

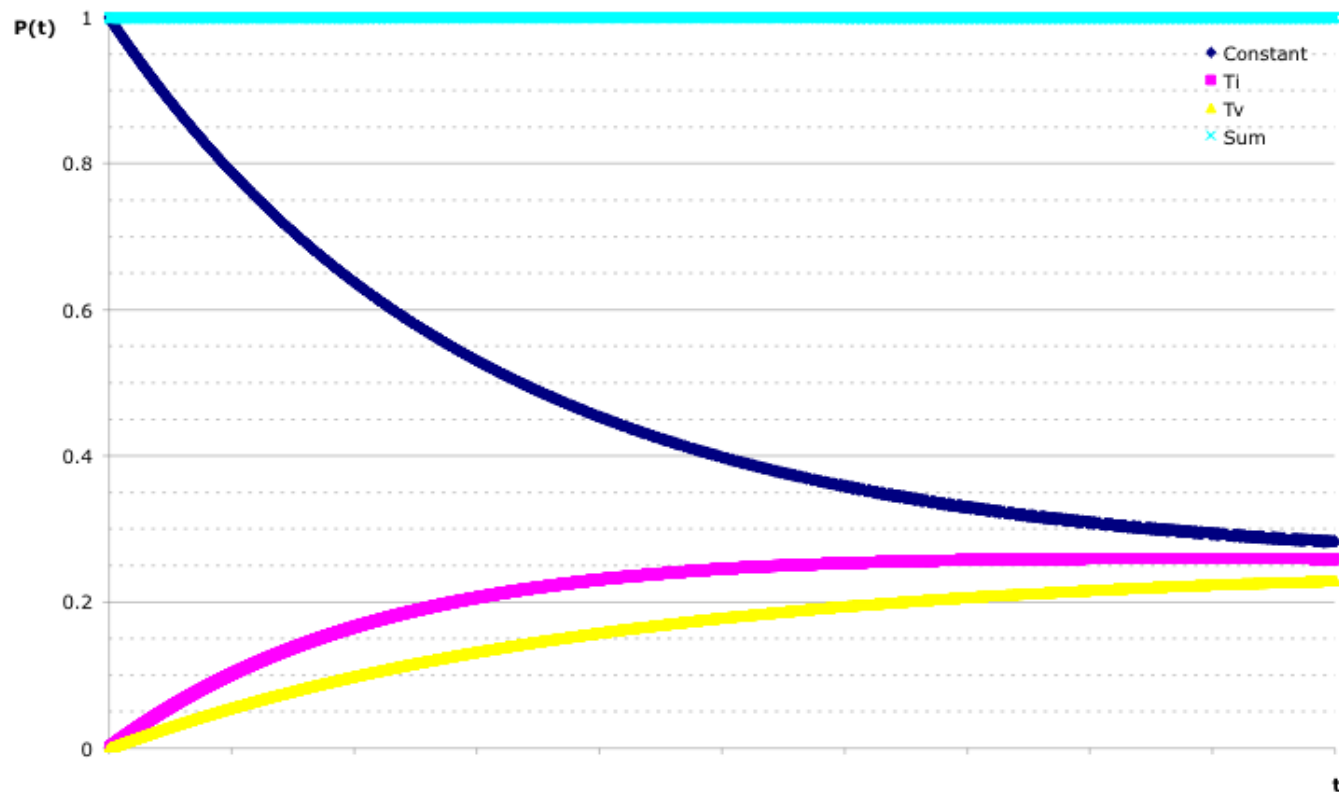


# *Evolutionary rate & saturation*



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**NHM UiO**

# *Increased chance of substitution*

