New spectral numerical characterization of DNA sequences

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Abstract: We present new numerical characterization of DNA sequences that is based on the modified graphical representation proposed by Hamori. While Hamori embeds the sequence into Euclidean space, we use analogous embedding into the strong product of graphs, $K_4 \bowtie P_4$, with weighted edges. Based on this representation, a novel numerical characterization was proposed in [14] which is based on the products of ten eigenvalues from the start and the end of the descending ordered list of the eigenvalues of the $L/L$ matrices associated with DNA. In this paper we compare two further numerical characterizations of the same type emphasizing the robustness of the approach.

Key-Words: numerical characterization, graph representation, graph invariant, DNA sequence

1 Introduction

Deoxyribonucleic acid (DNA) is the chemical inside the nucleus of all cells that carries the genetic instructions for making living organisms. A DNA molecule consists of two strands that wrap around each other to resemble a twisted ladder. The sides are made of sugar and phosphate molecules. The "rungs" are made of nitrogen-containing chemicals called bases. Each strand is composed of one sugar molecule, one phosphate molecule, and a base. Four different bases are present in DNA - adenine (A), thymine (T), cytosine (C), and guanine (G). The particular order of the bases arranged along the sugar - phosphate backbone is called the DNA sequence; the sequence specifies the exact genetic instructions required to create a particular organism with its own unique traits. Each strand of the DNA molecule is held together at its base by a weak bond. The four bases pair in a set manner: Adenine (A) pairs with thymine (T), while cytosine (C) pairs with guanine (G).

Nowadays the automated DNA sequencing techniques have led to an explosive growth in the number and the length of DNAs sequences from different organisms. This has resulted in a large accumulation of data in the DNA databases, but has also called for the development of suitable techniques for rapid viewing and analysis of the data. Graphical representations of DNA sequences were initiated by Hamori [6] and later expanded by many others, see the review [24] and a number of more recent papers, for example [10], [11], [12], [15], [16] [17][18],[24],[27], [3] the list being by no means exhaustive.

The advantage of graphical representation of DNA sequences is that they allow visual inspection of data, helping in recognizing major differences among similar DNA sequences. These techniques provide useful insights into local and global characteristics and the occurrences, variations and repetition of the nucleotides along a sequence which are not as easily obtainable by other methods.

Two-dimensional plots are obviously useful for visual communication of the results of an analysis, but can also be useful to help checking for the presence of an effect by human eye rather by a computer program, and finally, they are used for identifying unsuspected structures in the data. Recently, it has been shown that some of the graphical representations lead to numerical characterizations of DNA sequences and quantitative measures of the degree of similarity/dissimilarity between the sequences [[15], [16], [17], [18], [24], [27]]. Similarly as topological indices used as molecular descriptors can dramatically improve the search for synthesis of compounds with a desired property [23], it is hoped...
that the numerical descriptors of DNA may be used
to predict some properties of the DNA sequences.
An important advantage of a characterization of
structures by invariants, as opposed to use of codes,
is the simplicity of the comparison of numerical
sequences based on invariants. The price paid is a
loss of information on some aspects of the structure
that accompanies any characterization based on
invariants. The loss of the information, however,
can in part be reduced by use of larger number of
descriptors (invariants) [19], [20].

By a graph we mean a set V(G) of vertices, together
with a set E(G) of edges. A graph is the complete
graph K_n if any two of its distinct vertices are
adjacent. A graph is called the path P_n if it is
isomorphic to a graph on n distinct vertices
v_1, v_2, ..., v_n and n-1 edges v_i, v_{i+1}, 1 \leq i < n.

As the four bases A, G, C, and T are regarded
independent, at least four dimensions are needed for
an embedding that is free of using some arbitrary
conventions. A number of graphical representations
first embeds the DNA sequence into an Euclidean
space of some dimension, using a projection to 2-D
plot, where for the projection again some more or
less arbitrary choice has to be made.

In this paper, we essentially use a more dimensional
presentation, but instead of working with Euclidean
coordinates we rather embed the sequence into a
graph, more precisely into a strong product of
coordinates we rather embed the sequence into a
Euclidean space of some dimension, using a projection to 2-D
plot, where for the projection again some more or
less arbitrary choice has to be made.

