### Phylogenetics and the CBGM

@ CSNTM

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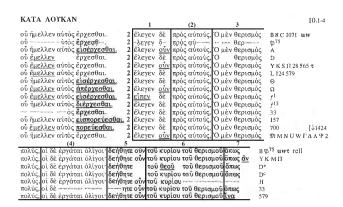


## **Preliminaries**





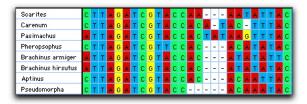
- To compare textual witnesses, align them at independent *variation units*
- Variant readings occur at variation units



Collation of Luke 10:2 with variation units numbered above text (Reuben J. Swanson, ed., New Testament Greek Manuscripts: Variant Readings Arranged in Horizontal Lines against Codex Vaticanus. Luke [Sheffield: Sheffield Academic Press, 1995], 183)



• Analogous to a DNA sequence alignment



- Rows: *taxa* = witnesses
- Columns: *sites* = variation units
- Cells: *states* = variant readings (including omissions)
  - Lacunae and uncertain retroversions correspond to fully or partially ambiguous states



 At the most basic level, a witness is just a sequence of readings, a row in the collation

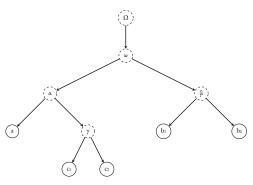
|         | 3Jo 1:1/2 | 3Jo 1:1/6 | 3Jo 1:1/8 | <br>3Jo 1:15/23 |
|---------|-----------|-----------|-----------|-----------------|
| GA 69   | a         | afl       | a         | <br>a           |
| GA 1739 | a         | a         | b         | <br>a           |
| GA 2243 | ь         | a         | a         | <br>a           |

- Paratextual features can be encoded in the same way
- In more complex phylogenetic approaches, age can also be incorporated

## Phylogenetics

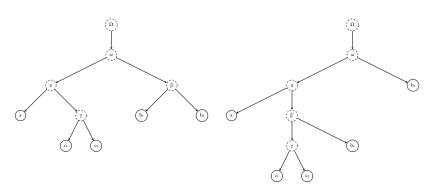






- A stemma is a "family tree" modeling the transmission of the text
- The leaves (solid circles) correspond to extant witnesses
- The hyparchetypes (dashed circles) are hypothetical (now-lost) ancestors, reconstructed from their descendents along the branches
- The *archetype* ( $\omega$ ) is the earliest reconstructible text
- The  $root(\Omega)$  represents the authorial text





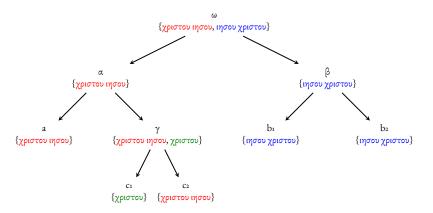
- A stemma represents a hypothesis about transmission history
- The goal is to determine which hypothesis (or hypotheses) best explain the extant data
- To do this, we need a numerical metric for the fitness of a given stemma



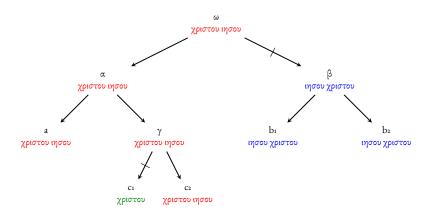
- One such metric is *parsimony*
- Cost = smallest number of changed readings along the branches of the stemma
- Motivated by Ockham's Razor
- Given a candidate stemma, we calculate its cost for each variation unit independently and add up the results
- We calculate it starting at the bottom of the stemma and working our way up

| Witness        | Reading       |
|----------------|---------------|
| a              | χριστου ιησου |
| b <sub>1</sub> | ιησου χριστου |
| $b_2$          | ιησου χριστου |
| C1             | χριστου       |
| C2             | χριστου ιησου |