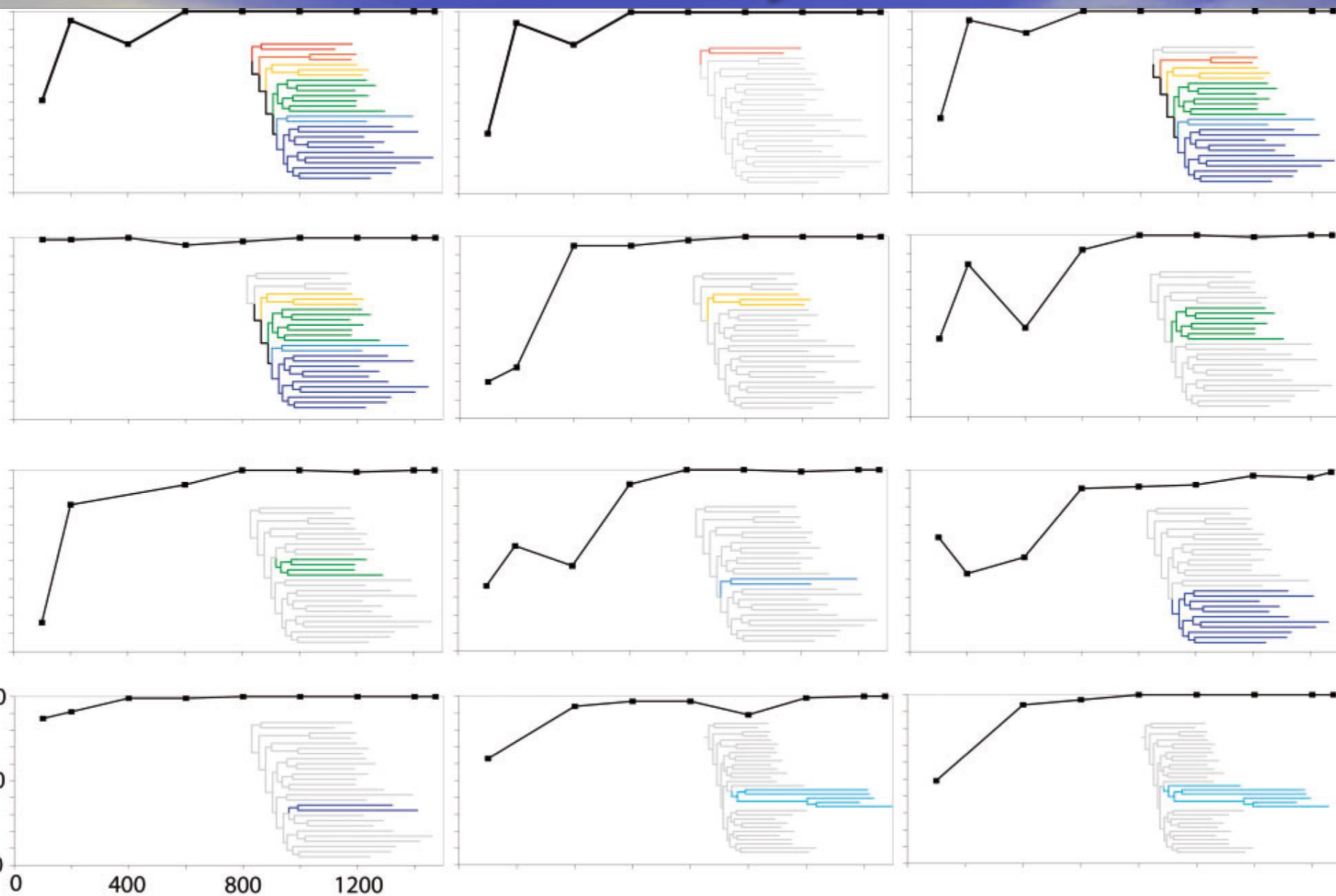


$$\mu = 0.25$$

$$\kappa = 2$$



# *Evolutionary rate*

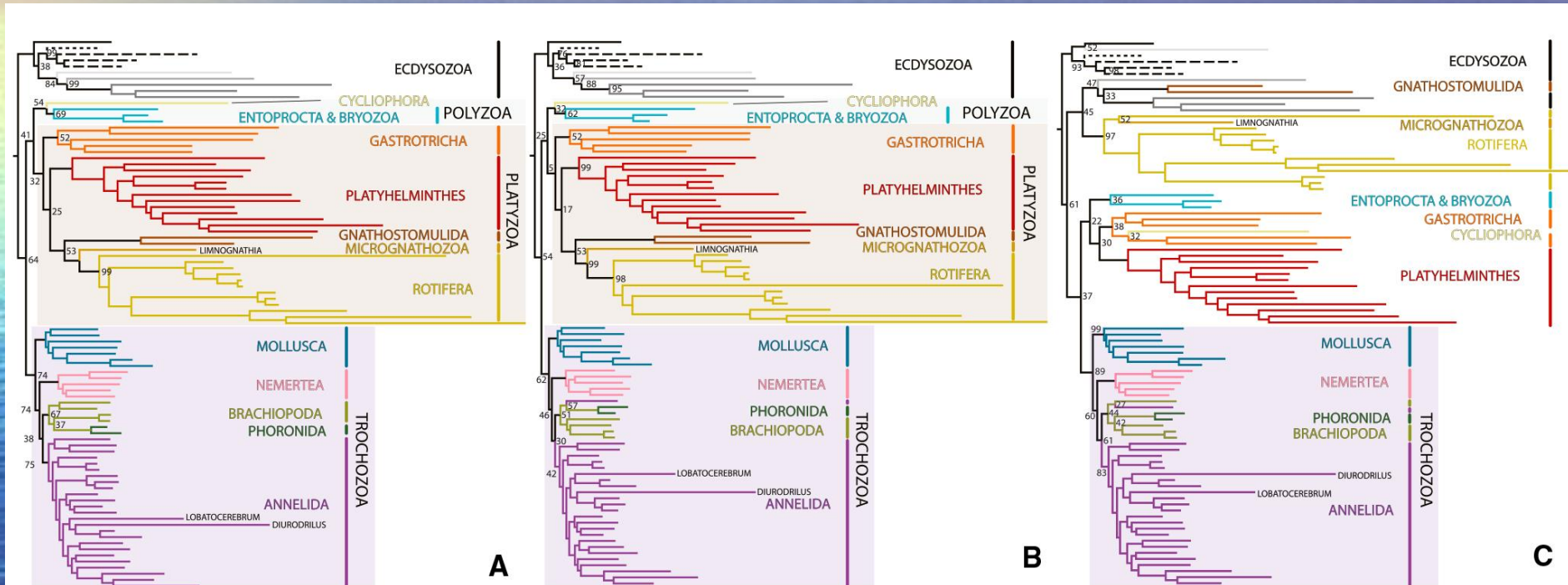


# Evolutionary rate

All

75% slowest

75% fastest

