

Research Staff

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Research Associates and PhD or Master students in our group during the last years

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Giannis Tsiagkas

Research Interests – general description

Probabilistic and statistical aspects in genome organization – Non-randomness at several length scales.

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Deviations from randomness at the level of nucleotide n -tuplets. Patterns related to the functionality of genomic regions and to global genome structure.

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Deviations from randomness at the “middle” length scale, expressed mainly through clustering of similar nucleotides. Distinction between protein-coding and non-coding functionalities.

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Long range correlations and Zipf laws in the genome structure. Power-laws in the distribution of exons and of other genomic functional localizations. Entropic scaling in the study of genomic sequences as an indication of long range order and fractality.

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DNA sequences seen as genomic text – Linguistic features in the genome: redundancy, multiple coding, polarity and asymmetry etc.

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“Conservation laws” at the genome structure. The case of “Chargaff’s 2nd parity rule”. The use

of deviations from this law in the study of genomic dynamics and evolution.

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Evolution at the genomic level. Formulation of minimal evolutionary scenarios compatible with the observed probabilistic features of genomes. Interpretation of the above mentioned probabilistic features either by selectionist or mutationist causality.

Pattern formation in biological systems – Self-organization and evolution.

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Early development – Left-right asymmetries – Mechanisms of activation of Hox genes during limb development.

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Reaction-diffusion systems – Spontaneous symmetry breaking and pattern-formation in systems with feedbacks and non-linear dynamics

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Prebiotic and early evolution as a complex self-organization procedure.

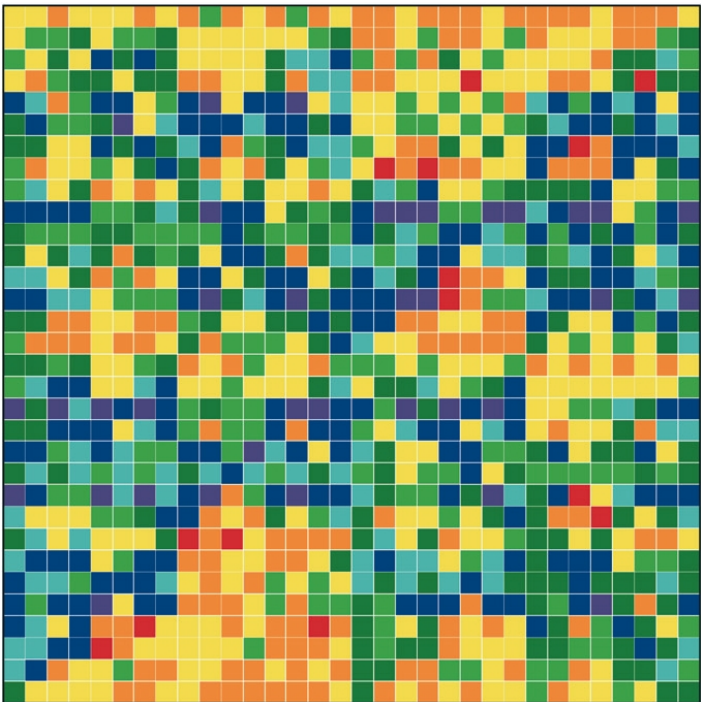
Findings in the field of Computational Genomics

“Word” preference in the genomic text and genome evolution

C
(b)

(a)
G

G C



T T A (dG)

