

# **Adaptation in genes, duplicates, families, functional modules and genomes**

François Serra

October 2011



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# Nomenclatura

BWT Burros-Wheeler transform

CR Complexity Ratio

# **1. Introduction**

- 1.1. Adaptive changes to evolutionary speed**
- 1.2. Evolution, and the detection at molecular level**
- 1.3. Grouping genes and finding evolutionary patterns**
- 1.4. What is DNA? How genes rose?**
- 1.5. Life in DNA, from genes to repetitive elements.**

## **Part I.**

# **Structure and dynamics of genomes**



## **2. Random-like structure of DNA**

### 2.1. Introduction

From a biological perspective it seems obvious that DNA is something else than random mix of A, T, G or C nucleotides. Genomes are composed of functional elements as can be genes or promoters but also repetitive elements that by definition can not be random when taken together. However to what extent can we state that genomes are not a random soup of 4 letters?

This question could be solved in some sense by measuring genomes entropy. This measure presents the disadvantage that extreme cases of high entropy could correspond to *a) a specially high content of information*, entropy-based algorithms are actually used to predict or confirm automatic detection of genes [Du *et al.*2006, Gerstein *et al.*2007], *b) an exact random structure*, some work in the sense of testing the random structure of DNA have been done using entropy [Loewenstern & Yianilos1999]. However this characteristic of entropy could be only a semantic problem if we use it as a measure of relative variation in DNA complexity in genomes, and try to discern statistical patterns in the DNA sequences of different genomic element such as interspersed repeats or functional element (like protein-coding genes). This kind of description of DNA sequence complexity was already done by [Holste *et al.*2001], but only in human chromosome 22.

### 2.2. Results and Discussion

### 2.3. Material and methods

#### 2.3.1. The complexity ratio and complexity value

Complexity Ratio CR is defined by a classical formula used in data compression [Adjero *et al.*2008], the Burros-Wheeler transform BWT [Burrows & Wheeler1994].

#### 2.3.2. Complexity in strings

#### 2.3.3. Simulations

### **3. Life inside genomes, dynamics and predictions**

*3. Life inside genomes, dynamics and predictions*

**3.1. Genomic elements, dispersion and abundance**

**3.2. Species Abundance Diversity in genomes**

**3.3. Neutrality of SAD**

**3.4. Material Methods**

**3.4.1. Ecology**

## **Part II.**

# **Detection of selective pressures in genomes**

#### **4. Searching for evolutionary patterns in functionally linked group of genes**

## 4.1. Introduction

## 4.2. Material and Methods

### 4.2.1. Dataset

#### Five mammals

Complete genomes of 5 mammals species (*Homo sapiens*, *Pan troglodytes*, *Mus musculus*, *Rattus norvegicus* and *Canis familiaris*) were retrieved from *Ensembl* [Flicek *et al.*2011]. Also orthology prediction between each pair of species possibly done between human and the others was retrieved from *Ensembl Compara* [Vilella *et al.*2009] using biomaRt [Kinsella *et al.*2011]. Only groups of orthologs *one-to-one* with one representative of each species were kept in the final dataset. 4.1 NUMBERS

#### 6 Drosophila

### 4.2.2. Alignments

Each of the group of orthologous sequences were aligned with Muscle [Edgar2004], and, once aligned sequences were cleaned with trimAl [Capella-Gutiérrez *et al.*2009] keeping all sequences but trimming alignment columns with the euristic1 method.

## 4.3. open on colocalization to not random

4. Searching for evolutionary patterns in functionally linked group of genes

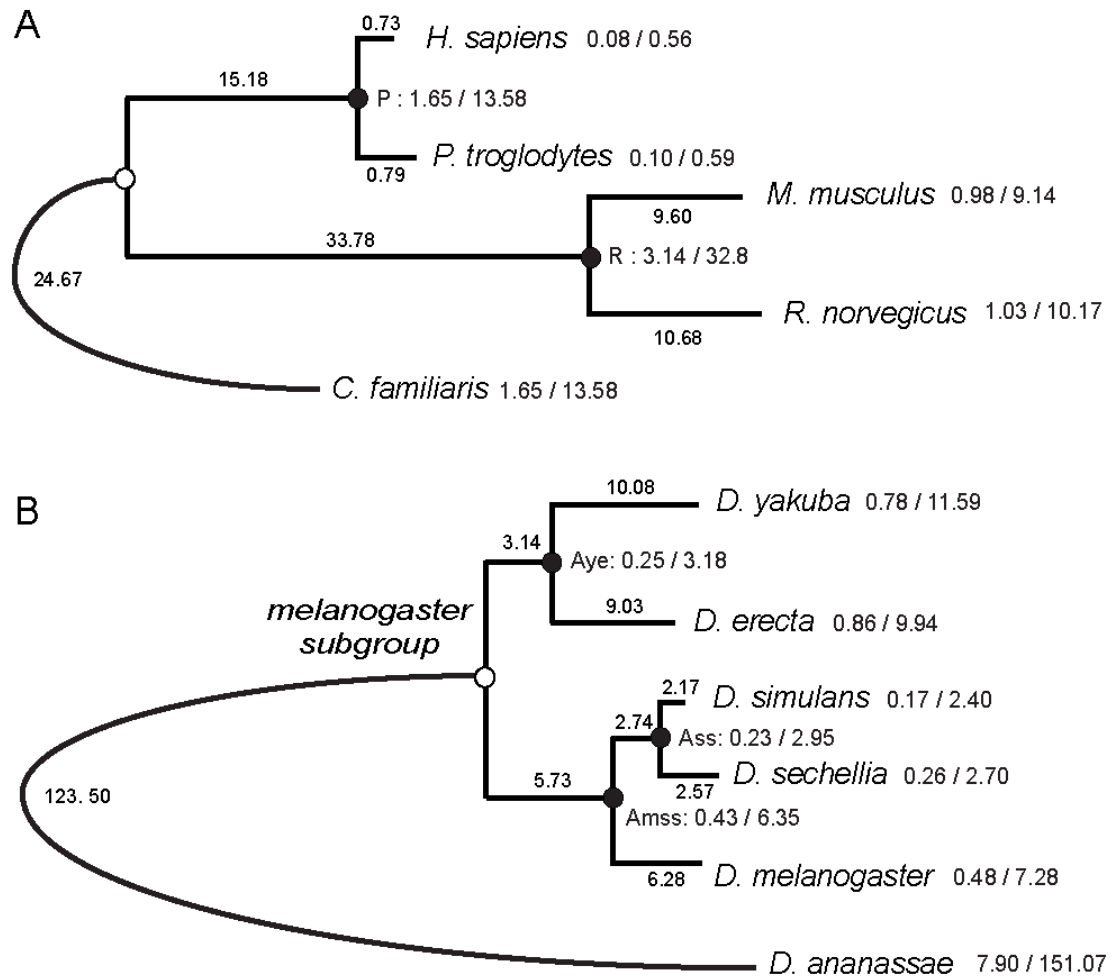


Figure 4.1.: Mammals and *Drosophila* phylogeny. blabli blob lu dkfnlskjdf



## **5. Tools, programs, methods**

*5. Tools, programs, methods*

**5.1. ETE-evol plugin**

**5.1.1. BRANCHED1**

**5.1.2. Protamines Rodents and Primates**

**5.2. Pipeline for study of adaptation at genomic scale**

**5.2.1. Selective pressure on duplicated genes in *Drosophila***

**5.3. Phylemon**

## 6. Conclusions

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