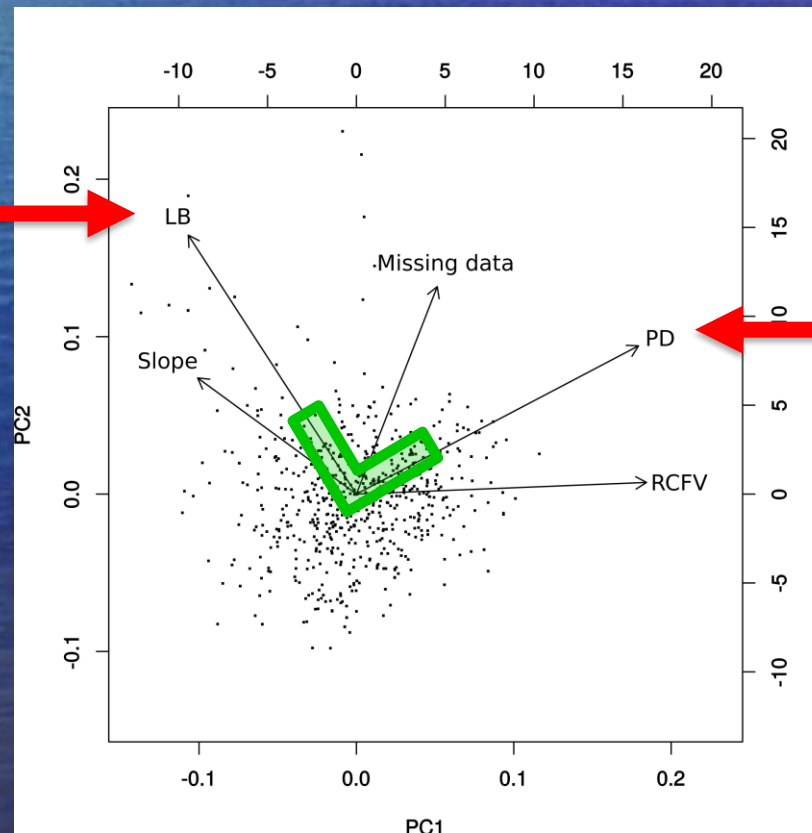


Laumer et al. (2015) Curr. Biol.

# *Evolutionary rate & Long branch attraction*

Overall evolutionary rate of a gene is often taken as a proxy for long branch attraction.