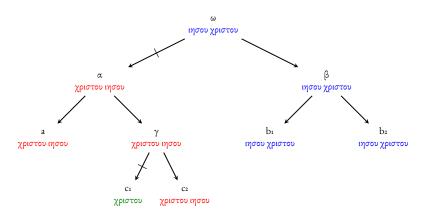




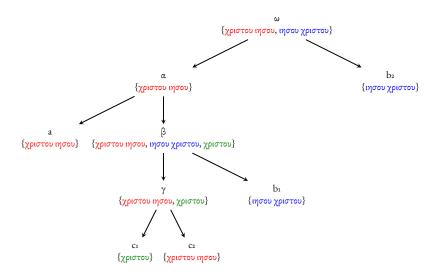




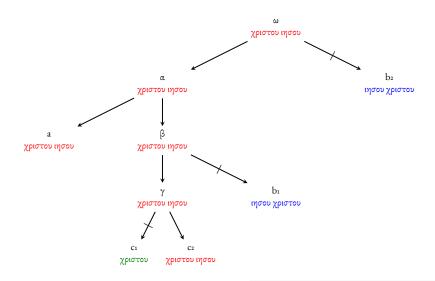




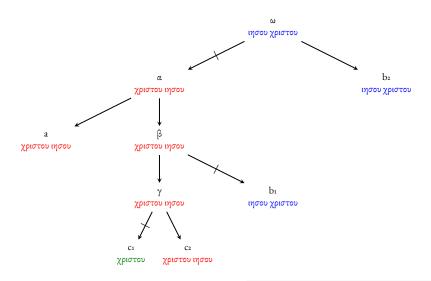








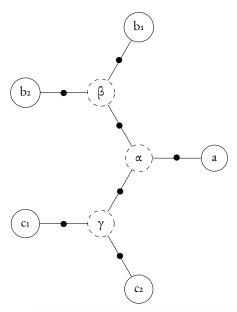




#### **Phylogenetics** | Parsimony

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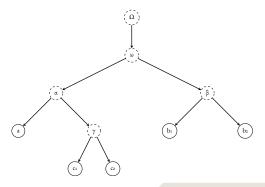
- Minimum number of changes is the same regardless of what the archetype reads
- A stemma's cost does not depend on where its root is
- This means the computer can calculate the costs of stemmata without knowledge about the root, and we can postpone the assessment of internal evidence of readings to the end, when we want to determine where the root of the tradition is
- The traditional approach for computer-assisted stemmatics



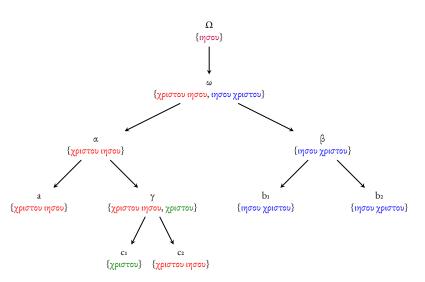
#### **Phylogenetics** | Extensions



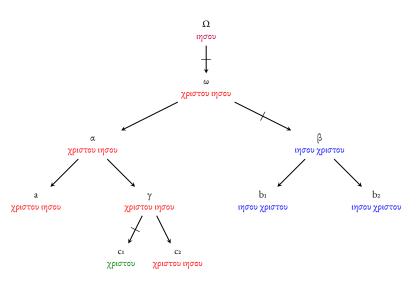
- But we may want to incorporate internal evidence up front:
  - Separation of concerns between intrinsic probabilities ("what would the author write?") and transcriptional probabilities ("how would later scribes/readers change it?")
  - Intrinsic probabilities can affect the backward pass
  - Transcriptionally, some changes are more likely than others or irreversible
- We can extend the approach to incorporate this evidence, but the stemma costs will now depend on where the root of the tradition is located



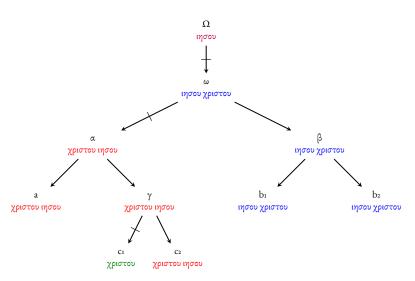




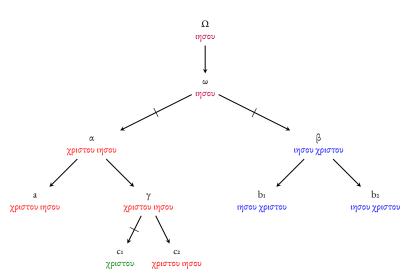










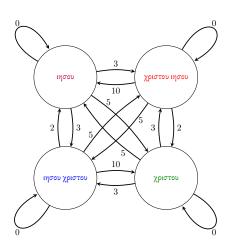


#### **Phylogenetics** | Extensions

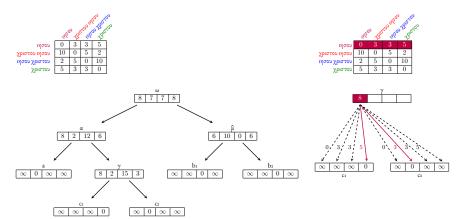


- We can likewise assign weighted costs to different transitions between readings using a *cost matrix*
- A model of the average scribe's behavior

