

Inhomogeneous Sequences of Letters in DNA, Proteins and Languages

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Abstract

1D, 2D and 3D ordering of different species like the bases A, C, G and T of DNA sequences is characterized by the self-coordination numbers T_i of $i = 1, 2, 3$ first, second, etc. neighbors and concentrations x and y . The T_i or α_i values ($\alpha_i T_i^{\max} = T_i - (T_i^{\max} - T_i) x/y$) of each species in a T_1, T_2, T_3 or α_1, α_2 structure map can be related with attractive or repulsive interactions of the species. Similar structure maps are obtained for a similar evolution process as outlined for different languages which were influenced by the Latin language. The DNA and protein sequences of enolase, secA or cytochrome enzymes can be ordered in different groups, which are supposed to be related by evolution.

Keywords: DNA sequences, languages, patterns, evolution

Abbreviations and notations:

T_i : self-coordination number of $i = 1, 2, 3, \dots$ first, second, third, etc. neighbors,

α_i : short-range order parameter.

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1 INTRODUCTION

The ordering of different species A, B, C, etc. can be characterized by the self-coordination numbers T_i with $i = 1, 2, 3$ of nearest, second and third neighbors and the ratio $r = y/x$ of concentrations x of the analyzed species and y of all other species. The one-dimensional row of A and B with maximum coordination numbers $T_1^{\max} = T_2^{\max} = T_3^{\max} = 2$ at distances $a, 2a$ or $3a$ is the simplest example for ordering at composition $A_x B_y$ with attractive or repulsive A–A interactions. The relation between structures and interactions can be outlined for A = ladies and B = gentlemen in a conference hall. Equal numbers of ladies and gentlemen ($r = y/x = 1$) can form three homogeneous structures with sequences AB, AB, etc. (opera house ordering), AABB, AABB, etc. (gay party) and $A_\infty B_\infty$ (moslem school). Each lady A (and each B) of the three structures has the self-coordination numbers 0 2 0 (opera), 1 0 1 (gay party) or 2 2 2 (moslem school) of other ladies. The complete segregation of ladies and gentlemen in a moslem school is achieved for infinite domains of ladies and gentlemen, where other T_i values at domain boundaries can be neglected. The T_i values or α_i values can be plotted in T_1, T_2, T_3 or α_1, α_2 structure maps with the three homogeneous structures at the corners of a triangle (first and second plot of Fig. 1).

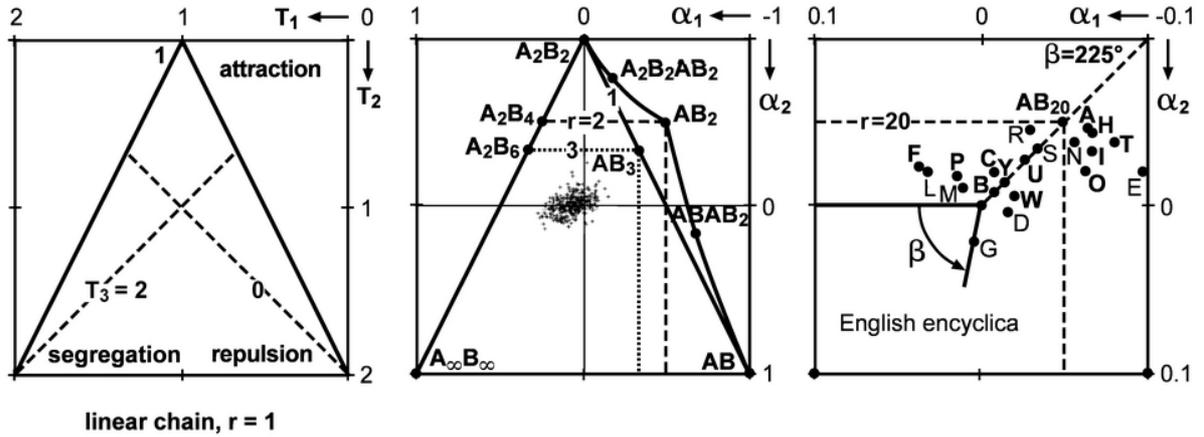


Figure 1. Structure map of linear chain with self-coordination numbers T_i of $i = 1, 2$ and 3 neighbors (first plot) or short-range order parameters α_1, α_2 as axes with α_1, α_2 values of 118 DNA sequences (+) and different periodic sequences of A and B (second plot). A section of this structure map with the α_1, α_2 values of each letter of the English translation of the encyclica *fides et ratio* is shown in the third plot. The letters V, K, Q, J, X, Z close to B position at $\beta = 225^\circ$ are not shown.

