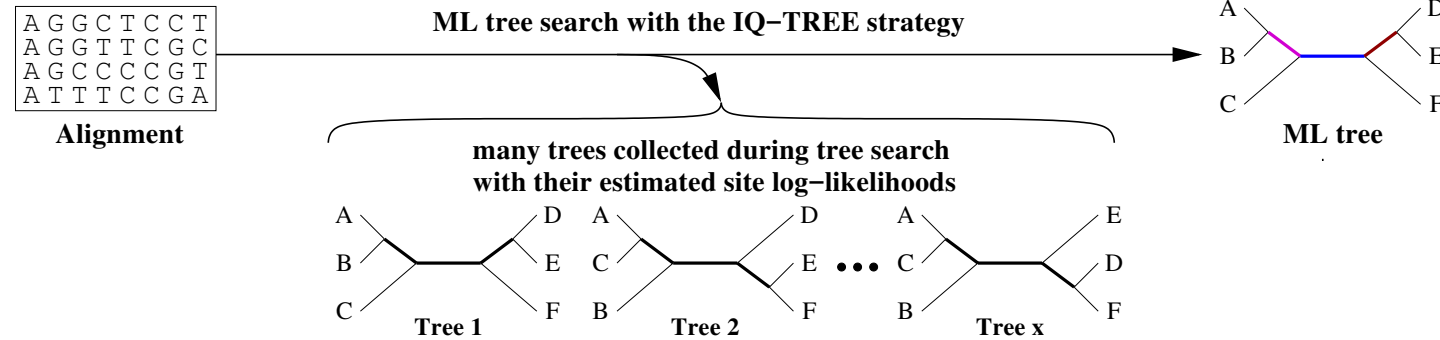
Consensus tree

Bootstrap analysis is
extremely time-consuming!

UFBoot: Ultrafast bootstrap approximation



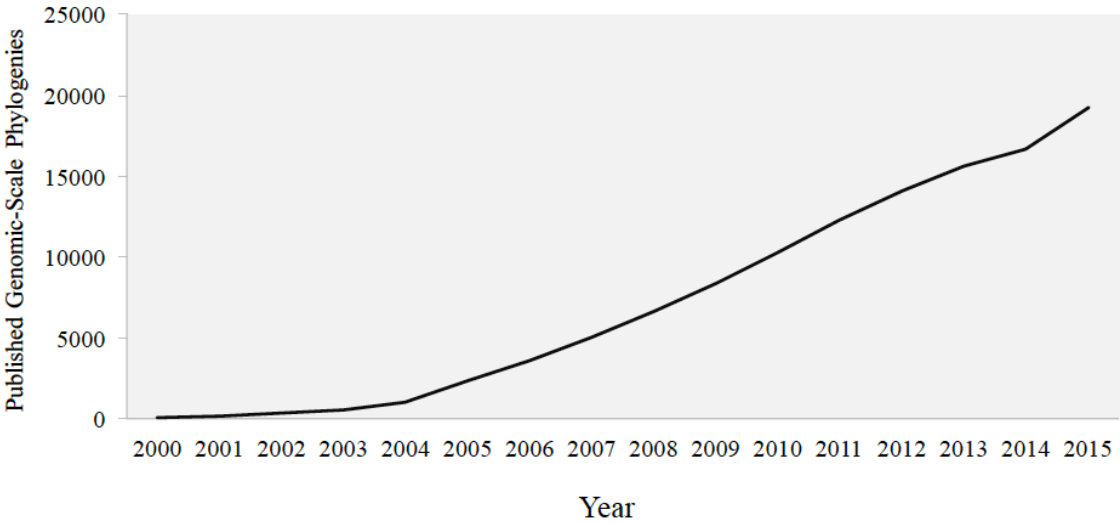
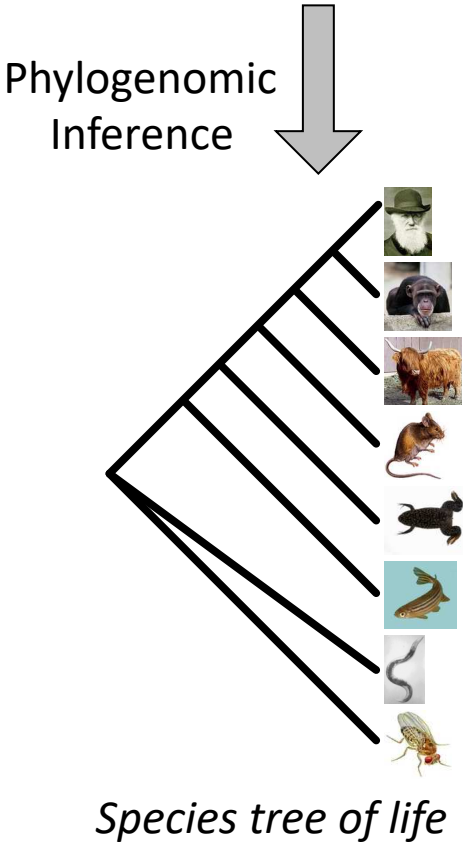
M.A.T. Nguyen, A. von Haeseler