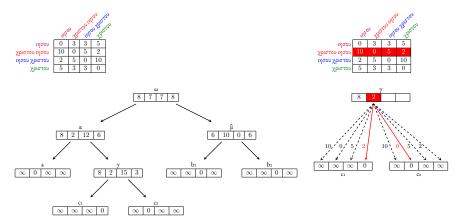
|               |    | ა .   | roote | 50° 1210°C | oo<br>oo |
|---------------|----|-------|-------|------------|----------|
|               | 40 | , 48, | , 46  | 120        |          |
| ιησου         | 0  | 3     | 3     | 5          |          |
| χριστου ιησου | 10 | 0     | 5     | 2          |          |
| ιησου χριστου | 2  | 5     | 0     | 10         |          |
| χριστου       | 5  | 3     | 3     | 0          |          |
|               |    |       |       |            |          |



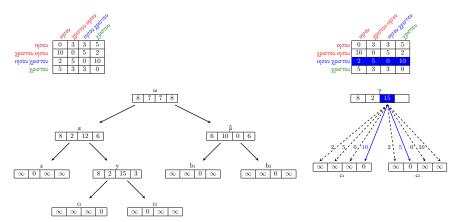




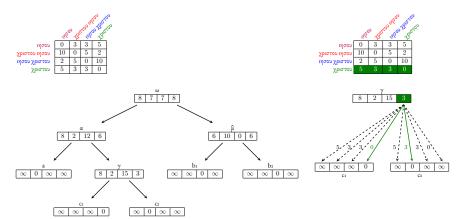




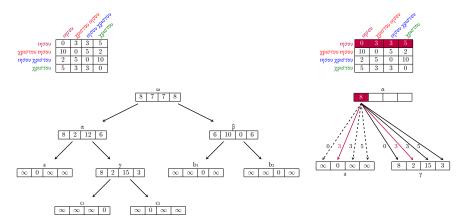




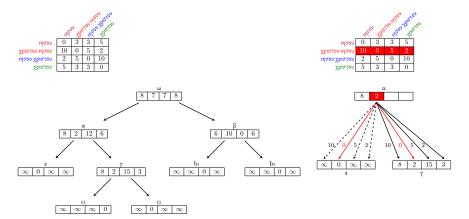




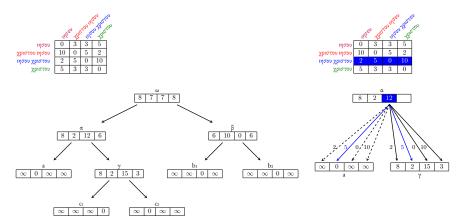




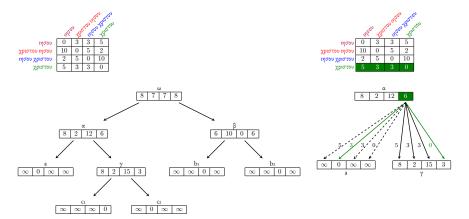








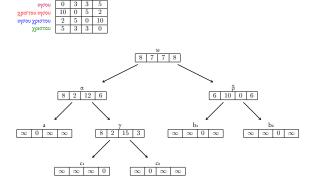




#### **Phylogenetics** | Extensions



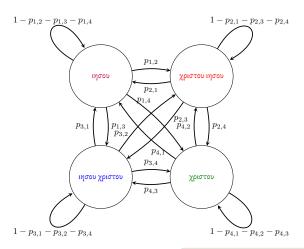
- We can also incorporate intrinsic evidence
- If the root reading is  $\eta \sigma ov$ , then the minimum cost of the stemma is  $\min(0+8,3+7,3+7,5+8) = \min(8,10,10,13) = \mathbf{8}$
- If the root reading is  $\chi\rho\iota\sigma\tau ov\ \iota\eta\sigma ov$ , then the minimum cost is  $\min(10+8,0+7,5+7,2+8)=\min(18,7,12,10)=7$