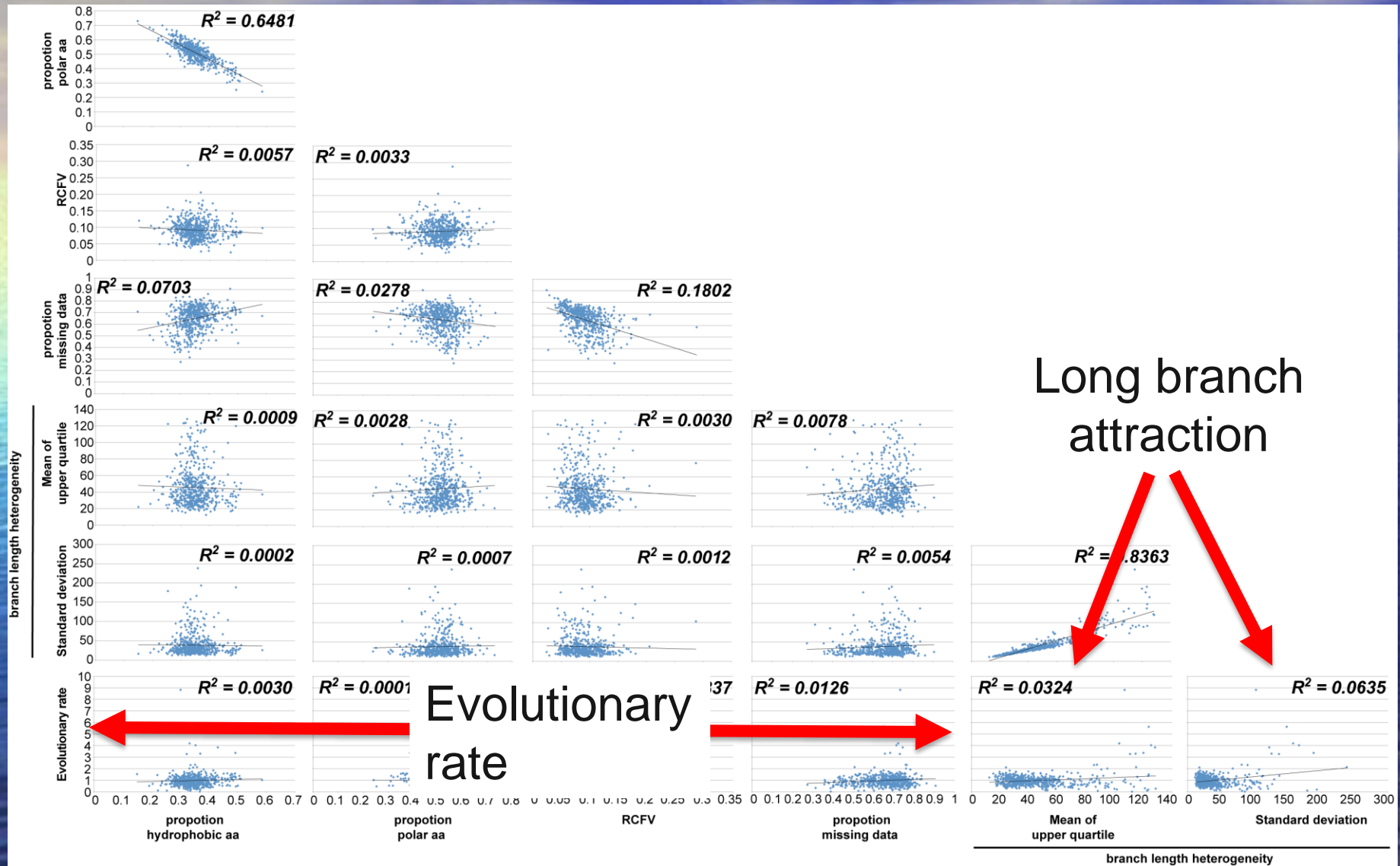
Long branch attraction



Evolutionary rate



# Evolutionary rate & Long branch



# *Evolutionary rate & Long branch attraction*

**Long branch attraction is caused by differences in branch lengths and NOT by an overall increased substitution rate.**

# *Saturation*

How can substitutions saturate?

Transitions are more likely to occur than transversions.

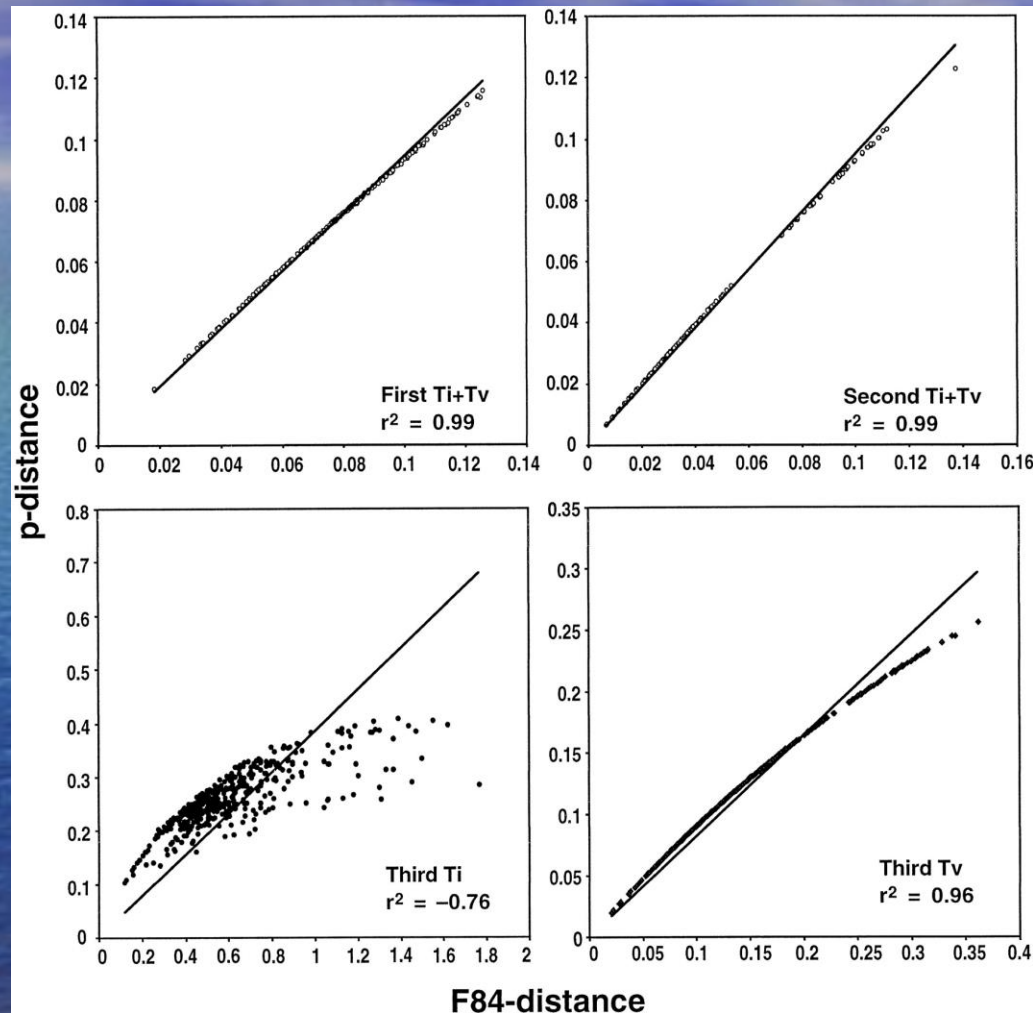
At a certain observed genetic divergence between sequences multiple substitutions become increasingly likely.