The α_i values obtained from T_i values by $\alpha_i T_i^{\max} = T_i - (T_i^{\max} - T_i)x/y$ are more appropriate for evaluation of an increased number of species like A = mother, B = father, C = daughter and D = son of families or A = Adenine, C = Cytosine, G = Guanine and T = Thymine of DNA sequences (second plot of Fig. 1). The three homogeneous structures at the corners of a triangle can be related with repulsive A–A interactions (opera house), attractive A–A interactions (gay party) and segregation (moslem school). The different corners of the α_1, α_2 structure map were obtained numerically for increasing numbers $n = x + y$ of seats with periodic boundary conditions. This can be compared with A and B sitting on a round table with n seats. The homogeneous AB, AB or AABB, AABB structures are obtained for $n = 2$ or $n = 4$ seats. The number of structures increases to 2^{n-1} as the round table is increased to $n = x + y$ seats and the first site is occupied by A. A maximum of $n = 24$ seats were considered at $y/x = 1$ with $\binom{23}{11} = 1\,352\,078$ structures. Few structures of the structure map like the $A_\infty B_\infty$ (moslem school) or $(A_2 B)_\infty (AB_2)_\infty$ (triplet cluster) with T_i values 2 2 2 or 2/3 2/3 2 are derived by extrapolation. Tables with infinite size or separate tables for A (or $A_2 B$) and B (or AB_2) are necessary, if all seats are occupied. Structures at the edges of the triangle are obtained by combination of corner structures like

ABA₂B₂, ABA₂B₂ (inhomogeneous boundary structures). This structure is inhomogeneous with different T_i values of the three A positions 0 1 2, 1 0 2 and 1 1 0, which are averaged to 2/3 2/3 4/3. Borders of the α_1, α_2 structure map connecting two structures with different $r = y/x$ values, like the border between the A₂B₂ and AB₂ (or ABC) structure, are curved. At increased r values the α_1, α_2 structure map is reduced by the boundary at $\alpha_1 = \alpha_2 = 1/r$ ($r \geq 2$). A random distribution is obtained for $\alpha_i = 0$ (all r values). The entropy is increased for all structures, which are not on the border of the structure map (inhomogeneous metastable structures) to a maximum at $\alpha_i = 0$. Combinations of clusters A_{*n*}B_{*n*} at attractive interactions and isolated groups AB at repulsive interactions are inhomogeneous border structures for infinite n and inhomogeneous metastable structures for finite n like A₅B₅AB with T_i values 4/3 4/3 1. The situation is similar to mobbing of a single A who is separated by empty B seats from the A₅ cluster of people in a conference. These inhomogeneous structures, which are not on the border of the structure map, are supposed to be metastable. They should vary to stable border structures by an evolution process. Artificial metastable structures, which are obtained by computer-assisted design, are varied after some time. Attractive or repulsive A–A forces or potentials can be calculated from homogeneous structures AB, AABB, AB₂ or A_∞B_∞ with maximum interactions, for inhomogeneous border structures and inhomogeneous metastable structures in the 1D row and the same method applied to 2D or 3D structures like the ordering of leaves (phyllotaxis), sunflower seeds, virus capsomeres or the coat patterns of zebras and giraffes [1,2]. Structures with maximum interactions can also be found by other forces as is outlined by the simple example of girls A and boys B in school. The attractive A–B interaction (or repulsive A–A interaction) is essential for the existence of the human being. Weak attractive or repulsive interactions could occur between the bases of DNA sequences or the amino acids of proteins containing atoms with different charges. The analyses of human languages and carols are probably somewhat easier to understand than the voices of animals like whales or birds. The evolution process of the Latin language could be studied in more detail, if old Italian, English or German texts would be analyzed by the present method.