Use UFBoot $\geq 95\%$
instead of 70% !

Genome-scale data: Concatenation methods

Supermatrix			
Gene 1	Gene 2	Gene 1,000
CACCTGTCGT	-----	-----	TCTGGTGCAG
CAGCTGTCGT	GCTCTTTCTG	TTGAGCCTGG	TCTGGTGCAG
CAGCTGCCGT	GTTTTCTCTG	TTGAGCCTGG	TCTGGTACAG
CAGCTGCCGC	GTTCTCTCCG	-----	TCTGGTGCAA
CTCCTGCCGG	GTGCTCTCAG	-----	-----
CTCCTGCCGG	-----	CTGAGCCGGG	TCTGGTGCAG
CTCTTGCCGG	-----	CTGAGCCTTG	-----



30 days of computation and 280 GB RAM for an insect data set!

Partition model

Supermatrix

Gene 1	Gene 2	Gene 1,000
CACCTGTCGT	-----	-----	TCTGGTGCAG
CAGCTGTCGT	GCTCTTTCTG	TTGAGCCTGG	TCTGGTGCAG
CAGCTGCCGT	GTTTTCTCTG	TTGAGCCTGG	TCTGGTACAG
CAGCTGCCGC	GTTCTCTCCG	-----	TCTGGTGCAA
CTCCTGCCGG	GTGCTCTCAG	-----	-----
CTCCTGCCGG	-----	CTGAGCCGGG	TCTGGTGCAG
CTCTTGCCGG	-----	CTGAGCCTTG	-----

Substitution
models:

JC

HKY+G

.....