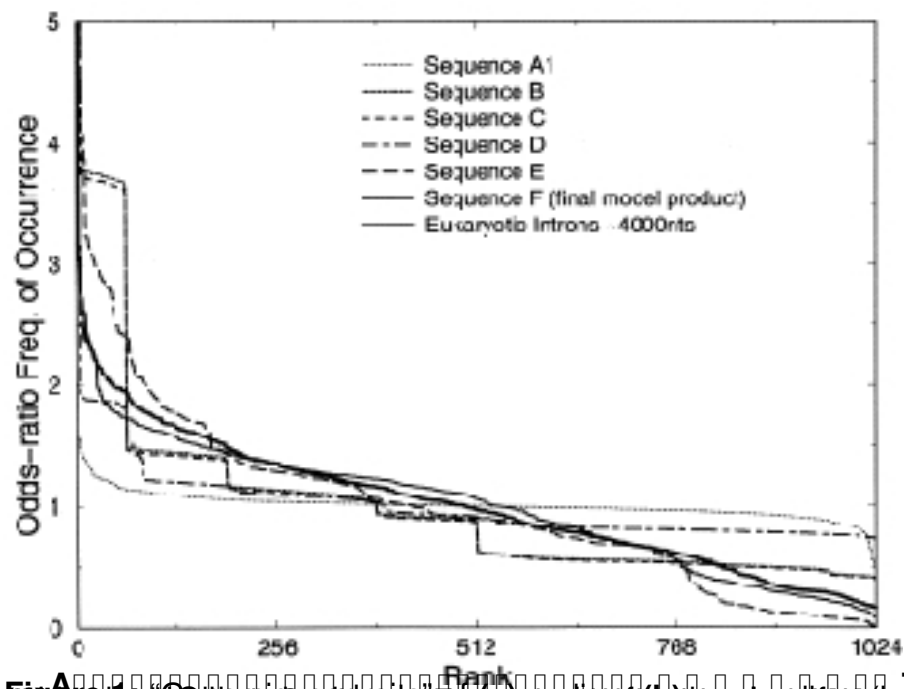
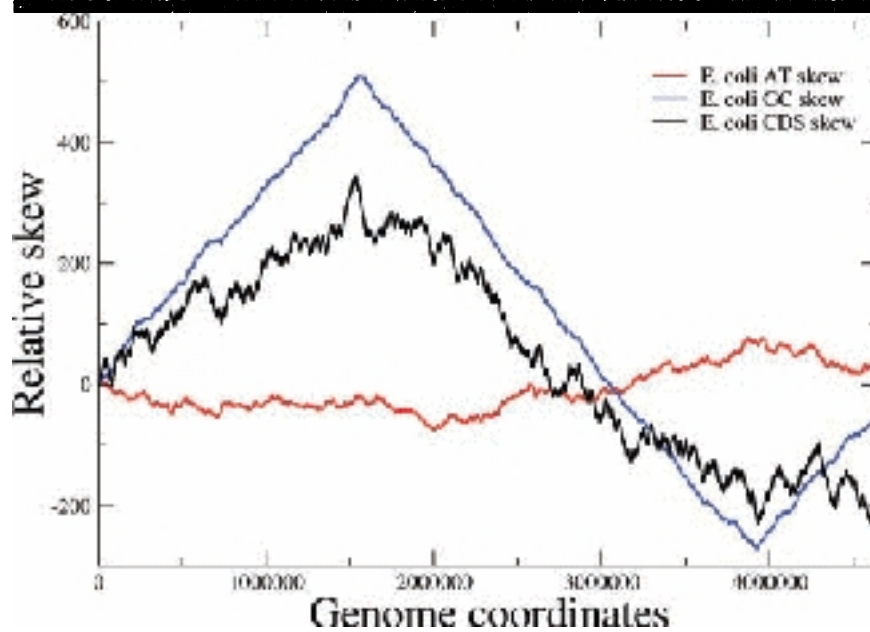


Figure 1: "Genomic patterns" of *E. coli* genome. The graph shows the relative skew of the *E. coli* genome, calculated as the difference between the number of A's and T's (or G's and C's) in a given region, normalized by the total number of bases. The x-axis represents the genome coordinates (0 to 40,000,000), and the y-axis represents the relative skew (from -200 to 600). The three lines represent the AT skew (red), GC skew (blue), and CDS skew (black) of the *E. coli* genome.