2 RESULTS

Usually different T_i or α_i values are obtained for the different letters. The dots close to $\alpha_1 = \alpha_2 = 0$ in Fig. 1 (second plot) correspond to the α_i values of the four bases A (Adenine), T (Thymine), C (Cytosine) and G (Guanine) in 118 enzym DNA sequences of different species [3]. These values deviate from a random distribution. Some values are close to the borders at increased r values (not shown) of the α_1, α_2 structure map. The positions on the structure map are assumed to vary with time as the ordering process of ladies and gentlemen can vary with time. The T_i values of school children are usually varied from a point close to 2 2 2 ; (1) (moslem school of children at an early age) to a point 0 2 0 ; 1 (opera house) for teenagers. The DNA sequences can be ordered in different groups (Table 1). The ordering of the bases can be compared to the ordering of school children at an intermediate age. Most bases of the enzym enolase have a tendency to clustering (moslem school). A tendency to repulsion (opera house) is preferred by the bases of the chicken (*Gallus gallus*) and short human or rat DNA sections. The groups of the enzym secA are somewhat different with repulsive base interactions for the species *Halobacterium sp.* The evolution process of most species of one group is supposed to be similar. In that case the present groups can be compared with results obtained by other methods [4,5]. An increased number of clusters are suggested for the early evolution process [6]. Another example of evolution are the 26 letters of the Latin language compared to different other languages which were influenced by the Latin. Fig. 1 (third part) shows a section of the α_1, α_2 structure map for the English translation of the encyclica *fides et ratio* (vatican.de). Many letters are located on or close to the line $\alpha_1 = \alpha_2$ (or $T_1 = T_2$). These conditions are obtained for isolated A's. The $T_1 = T_2 = 0$ values correspond to negative α_i values. The letters S U Y V B etc. of the English language (values in brackets, Table 1) are usually in isolated positions. (The word 'mobbing' containing bb does not occur in the analyzed section of the encyclica with about 4000 letters.) These values at $\alpha_1 = \alpha_2$ are on a line with $\beta = 225^0$ of 2D polar coordinates (Fig. 1). The letters at increased β values R, C, P, M, L, F are also occurring as miniclusters like C

Table 1

Characterization of sequences of letters A, B, C, ... in languages, proteins, carols or nucleic acids for example (GenBank of the National Center for Biotechnological Information) by the location of Adenine (A), Thymine (T), Cytosine (C) and Guanine (G) bases on the α_1 , α_2 structure map (Fig. 1) in 2D polar coordinates $\alpha = \sqrt{\alpha_1^2 + \alpha_2^2}$ and angle β ($\beta = 0^0$ or 90^0 for the values on the positive α_1 and α_2 axis). The letters written in bold are on the border of the structure map (inhomogeneous border structures). The F, G, etc. in carols can be FIS, GIS, etc. in some cases.

Species	Repulsive [45 ⁰] 45 ⁰ < β < 225 ⁰	single (225 ⁰)	attractive 225 ⁰ < β
Languages			
Latin encyclica	VOIESTARNPU	(□QHGBXKWXYZJ)	MCDLF
Spanish encyclica	VDOIEAST□MNU	(CPHBFQGYZXJKW)	RL
Italian encyclica	VOIEA□UN	(RHQJKWXY)	DCMTSPLFGZB
Portuguese encyclica	DMVOAEI□TNSC	(UPLQFHGBZXJKWY)	R
French encyclica	MGOEI□AUT	(QHVJBXYKWZ)	RDSCNPLF
German encyclica	GHTOEDAIR	(□UCWBZKVJQXY)	NSLFPM
English encyclica	GDEOWTIHNA□	(SUYVBKQJXZ)	RCPMLF
Proteins			
Rice Enolase	A	(LNQMFPSEHVTYC)	IGK
<i>Drosophila</i> Enolase	[E]LCTMADPI	(NSGFQYRW)	KVH
<i>Ricinus</i> Enolase	[L]WNRPA	(VTYQIKFC)	GSEDH
Yeast Enolase	[ET]AKNPISV	(RQYLMWC)	GFHD
<i>E.coli</i> Enolase	[A]TPDSLK	(FRYQWC)	GVNIEMH
<i>Staphylococcus</i> Enolase	ATSYPENDVK	(FRQHC)	GLIM
<i>Actinobacillus</i> SecA	[HPI]VCGFATK	(W)	LNEDSYQMR
<i>Borrelia</i> SecA	VCNFGAKT	(QMPYHW)	ELSDIR
<i>Phormidium l.</i> SecA	GFDTAEN	(QSPMKHWC)	RLYIV
<i>Streptomyces l.</i> SecA	[FM]AQHVPTGL	(SYWC)	IEDKNR
<i>Mycobact. smeg.</i> SecA	[PQ]AGFTVSE	(YKHC)	LNMIDR
Human Cytochrome	[HRN]KVSDAG	(PQMYWEC)	FILT
<i>Camponotus f.</i> Cytochr.	DFLI	(AMYGNVEPR)	THKWS
Silk GAGAG(SG(AG) _n) ₈	GA	(SY)	
SGAAGY			