In figures here we use a particular drawing of the
graph, which in our opinion seems to give a good
impression of the sequence to the observer.
The one dimensional plot of K_4 is of course not
isometric (i.e. the edges in the plot have different
lengths) but we believe that the resulted drawing
may be a reasonable compromise between the
arbitrary projection(s) and a unique more
dimensional embedding which can, of course, easily
be found by an isometric embedding of the complete
graph K_4 into Euclidean space, for example by
mapping A, C, G, and T to the edges of a tetrahedron
in 3D or to the four unit vectors in 4D.
Furthermore, based on this graph representation we
propose a novel numerical characterization of the
DNA sequence.

In contrast to some other numerical character-
izations that are based on the graphical
representations [12], [18], [27], our representation
is free of arbitrary choices because it is based on the
graph and not on its drawing, i.e. embedding and
projection. The numerical characterization uses
eigenvalues of a matrix that is based on the graph
distances.

The numerical invariant is computed for the first
exon of the β-globin gene for the 10 different
species, a dataset shown in Table 1, that is used in
many recent studies [10], [11], [12], [15], [16],
[17], [18], [24], [27] and is taken from EMBL-EBI
database [29]. This dataset is one of the primary
tools for comparison of different graphical and
numerical characterizations and was first used by
Nandy [13] and later by other authors [10], [15],
[16], [18], [24]. The reason why Nandy decided to
use this gene lies in the fact that β-globin sequences
represent a conservative gene, that is, the
gene that changes little from one species to another.
The differences between the values of the invariant
are used as a measure of similarity/dissimilarity
among the species.

We do not attempt to extensively comment the
results because this is not an area of our expertise.
However we wish to note that our results are not
like those obtained by similar computations which
are based on eigenvalues of the graphical
representations [15], but are based on graphs,
therefore our approach is using less computational
effort.

For example in [15] one has to compute 12 different
permutations of the graphical representation before
the actual characterization, while our approach
computes only one.

2 Modified Hamori curve representation

We based our research on DNA sequence
representation introduced by Hamori [6]. In this
method, the information content of a DNA sequence
is mapped into a three-dimensional space function
(H curve). The positive x-direction is used to count
the number of bases in the sequence. At each point
of x on the corresponding yz plane the four corners
(NW, NE, SE and SW as four points on the
compass) are taken to represent the four bases A, C,
G and T. Basic rule for the construction of the
sequence map is to move one unit in the corresponding direction depending on which nucleotide (base) is being plotted and to draw a connected line of all such points plotted, one for each unit in the x-direction. Thus a sequence like ATGGTGCACCTGACT... will generate a spiral along the x-axis.

H-curve representation is sensitive to the directions chosen for four bases. For example representation with bases ACGT corresponding to four corners is different from AGCT, since the distance from base A to base G is different in this two cases.

We modified this approach by putting the corners of four bases on the complete graph $K_4$ and weighted all the edges in $K_4$ with 1. This way we avoided the drawback of the original representation. Edges in the x direction or along $P_n$ are weighted with 1 if the base in the coding sequence is the same as the previous one and with $\sqrt{2}$ otherwise.

Formally, a sequence of the length n in this paper is a path in the strong product of the graphs $K_4$ and $P_n$. The strong product $G_1 \boxtimes G_2$ of graphs $G_1$ and $G_2$ has as vertices the pairs $(g,h)$ where $g \in V(G_1)$ and $h \in V(G_2)$. Vertices $(g_1,h_1)$ and $(g_2,h_2)$ are adjacent if either $\{g_1,g_2\}$ is an edge of $G_1$ and $h_1 = h_2$ or if $g_1 = g_2$ and $\{h_1,h_2\}$ is an edge of $G_2$ or if $\{g_1,g_2\}$ is an edge of $G_1$ and $\{h_1,h_2\}$ is an edge of $G_2$. The strong product is one of the standard graph products [9]. For example, the strong product of two edges (complete graphs on 2 vertices, $K_2$) is the complete graph on four vertices, $K_4$). Another example is the product of two paths of length 2.

Below is depicted a product of two copies of a general graph together with the factors.