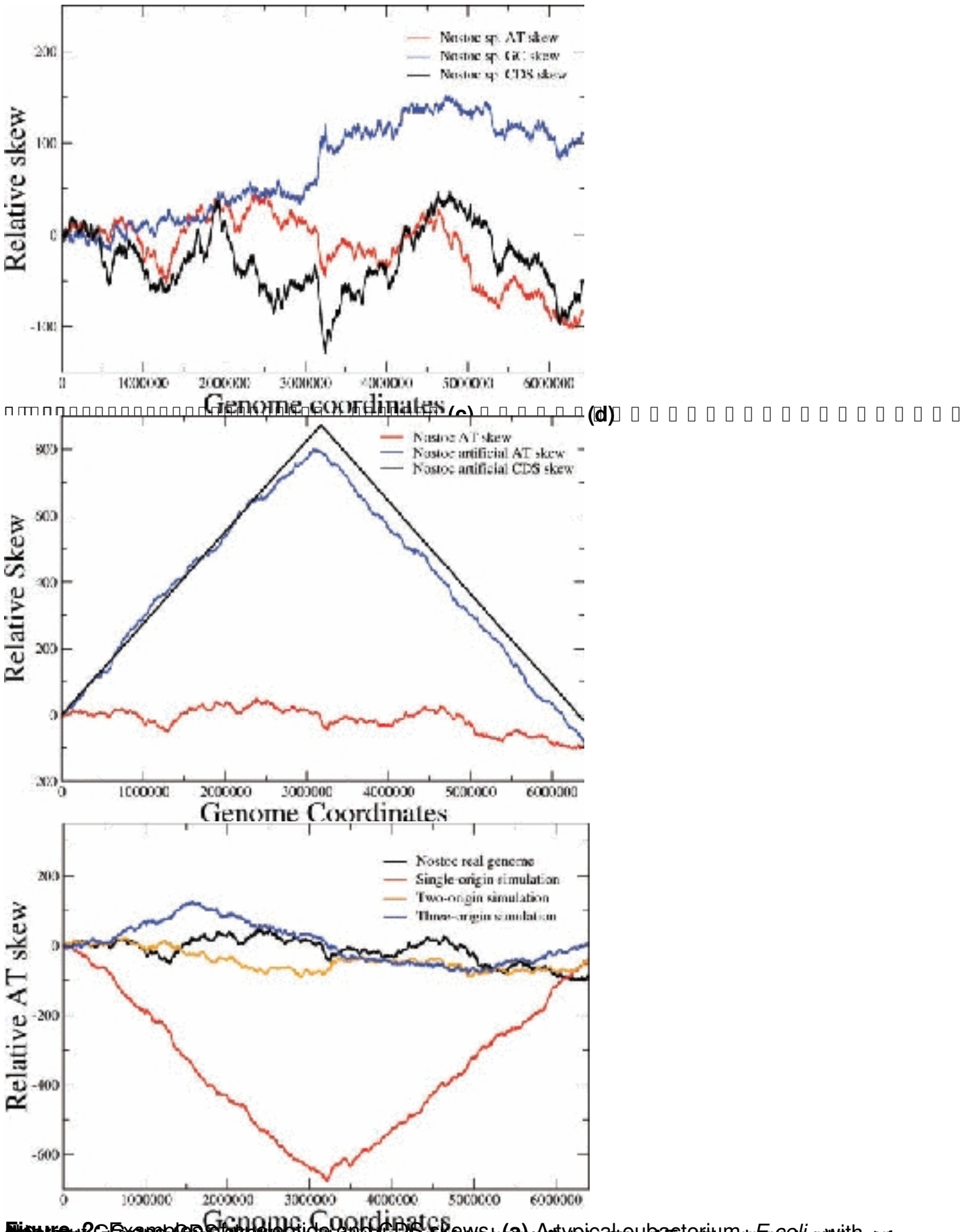
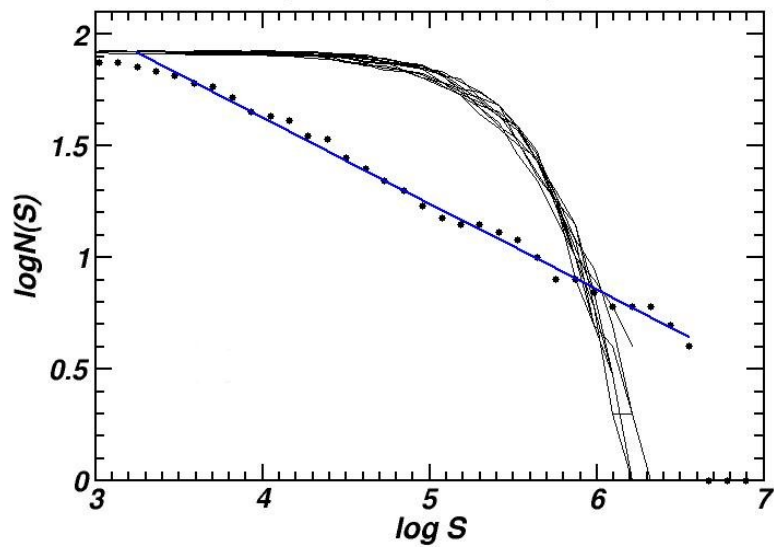
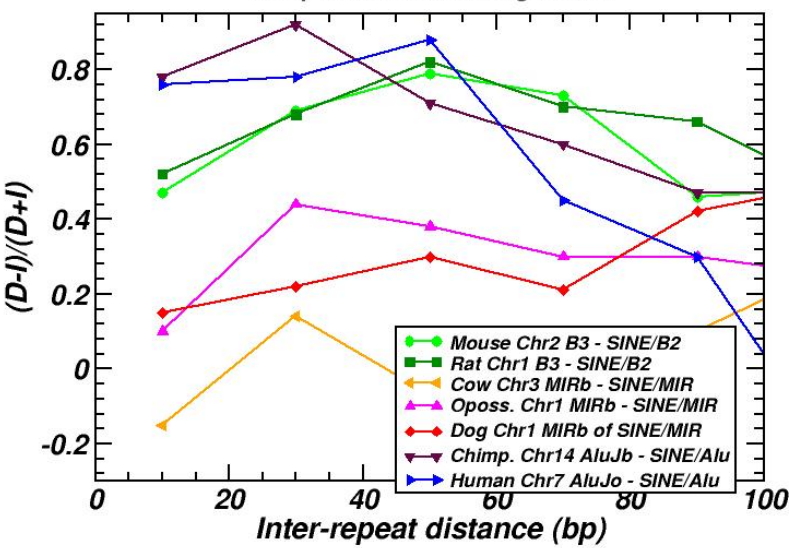