Because transitions occur more often than transversions this effect will be seen much earlier in transitions leading to saturation:

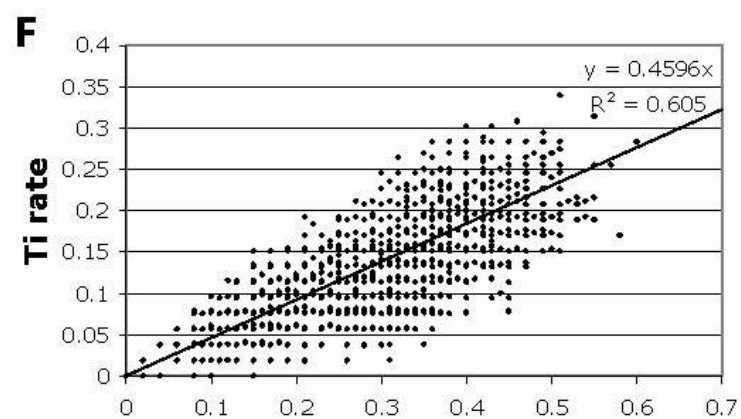
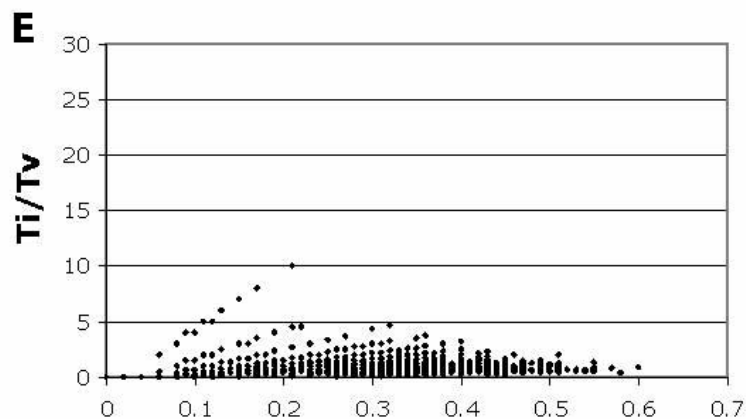
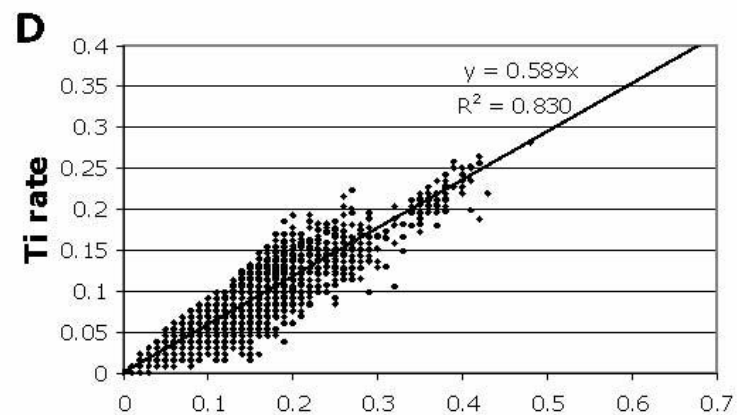
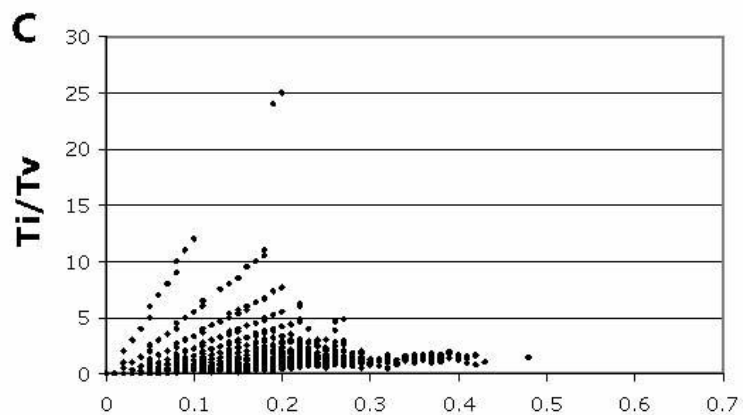
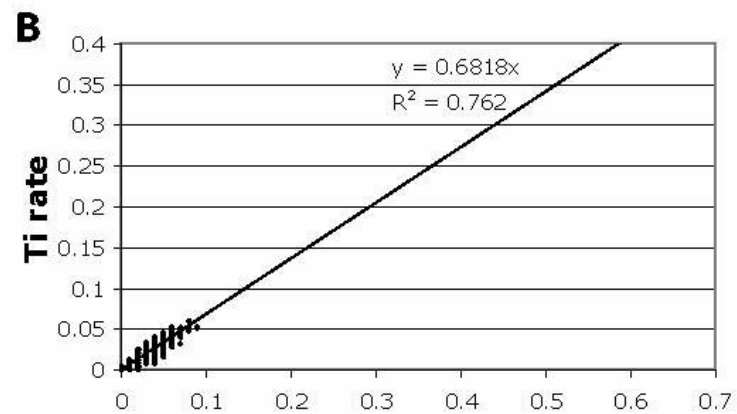
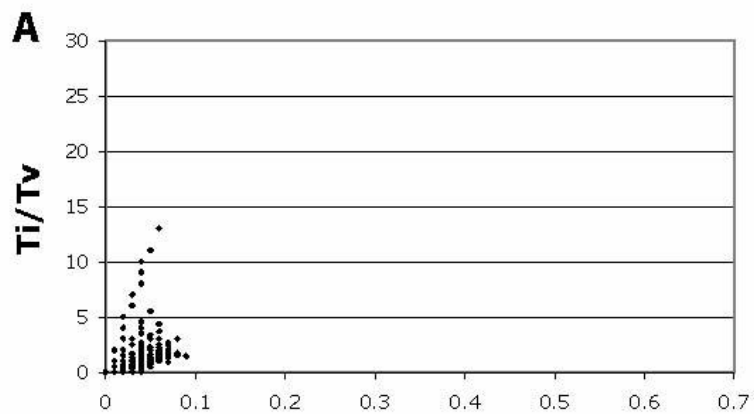
*While the genetic divergence is still increasing the transition rate is not.*