GTR+G

**Model of
branch lengths**

Gene trees

Universally
shared



Proportionally
linked

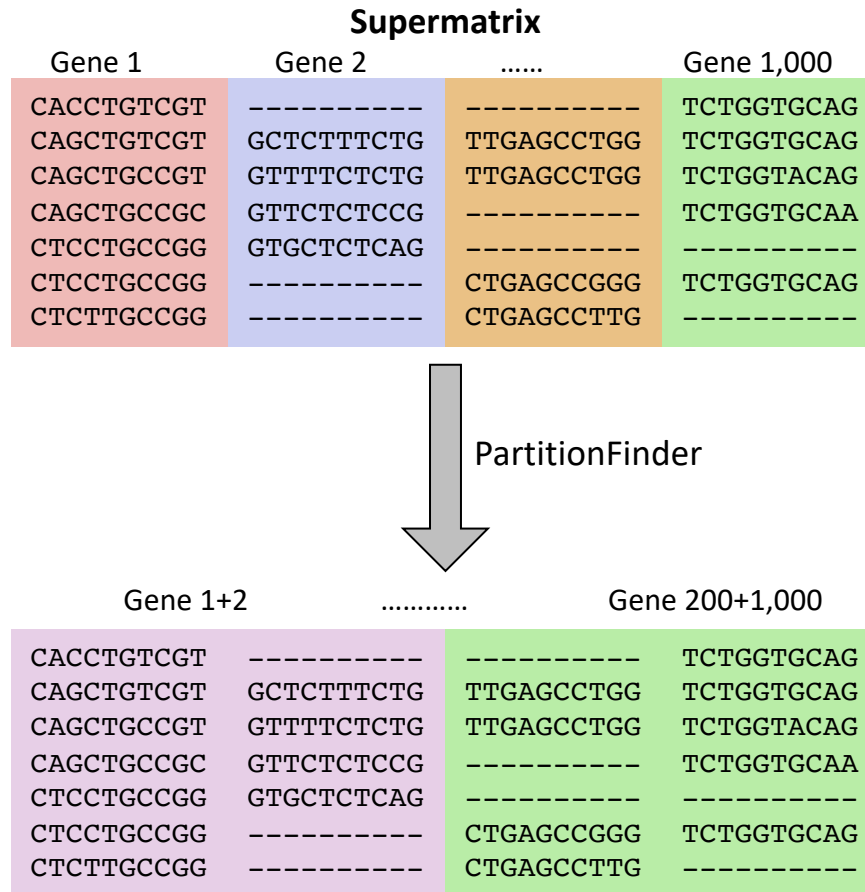


Unlinked



Recommended for typical analysis
([Duchene et al. 2020](#))

How to reduce potential model overfitting?



PartitionFinder algorithm (Lanfear et al. 2012):

1. Evaluate to merge all pairs of genes.
2. Choose the pair with the best score.
3. If score improves, merge two genes and repeat steps 1-3.
4. Otherwise, stop.

Relaxed clustering algorithm (Lanfear et al. 2014):

In step 1: only examine the top k% of most “promising” pairs.

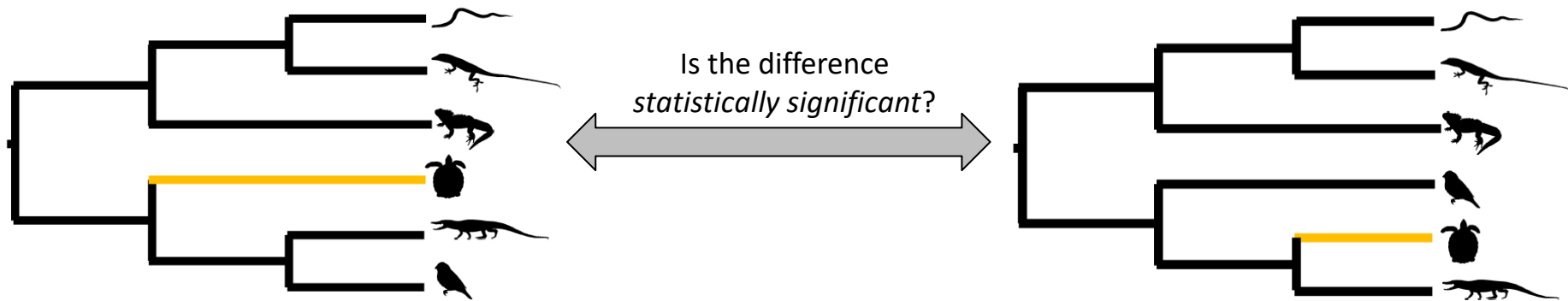
Substitution
models:

HKY

.....