Figure 2. Example of nucleotide and CDS skews. (a) Atypical eubacterium, *E. coli*, with artificial AT skew. (b) *E. coli* with artificial GC skew. (c) *E. coli* with artificial CDS skew. (d) *E. coli* with artificial AT skew. (e) *E. coli* with artificial CDS skew.

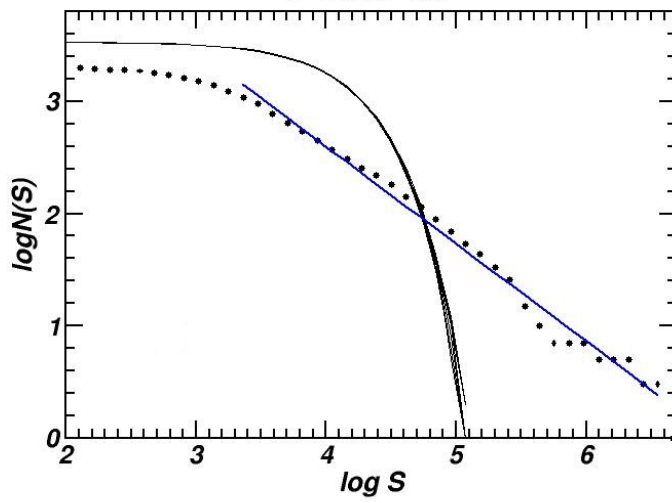
D. melanogaster chr. 3R, DNAREP1_DM - Helitron
Length min. = 200, E=3.31, $\mu=0.38$



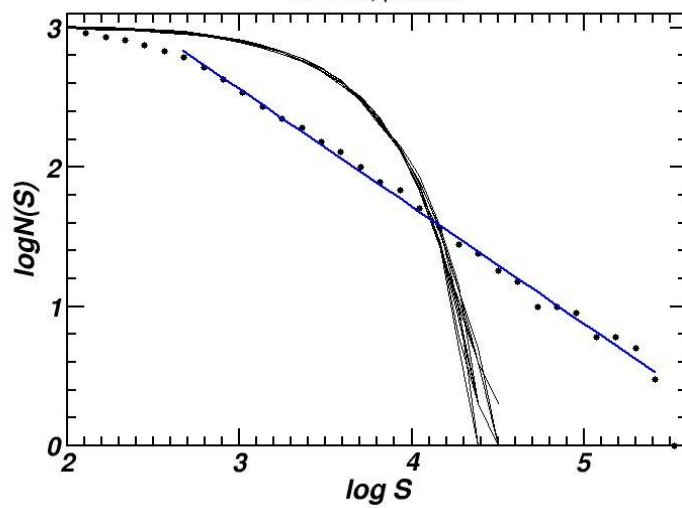
Inverted vs. Direct SINE pairs
Comparison between organisms