Species	Repulsive [45 ⁰] 45 ⁰ < β < 225 ⁰	single (225 ⁰)	attractive 225 ⁰ < β
music notes			
O come, all ye faithful (Latin)	CADG	(HF)	E
Silent Night (F. Gruber)	[GH]F	(DE)	AC
Joy to the World (G.F. Handel)	FG	()	HACDE
In Dulci Jubilo (German)	GC	(H)	ADFE
Hark, the Herald Angels Sing (F. Mendelssohn)	C	(A)	HFDGE
O Christmas Tree (German)		(AFE)	DCGH
Jingle Bells (J. Pierpont)		(GA)	FDEHC

Table 1 continued

Species	rep.	sgl	att.	Species	rep.	sgl	att.
enolase				secA			
<i>Drosophila melanogaster</i>		()	TGCA	<i>Actinobacillus actinomycet</i>		()	CGTA
<i>Loligo pealii</i>		()	TGCA	<i>Synechocystis sp.</i>		()	CGTA
<i>Sceloporus undulatus</i>		()	TGCA	<i>Cyanidium caldarium</i>		()	CGTA
<i>Eummeceus inexpectatus</i>		()	TGCA	<i>Phormidium laminosum</i>		()	CGTA
<i>Oryza sativa</i>		()	TGCA	<i>Rickettsia prowazekii</i>		()	CGTA
<i>Pneumocystis carinii</i>		()	CGTA	<i>Bacillus subtilis long</i>		()	GCTA
<i>Saccharomyces cerevisiae</i>		()	CGTA	<i>Odontella sinensis</i>		()	GCTA
<i>Zymomonas mobilis</i>		()	CGTA	<i>Guillardia theta</i>		()	GCTA
<i>Entamoeba histolytica</i>		()	CGTA	<i>Staphylococcus carnosus</i>		()	CGAT
<i>Schistosoma japonicum</i>		()	CGTA	<i>Porphyra purpurea</i>		()	CGAT
<i>Python regius</i>		()	GCAT	<i>Mycobacterium tuberculosis</i>		(C)	GAT
<i>Arabidopsis thaliana</i>		()	GCAT	<i>Heterosigma akashiwo</i>		()	GCAT
<i>Lycopersicon esculentum</i>		()	GCAT	<i>Mycoplasma pneumoniae</i>		()	TGAC
<i>Zea mays</i>		()	GCAT	<i>Borrelia burgdorferi</i>		()	CTAG
<i>Streptococcus therm.</i>		()	CGAT	<i>Aquifex aeolicus</i>		()	GTAC
<i>Nitrosomonas europea</i>		()	CGAT	<i>Mycoplasma genitalium</i>		()	ATCG
<i>Lycopersicon esculentum</i>		()	CGAT	<i>Prochlorothrix hollandica</i>		()	ACGT
<i>Ricinus communis</i>		()	GTCA	<i>Chlamydia pneumoniae</i>		()	GATC
<i>Chlamydomonas reinhardtii</i>		()	GTCA	<i>Streptomyces galbus</i>	A	()	CTG
<i>Candida albicans</i>		()	GCTA	<i>Prochloron didemni</i>	T	()	GCA
<i>Schizosaccharomyces pombe</i>		()	GCTA	<i>Mycoplasma capricolum</i>	C	()	GAT
<i>Rice m RNA</i>		()	CTGA	<i>Mycobacterium leprae</i>	C	()	GAT
<i>Neocallimastix frontalis</i>		()	TGAC	<i>Anabaena variabilis</i>	C	()	GAT