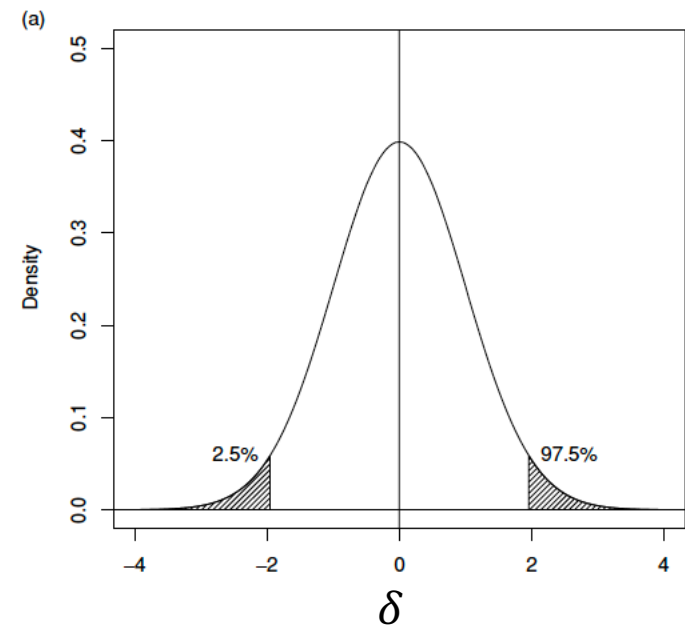
GTR+G

Tree topology tests



Testing two trees (Kishino & Hasegawa, 1989):

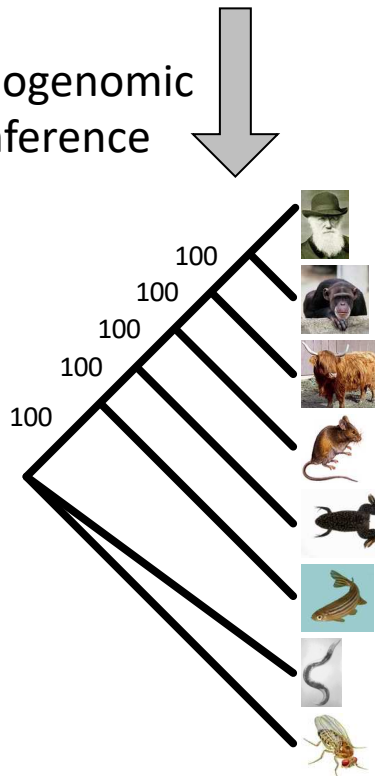
1. Statistic: $\delta = \log(\text{likelihood}(T_1)) - \log(\text{likelihood}(T_0))$.
2. Generate distribution of δ from many "random" data (e.g. by 1000 bootstrap resampling).
3. Compare the statistic between original and random data to obtain *p-value*.
4. If **p-value < 0.05**: YES! two trees are significantly different.
5. If p-value ≥ 0.05 : NO! they are not.



Concatenation methods: Limitation

Supermatrix			
Gene 1	Gene 2	Gene 1,000
CACCTGTCGT	-----	-----	TCTGGTGCAG
CAGCTGTCGT	GCTCTTTCTG	TTGAGCCTGG	TCTGGTGCAG
CAGCTGCCGT	GTTTTCTCTG	TTGAGCCTGG	TCTGGTACAG
CAGCTGCCGC	GTTCTCTCCG	-----	TCTGGTGCAA
CTCCTGCCGG	GTGCTCTCAG	-----	-----
CTCCTGCCGG	-----	CTGAGCCGGG	TCTGGTGCAG
CTCTTGCCGG	-----	CTGAGCCTTG	-----

Phylogenomic
Inference



Species tree of life

Bootstrap supports and Bayesian posteriors
tend to 100% as #genes increases!