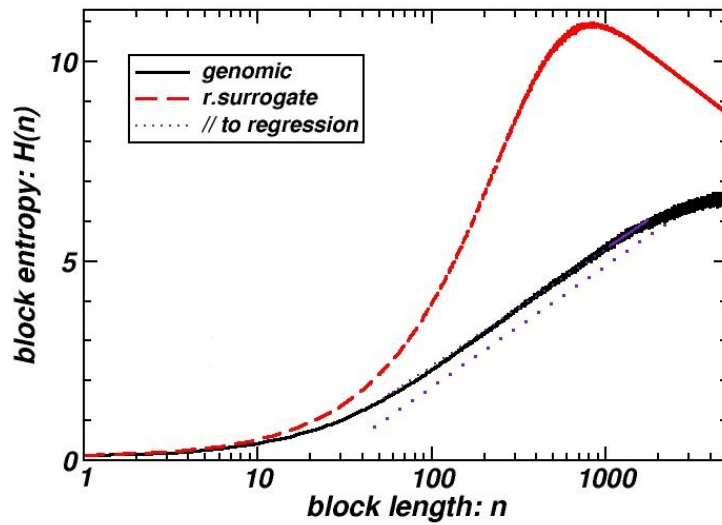
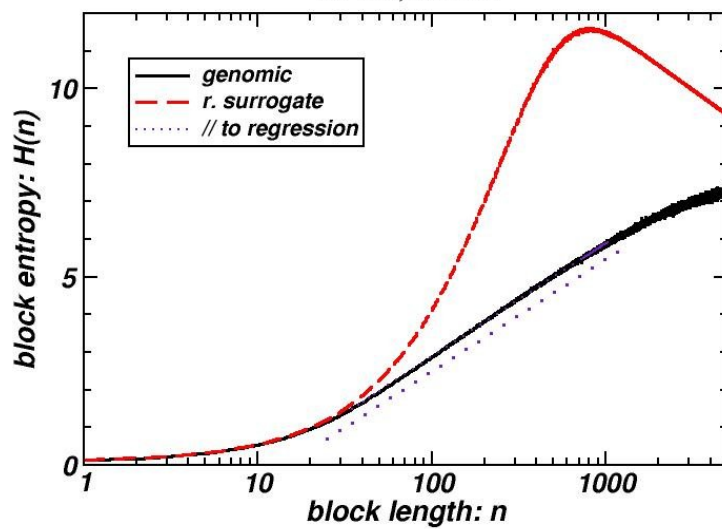
H. sapiens chr. 21, Protein Coding Segments
 $E = 3.17, \mu = 0.87$



G. gallus chr. 22, Protein Coding Segments
 $E = 2.74, \mu = 0.84$

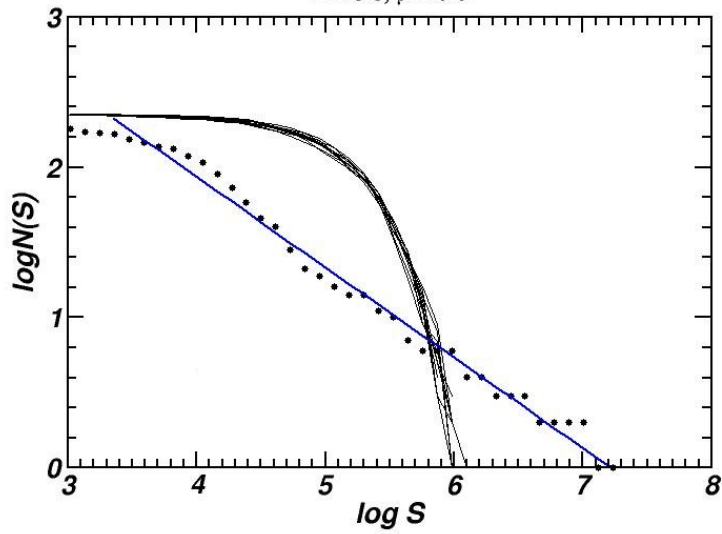


(a) (b)

H. sapiens chr. 20, Protein Coding Segments $E = 2.03, R = 2.38$ ***G. gallus chr. 4, Protein Coding Segments*** $E = 1.94, R = 2.12$ 