#### **Phylogenetics** | Extensions

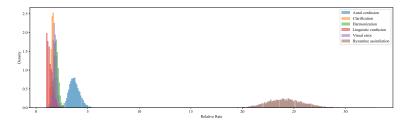


- We can even use probabilities for intrinsic and transcriptional evidence
- Intrinsic probabilities = *prior probabilities* at root
- Transcriptional probabilities = probabilities of copying the same reading or changing it to another reading, modeled as a *Markov chain*





- In a probabilistic setting, we can incorporate and estimate other parameters of interest:
  - Probabilities for classes of transitions between readings (scribal habits)
  - Lengths of branches (how many copying events separated an ancestor from a descendent, and how error-prone were the scribes involved?)
  - Dates of witnesses (using clock models)
  - Measurements of how certain we can be about the best stemmata found in the process
- More complex, but feasible with modern computers!

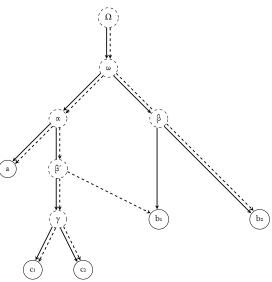


# CBGM





• Developed to solve *contamination*, or mixture across branches of the textual tradition

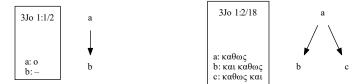




- Foundational principles:
  - 1. Scribes typically copied their exemplars with fidelity.
  - 2. If a scribe introduced a variant, then it came from some other reading.
  - 3. Scribes typically used fewer sources rather than many.
  - 4. Scribes typically used closely related sources rather than distant ones.
- Witnesses are texts (sequences of readings), minus the material baggage (date, provenance, etc.)
  - "How texts relate" \( \neq \) "How manuscripts relate"
- No hypothetical ancestors (except for the Ausgangstext A)
  - Contamination would (presumably) hinder their reconstruction
  - Instead, we use extant witnesses as proxies for different states of the text
  - This yields a much smaller and more manageable space of solutions compared to the space of all possible stemmata for a given set of witnesses



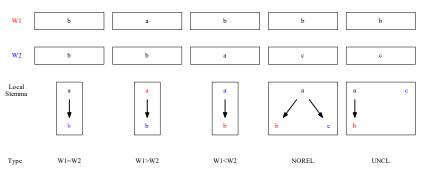
- The basic unit of comparison
- One for each variation unit
- A graphical representation of our judgments of readings
- Similar to cost graphs in function, but in principle, represents a judgment about what *did* happen, not what *could* happen
- · Thus, no bidirectional edges or cycles like we have in cost graphs



## **CBGM** | Genealogical Relationships



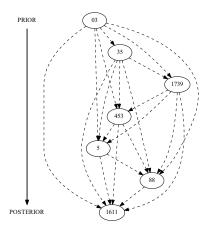
• The relationship of two witnesses is the overall pattern *of the relationships of their readings* at all variation units where both are extant



The first three are the most important



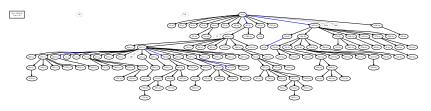
• Potential ancestor = "more prior than posterior readings"



### **CBGM** | Textual Flow

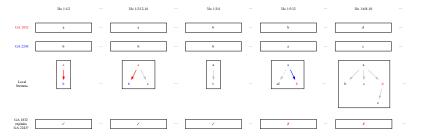


- Textual flow diagram = a tree that relates each witness to its closest potential ancestor, with as few changes in reading as possible
- Similar to a most-parsimonious stemma for a specific variation unit
- We specify a *connectivity limit*  $\kappa$  (i.e., a radius of "close-enough" neighbors)
- Then, for each witness:
  - 1. List its potential ancestors, sorted from most agreement to least
  - 2. If one of the first  $\kappa$  has the same reading at this unit, pick the first that does
  - 3. If not, pick the first (non-lacunose) potential ancestor
- Core idea: use *general relationships* between witnesses to find *specific relationships* between readings, so local stemmata can be refined