# Saturation



First, second, and third codon positions in *rbcL* of land plants  
(Nickrent et al. 2000 Mol. Biol. Evol. 17:1885-1895)



# *C (=convergence) index*

**What are the properties of convergence?**

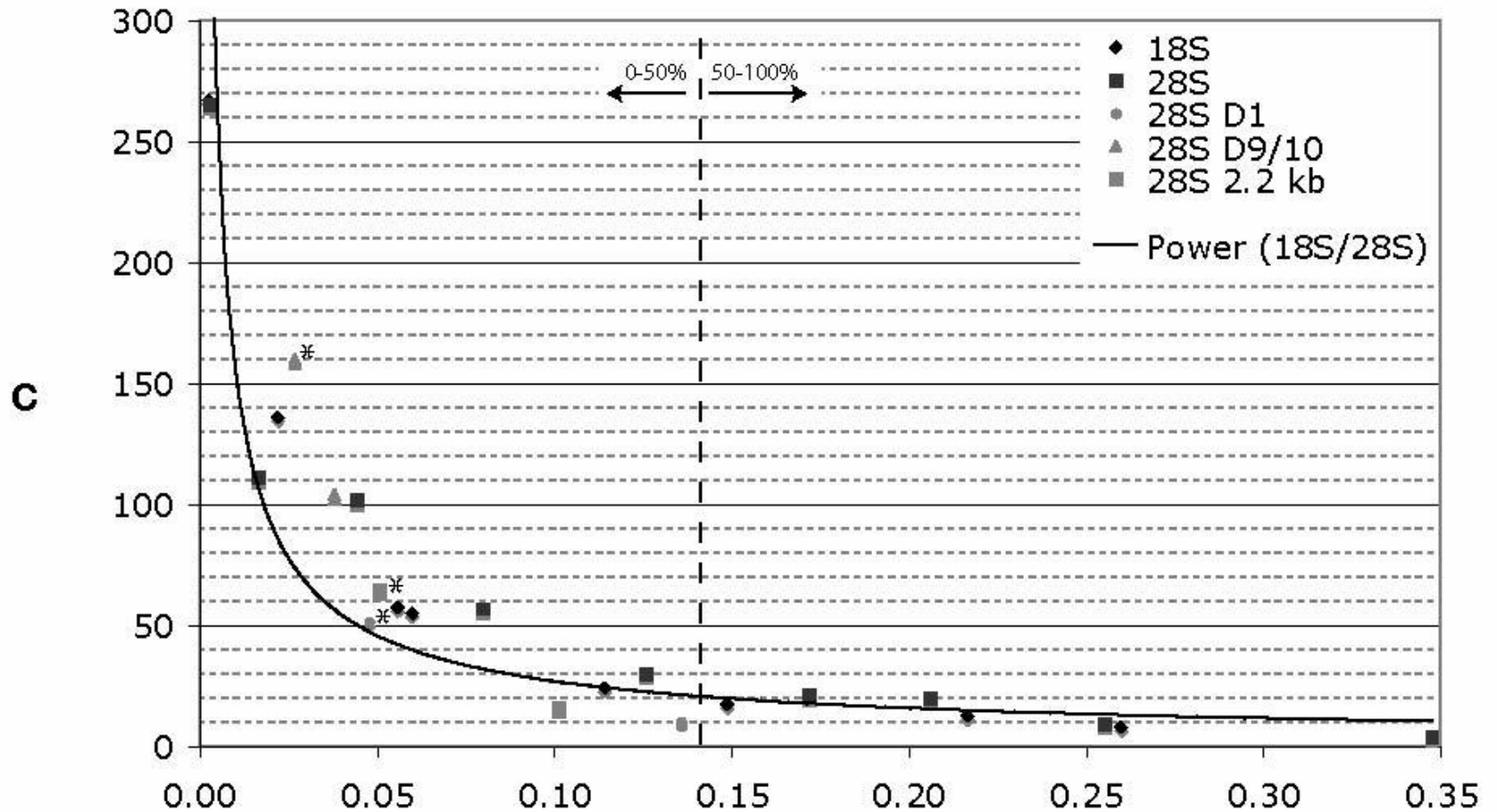
One factor is increasingly spread out, whereas the other becomes more and more condensed around a single value. Therefore, the standard deviation should increase as well as decrease, respectively.