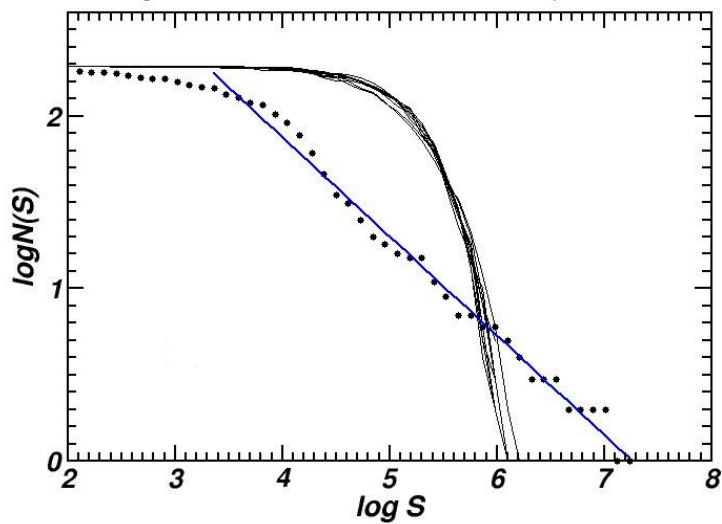
H. sapiens chr.19, CNEs (type: Amniotic)

$E = 3.8, \mu = 0.6$



H. sapiens chr.19, CNEs (type: Amniotic)

genes and flanks are masked; $E = 3.9, \mu = 0.57$



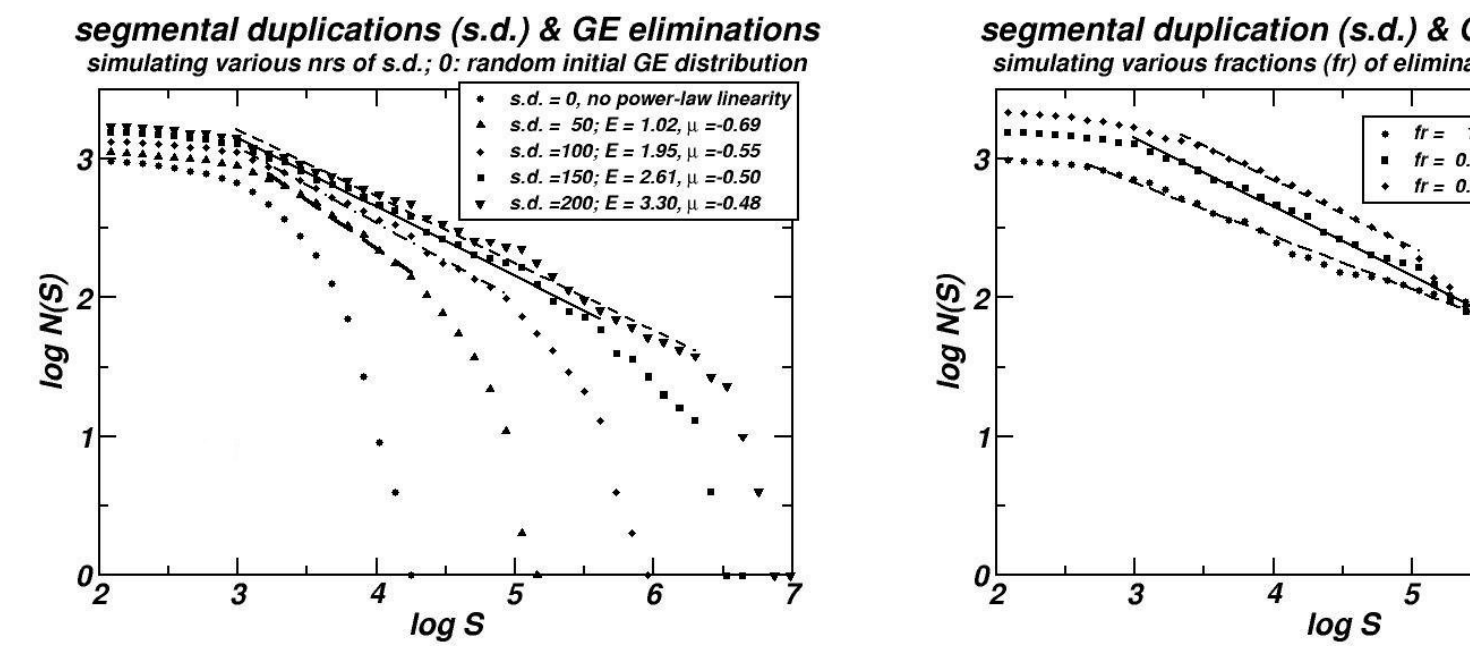


Figure 6: (a) $N(S)$ vs S for varying $s.d.$ values (0: random initial GE distribution, 50, 100, 150, 200) and (b) $N(S)$ vs S for varying fr values (0.0, 0.5, 0.9).