Concatenation assumes a single tree
across all loci

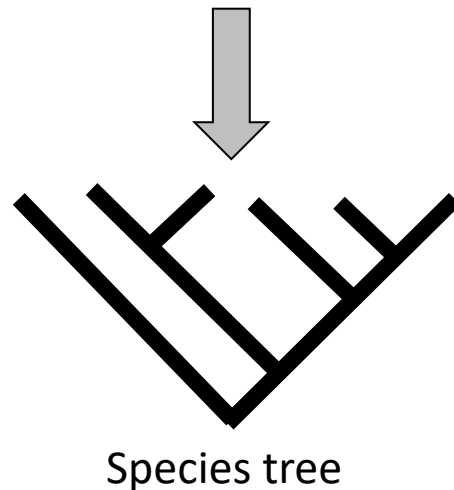
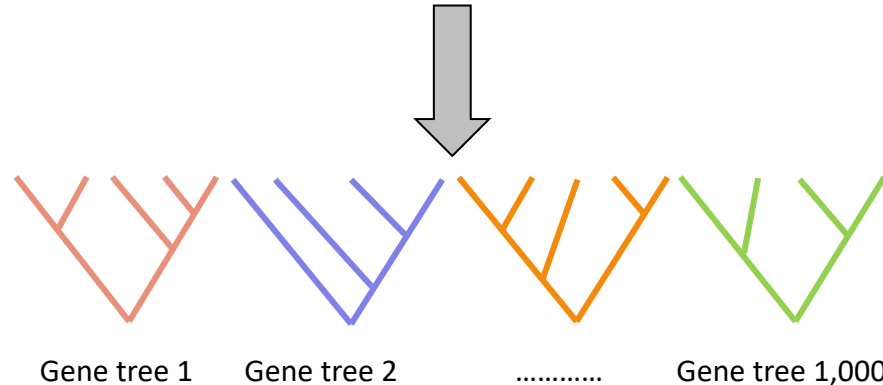
Potential *systematic bias*

Felsenstein (1985):

which not. Where the method of inferring
phylogenies is one with undesirable sta-
tistical properties such as inconsistency,
the bootstrap does not correct for these.

Coalescent/reconciliation methods

Supermatrix			
Gene 1	Gene 2	Gene 1,000
CACCTGTCGT	-----	-----	TCTGGTGCAG
CAGCTGTCGT	GCTCTTTCTG	TTGAGCCTGG	TCTGGTGCAG
CAGCTGCCGT	GTTTTCTCTG	TTGAGCCTGG	TCTGGTACAG
CAGCTGCCGC	GTTCTCTCCG	-----	TCTGGTGCAA
CTCCTGCCGG	GTGCTCTCAG	-----	-----
CTCCTGCCGG	-----	CTGAGCCGGG	TCTGGTGCAG
CTCTTGCCGG	-----	CTGAGCCTTG	-----



Gene Concordance Factor (gCF):
How often a branch in species
tree is found among gene trees?
 $0\% \leq gCF \leq 100\%$

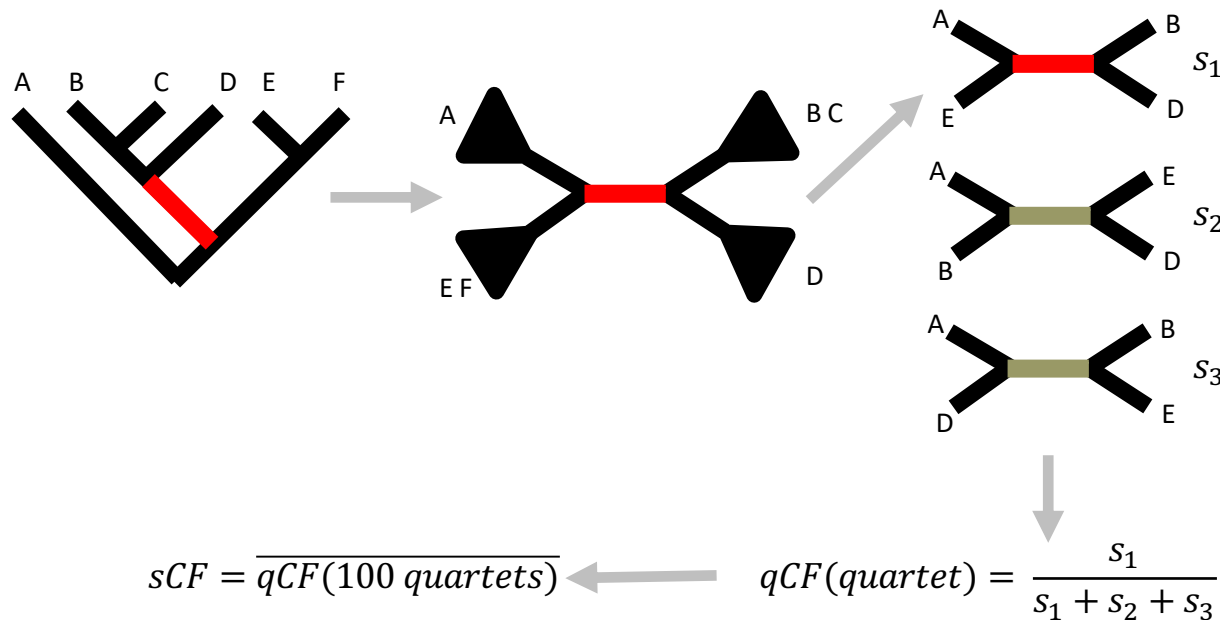
Implementation in IQ-TREE fully
accounts for missing data