$$C = \frac{\sigma(\frac{ti}{tv})}{\sigma(p)}$$

Struck et al. (2008) MPE



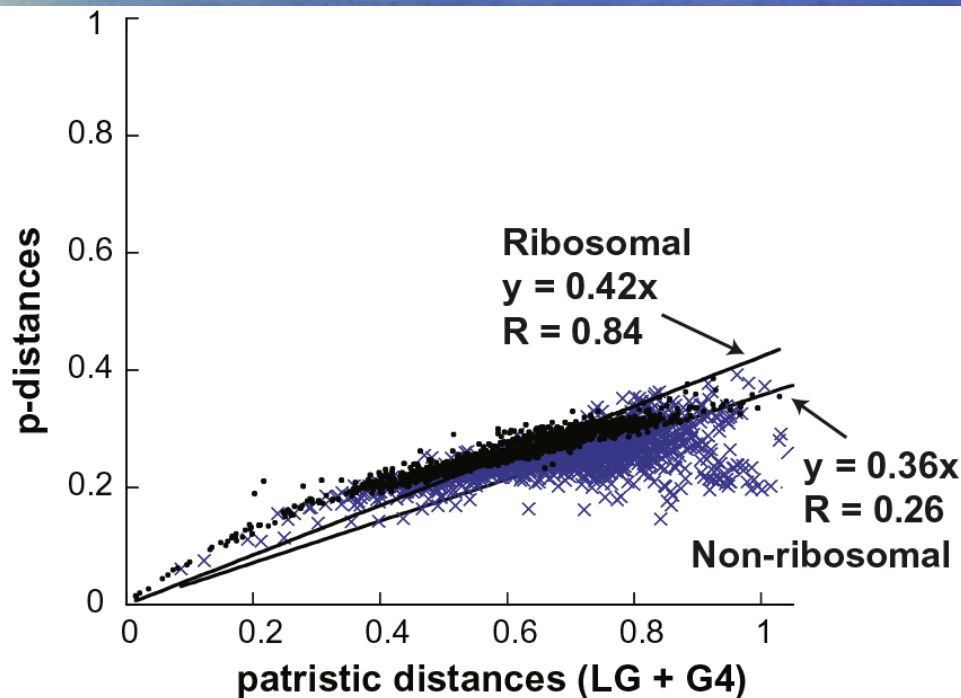
# *C (=convergence) index*



# *R<sup>2</sup> and slope*

## Deep metazoan phylogeny: When different genes tell different stories

Tetyana Nosenko<sup>a</sup>, Fabian Schreiber<sup>b</sup>, Maja Adamska<sup>c</sup>, Marcin Adamski<sup>c</sup>, Michael Eitel<sup>d,1</sup>, Jörg Hammel<sup>e</sup>, Manuel Maldonado<sup>f</sup>, Werner E.G. Müller<sup>g</sup>, Michael Nickel<sup>e</sup>, Bernd Schierwater<sup>d</sup>, Jean Vacelet<sup>h</sup>, Matthias Wiens<sup>g</sup>, Gert Wörheide<sup>a,i,j,\*</sup>



**Fig. 3.** Saturation analysis. The relative saturation levels were estimated for the ribosomal and non-ribosomal gene matrices containing 50 taxa by computing the Pearson correlation coefficient  $R$  and slope of the regression line of patristic vs.  $p$ -distances. The patristic distances between pairs of taxa were inferred from the branch lengths of ML trees constructed under the LG +  $\Gamma$ 8 + I model.