<i>Cunninghamella elegans</i>		()	TCGA	<i>Escherichia coli</i>	C	()	GTA
<i>Mesembryanth. crystall.</i>		()	GACT	<i>Haemophilus influenzae</i>	C	()	ATG
<i>Plasmodium falciparum</i>		()	GTAC	<i>Synechococcus sp.</i>	G	()	CTA
<i>Gluconobacter oxydans</i>	T	()	GCA	<i>Anacystis nidulans</i>	G	()	CTA
<i>Pelusios subniger</i>	T	()	GCA	<i>Bacillus firmus</i>	G	()	CTA
<i>Caiman crocodilus</i>	T	()	GCA	<i>Listeria monocytogenes</i>	G	()	CAT
<i>Alligator mississippiensis</i>	T	()	GCA	<i>Pisum sativum</i>	G	()	CAT
<i>Rattus norvegicus</i>	T	()	GCA	<i>Phormidium laminosum</i>	G	(C)	AT
<i>Peking duck</i>	T	()	GCA	<i>Helicobacter pylori</i>	G	()	TAC
<i>Xenopus laevis</i>	T	()	GCA	<i>Spinacia oleracea</i>	[C]G	()	AT
<i>Trachemys scripta</i>	T	(G)	CA	<i>Borrelia burgdorferi</i>	CG	()	AT
<i>Escherichia coli</i>	T	()	CGA	<i>Treponema pallidum</i>	CG	()	AT
<i>Mus musculus long</i>	T	()	CGA	<i>Staphylococcus aureus</i>	CG	()	AT
<i>Bos taurus</i>	T	()	CGA	<i>Arabidopsis thaliana</i>	CG	()	AT
<i>Mus musculus short</i>	T	()	AGC	<i>Bacillus subtilis short</i>	CG	()	TA
<i>Aspergillus oryzae</i>	T	()	AGC	<i>Zea mays</i>	CG	()	TA
<i>Homo sapiens long</i>	T	()	CAG	<i>Streptomyces coelicolor</i>	GT	()	CA
<i>Homo sapiens alpha short</i>	T	(C)	AG	<i>Streptomyces lividans</i>	GT	()	CA
<i>Cladosporium herbarum</i>	T	()	GAC	<i>Streptomyces griseus</i>	GT	()	CA
<i>Staphylococcus aureus</i>	C	()	GAT	<i>Stylosanthes scabra</i>	AT	()	CG
<i>Alnus glutinosa</i>	C	()	GAT	<i>Antithamnion spec</i>	TA	()	CG
<i>Streptococcus intermedius</i>	C	()	GTA	<i>Pavlova luthergerii</i>	A	(G)	CT
<i>Aspergillus oryzae</i>	TC	()	GA	<i>Caulobacter crescentus</i>	TG	()	CA
<i>Mastigamoeba balamuthi</i>	TC	()	GA	<i>Mycobacterium bovis</i>	GC	()	TA
<i>Fasciola hepatica</i>	TG	()	CA	<i>Vibrio alginolyticus</i>	GC	()	TA
<i>Sphenodon punctatus</i>	TG	()	CA	<i>Deinococcus radiodurans</i>	GC	()	AT
<i>Bacillus subtilis</i>	GC	()	TA	<i>Chlamydia trachomatis</i>	AGC	()	T
<i>Campylobacter fetus</i>	CG	()	TA	<i>Thermotoga maritima</i>	ACG	()	T
<i>Human m RNA very short</i>	TCA	()	G	<i>Mycobacterium smeg.</i>	GCT	()	A
<i>Rat neuron-specific short</i>	CAT	()	G	<i>Rhodobacter capsulatus</i>	TGC	()	A
<i>Gallus gallus</i>	ATCG	()		<i>Halobacterium sp.</i>	CGAT	()	