- We say that one reading *explains* another if
  - it is the same reading ("explanation by agreement"), or
  - there is an edge in the local stemma from it to the other reading ("explanation by descent")

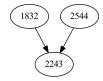


Lacunae do not have to be explained, and they cannot explain readings



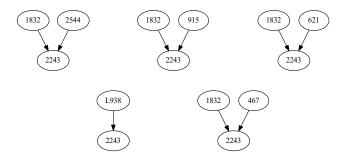
- The *substemma* of a witness is the portion of the global stemma consisting of the witness and its ancestors in the stemma
- Requirement: every extant reading in the witness must be explained by a reading in at least one of its ancestors

| Explained by GA 1832 | <br>X | 1 | 1 | 1 |  |
|----------------------|-------|---|---|---|--|
| Explained by GA 2544 | <br>1 | Х | Х | 1 |  |
| Explained by Either  | <br>✓ | 1 | 1 | 1 |  |



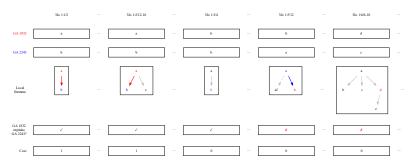


- A witness may have multiple valid substemma (i.e., ones that explain all of its readings), but some are better than others
- Two of the CBGM's methodological assumptions are important here:
  - 3. Scribes typically used fewer sources rather than many.
  - 4. Scribes typically used closely related sources rather than distant ones.
- Thus, we need a cost function to distinguish between candidate substemmata





- A simple cost function: "the number of variation units where the ancestor explains the witness by descent and not agreement"
- Thus, in the example below, GA 1832 is a stemmatic ancestor of 2243 with a cost of 2 (but it cannot be its only stemmatic ancestor)



## **CBGM** | Substemma Optimization

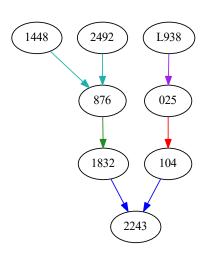


- The process of finding the best substemma for a witness
- $\bullet\,$  For n potential ancestors, a weighted set cover problem with n sets
- $2^n-1$  possible combinations to check (!), but fast heuristics exist

| Substemma | Variation | Cost     |          |          |         |
|-----------|-----------|----------|----------|----------|---------|
|           |           |          |          |          |         |
| {A}       | ✓         | <b>√</b> | <b>✓</b> | <b>✓</b> | 4       |
| {B}       | <b>√</b>  | 1        | X        | X        | 1       |
| {C}       | X         | 1        | <b>✓</b> | ✓        | 2       |
| {A, B}    | ✓         | 1        | <b>✓</b> | ✓        | 4+1=5   |
| {A, C}    | 1         | 1        | 1        | 1        | 4+2=6   |
| {B, C}    | <b>√</b>  | <b>√</b> | <b>√</b> | <b>√</b> | 1+2=3   |
| {A, B, C} | 1         | 1        | 1        | 1        | 1+2+4=7 |

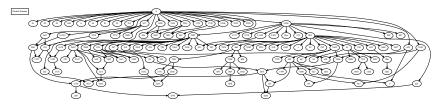


- Just as the local stemma relates readings, the global stemma relates witnesses
- Combination of all substemmata into a single graph
- Analogous to a phylogenetic stemma





- How does this differ from a phylogenetic stemma?
  - In practice, it takes much less time to produce a global stemma (minutes) than it does to do a satisfactory phylogenetic search for promising stemmata (> a day)
  - Converging branches (reflecting contamination) are allowed
  - No hyparchetypes, and texts found in later manuscripts can be ancestors to texts found in earlier manuscripts
  - Can it model a history of the text?



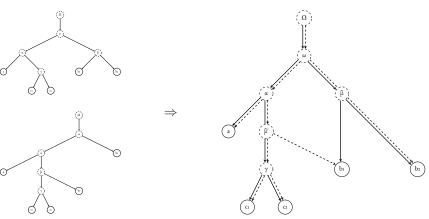
## Conclusions



#### Conclusions



- Can we get the advantages of both approaches?
- The CBGM could use cost graphs instead of local stemmata
- Phylogenetics could use the local-genealogical principle to model contamination
  - Different low-cost stemmata at different variation units



# Questions?