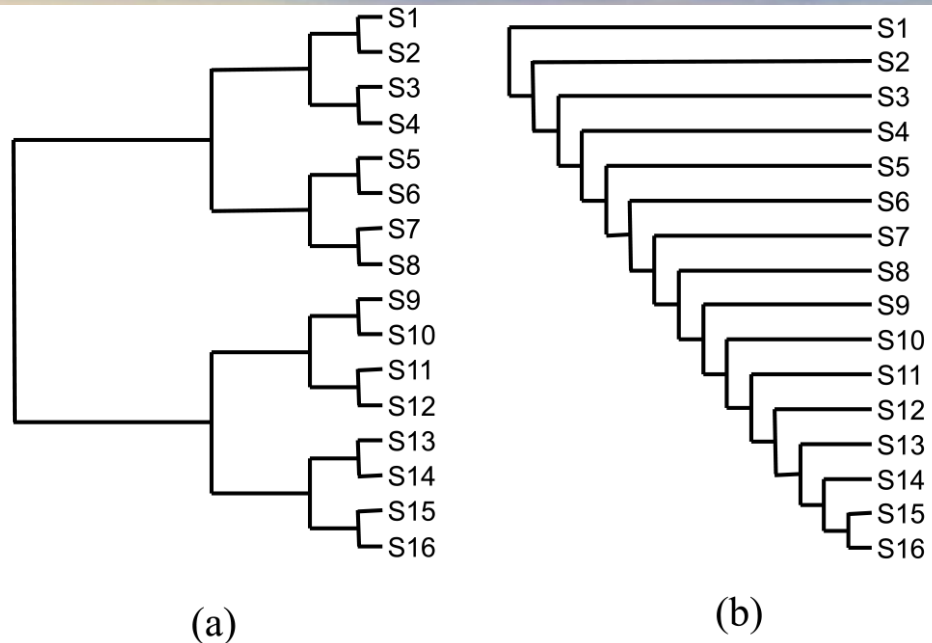
# $I_{ss}$

## An index of substitution saturation and its application

Xuhua Xia,<sup>a,d,\*</sup> Zheng Xie,<sup>b</sup> Marco Salemi,<sup>c</sup> Lu Chen,<sup>d</sup> and Yong Wang<sup>d</sup>

- ancestral state reconstruction of DNA sequences and comparison with simulated data
- estimation of substitution numbers of neighboring nodes along the tree
- $I_{ss}$  is the ratio of the average information entropy of positions to the expected entropy of the alignment
- the smaller  $I_{ss}$ , the less substitution saturation has occurred





For simulation, only two possibilities of tree topology are considered. The two extremes:

- Absolutely symmetrical
- Absolutely asymmetrical

Fig. 1. Two extreme topologies used in simulation: (a) symmetrical; (b) asymmetrical.

# $I_{ss}$

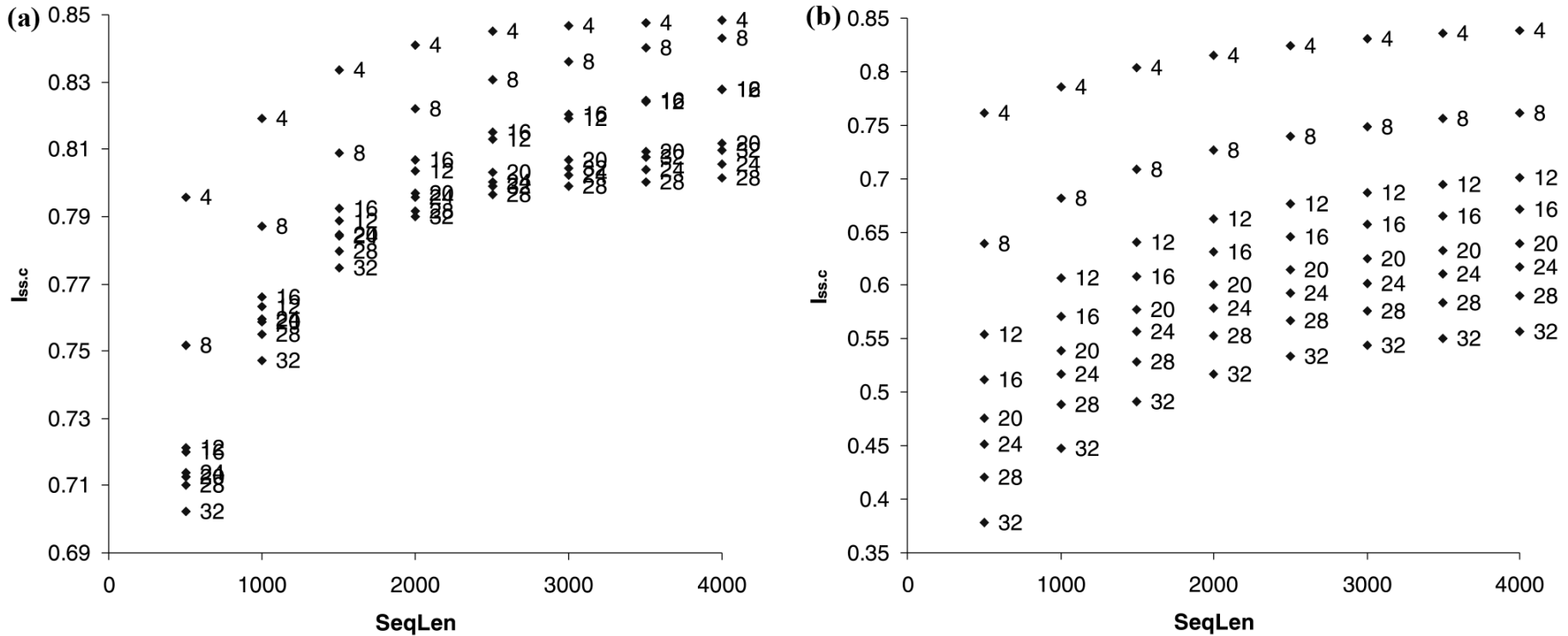


Fig. 3. The critical index of substitution saturation ( $I_{ss,c}$ ) depends on the sequence length (SeqLen) and the number of OTUs ( $N_{OTU}$ ). Data labels are  $N_{OTU}$  values: (a) with a symmetrical topology; (b) with an asymmetrical topology.

These are then used for all similar datasets to get the critical  $I_{ss}$  value indicating saturation.