Species	rep.	sgl	att.	Species	rep.	sgl	att.
cytochrome				beta-globin [12]			
<i>Saccharomyces cerevisiae</i>		()	GCTA	<i>goat</i>		()	TGCA
<i>Bison bonasus</i>	A	()	CTG	<i>bovine</i>		()	GTAC
<i>Rat cytochrome</i>	A	()	CGT	<i>opossum</i>		(A)	TCG
<i>Chimera cytochrome</i>	A	()	TCG	<i>chimpanzee</i>		(T)	GAC
<i>Rhodobacter sphaeroides</i>	T	()	CGA	<i>human</i>		(GT)	AC
<i>Streptomyces tendae</i>	TA	()	CG	<i>gorilla</i>		(GT)	AC
<i>Capreolus capreolus</i>	TA	()	CG	<i>rat</i>	T	()	GAC
<i>Camponotus atriceps</i>	CA	()	TG	<i>mouse</i>	T	()	GAC
<i>Camponotus floridanus</i>	CA	()	TG	<i>gallus</i>	T	()	GAC
<i>Camponotus pennsylv.</i>	CA	()	GT	<i>rabbit</i>	TG	()	AC
Human cytochrome	AT	()	CG	<i>lemur</i>	[A]	CTG	()

and R in ‘occurring’. The other letters at decreased β values are frequently occurring at alternating positions like I in ‘minicluster’. The letters G, D, E, N, S, R, M and L of the English language are either isolated, alternating or miniclusters (inhomogeneous metastable structures). The coexistence of isolated A’s and clusters of A’s (mobbing situation) with $T_1 = T_2$ at $\beta = 45^0$ are observed for some music notes or amino acids in proteins (values in squared brackets, Table 1). The sequence of four amino acids in silk to a structure close to an inhomogeneous border structure (last example of proteins in Table 1) can be related to the high strength. The deviation of Alanine (A) is essential for the elasticity of silk [7].

3 CONCLUSION

The relation between the type of ordering and attractive or repulsive interactions is outlined for the most simple example of the one-dimensional row. The homogeneous sequences $AB, A_2 B_2$ of A and B are related to repulsive or attractive interactions. The A positions of boundary structures like $ABA_2 B_2$ are inhomogeneous with two neighboring values $T_1 = 0$ or 1. Other inhomogeneous structures usually have more than two neighboring values like $T_1 = 0, 1$ and 2 for $A_5 B_5 AB$ (inhomogeneous metastable structures).

Most letters of different languages and proteins are at the border of the structure map (inhomogeneous border structures written in bold in Table 1). The four bases of enzym DNA A, C, G and T are inhomogeneous metastable structures, which are not on the border of the structure map.

The human languages and DNA sequences of enolase of different species can be ordered to different groups. The similar sequences of Italian, Spanish and Latin languages or English and German languages can be explained by the evolution process. The voices of different animals like whales or birds could be analyzed by the same method as the music notes or the letters. Many animals have a single (A_∞) or two different sounds (AB), few like the voice of the cock or pigeon have the homogeneous AABB or ABC sequences. Homogeneous structures and border structures can also be found in coat patterns in an extended sense on surfaces of spheres (virus capsomeres, blackberries), cylinders (phyllotaxis, cylindrical viruses [1,8]) or other curved surfaces of animals. The capsomeres of the icosahedral microvirus have $T_1 = T_2 = 5$ and $T_3 = 1$ neighbors [2]. The capsomeres of most other viruses like simian virus 40, herpes virus or adenovirus are forming inhomogeneous border structures with $T_1 = 5$ and 6 [9,10]. The same applies to blackberries or sunflower seeds. The maximum density of the virus capsomeres on the surface of a sphere is essential to protect the virus genome. The cactus spines of some

cacti of the mamillaria family have a similar structure with $T_1 = 5$ or 6 neighbors for protection. The density of spines is decreased in *opuntiae cacti* with $T_1 = 4$ or columnar cacti with $T_1 = 2$. The coats of giraffes [11] contain white or yellow stripes with $T_1 = 3$ or 4 connections ($T_1 = 1$ or 2 for zebras, inhomogeneous border structures).

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