**Problem: Uncertainties in
gene trees!**

Site Concordance Factor (sCF)

Supermatrix			
Gene 1	Gene 2	Gene 1,000
CACCTGTCGT	-----	-----	TCTGGTGCAG
CAGCTGTCGT	GCTCTTTCTG	TTGAGCCTGG	TCTGGTGCAG
CAGCTGCCGT	GTTTTCTCTG	TTGAGCCTGG	TCTGGTACAG
CAGCTGCCGC	GTTCTCTCCG	-----	TCTGGTGCAA
CTCCTGCCGG	GTGCTCTCAG	-----	-----
CTCCTGCCGG	-----	CTGAGCCGGG	TCTGGTGCAG
CTCTTGCCGG	-----	CTGAGCCTTG	-----

Site Concordance Factor (sCF):
How often a branch is
“supported” by alignment sites?
 $33.3\% \leq \text{sCF} \leq 100\%$



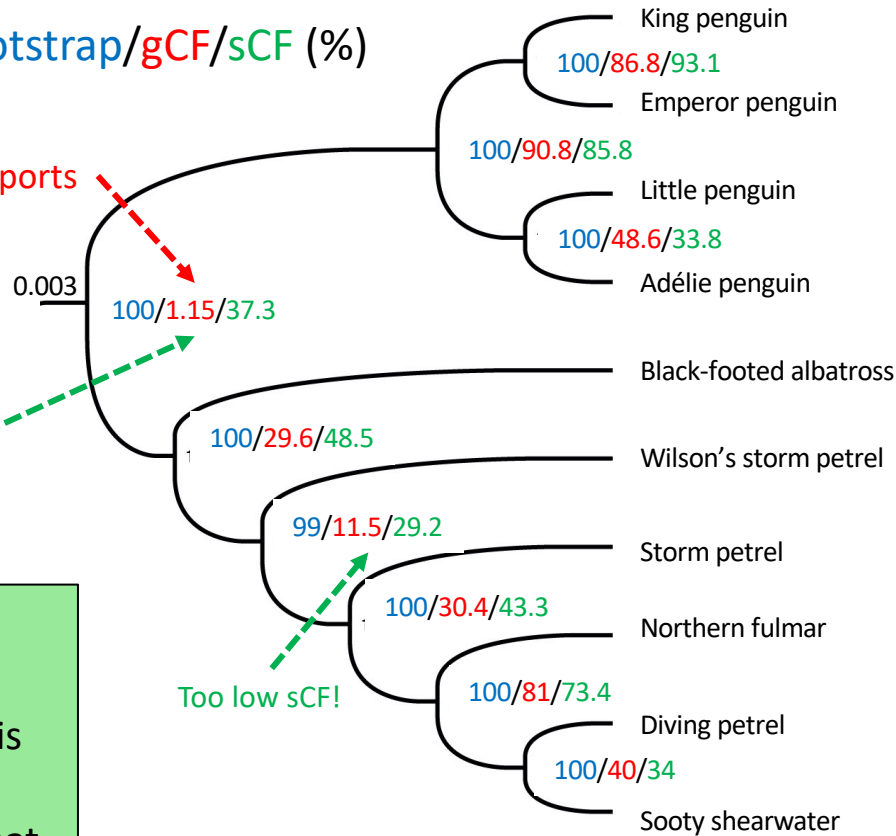
An example birds data set (Reddy et al., 2017)

Bootstrap/gCF/sCF (%)

Only 1 (of 88)
gene tree supports
this branch!

- 131 sites support this branch
- 105 sites support NNI branch 1
- 114 sites support NNI branch 2

Felsenstein (1985): a difference of 20 sites favouring one topology is enough to give 100% bootstrap support for that one topology!



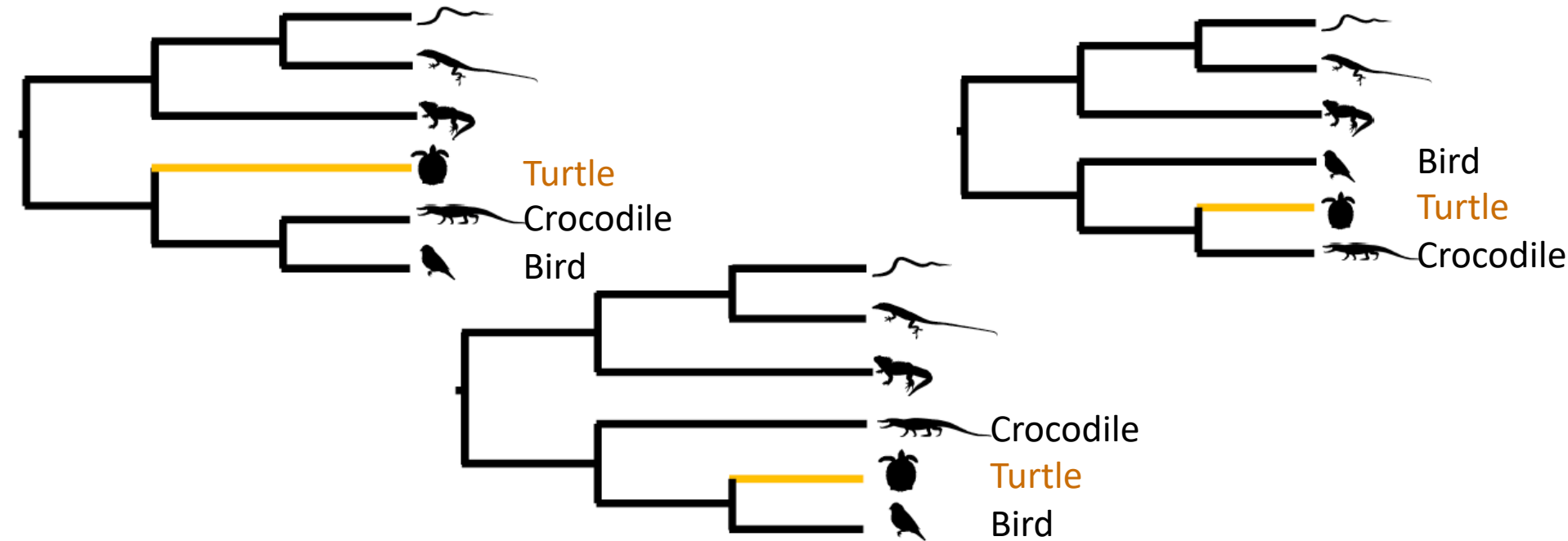
Penguins



Tubenoses

- gCF and sCF are useful when bootstrap supports reach 100%.
- CAUTION when gCF ~ 0% or sCF ~ 33%, even if BS ~ 100%.
- GREAT when gCF and sCF > 50%.

Dataset for IQ-TREE lab: Where is Turtle in the tree?



Chiari et al.
Crawford et al.
Fong et al.

Wang et al.
Lu et al.
Shaffer et al.

2012 2013 2014

Different studies led to different trees!

Dataset: 16 species, 29 genes,
20,820 bp
(a subset of Chiari et al. 2012)

Thanks Jeremy Brown

1. Input data
2. Inferring the first phylogeny
3. Applying partition model
4. Choosing the best partitioning scheme
5. Tree topology tests
6. Identifying most influential genes
7. Removing influential genes
8. Concordance factors (*advanced)

<http://www.iqtree.org/workshop/molevol2022>

Fill out your answers in a Google form (shared via Slack)