Here $K_4$ is a complete graph on vertices $A,C,G,T$ and $P_n$ is a path on the vertices 1,2,...,n. The edges of the product are weighted as follows:

\[
W((i,j)(k,\ell)) = \begin{cases} 
1 & \text{if } i = k \text{ or } j = \ell \\
\sqrt{2} & \text{if } i \neq k \text{ and } j \neq \ell
\end{cases}
\]

Figure 1 shows modified Hamori curve, where first few edges between the $K_4$'s have weights indicated with the numbers on gray background. The factor $K_4$ is drawn on a circle and projected to obtain a 2-D drawing. Any other possibly nicer drawing of the final graph can be used [1]. However, we find our way of drawing the graph and the path a reasonable compromise that can be used as a help for easier understanding of our concept. Note that all the edges within the vertical factor ($K_4$) and all the horizontal edges have weight 1 while all edges between $K_4$ factors that are not horizontal have weight $\sqrt{2}$.

The motivation for choosing $\sqrt{2}$ is the intuitive assumption that the two factors in the product are orthogonal, hence the corresponding edge is the diagonal of a unit square.
While Hamori embeds the sequence into Euclidean space, we use analogous embedding into the strong product of graphs, $K_n \otimes P_k$, with weighted edges. Based on this representation, a novel numerical characterization was proposed in [1] which is based on the products of ten eigenvalues from the start and the end of the descending ordered list of the eigenvalues of the $L/L$ matrices associated with DNA. Below we explain this and two further numerical characterizations of the same type.

3 Numerical characterization of DNA sequences

In order to numerically characterize a DNA sequence given by the 2-D graphical representation based on our approach one can associate with a corresponding zigzag curve a matrix and consider matrix invariants that are sensitive to the form of the curve. This approach was first outlined and used by Randić, Vračko, Lerš, and Plavšič [16]. One of the possible matrices they use is the $L/L$ matrix (the length/length matrix) whose elements are defined as the quotient of the distance between a pair of the vertices (dots) of the zigzag curve and the sum of distances between the same pair of vertices measured along the zigzag curve. Here we use analogous matrix based on the weighted graph representation of DNA, i.e. the entries of the $L/L$ matrix are the quotients between the graph distance and the weighted graph distance.

Using this weights we can construct the $L/L$ matrix as is shown in Table 1 where we used the first 6 bases of the first exon of $\beta$-globin gene of human.

<table>
<thead>
<tr>
<th>Species &amp; Coding sequence</th>
<th>Gorilla (93 bases)</th>
<th>Bovine (86 bases)</th>
<th>Chimpanzee (105 bases)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Human (92 bases)</td>
<td>ATGGTGCACTGACTCCTGAGGAGAAGTCT-GCCGTTACTGCCCCGTGGGGCAGGTAACGTGGAATGAGGGTTGGTGGAGGCCCCTGGGC-AG</td>
<td>ATGCTGACTGCTGAGGAGAAGCTGCCGCT-ACCCTCTTCTGCGGGCAAGGTGAAGTGAATGTAAGGTTGCTGAGGCCCCTGGGCAG</td>
<td>ATGGTGCACTGACTCCTGAGGAGAAGTCT-GCCGTTACTGCCCCGTGGGGCAGGTAACGTGGAATGAGGGTTGGTGGAGGCCCCTGGGCAGGTTGGTATCAAGG</td>
</tr>
<tr>
<td>Opossum (92 bases)</td>
<td>ATGGTGCACTGACTCTTCTGAGGAGAAGAC-TCATACCTACCACATCTGCTAAGGTGCA-GGTGACCAAGACTGTTGAGGCCCTTTGGC-AG</td>
<td>ATGGTGCACTGACTCTTCTGAGGAGAAGAC-TCATACCTACCACATCTGCTAAGGTGCA-GGTGACCAAGACTGTTGAGGCCCTTTGGC-AG</td>
<td>ATGGTGCACTGACTCTTCTGAGGAGAAGAC-TCATACCTACCACATCTGCTAAGGTGCA-GGTGACCAAGACTGTTGAGGCCCTTTGGC-AG</td>
</tr>
<tr>
<td>Gallus (92 bases)</td>
<td>ATGGTGCACTGACTGCTGAGGAGAAGAC-TCATACCTACCACATCTGCTAAGGTGCA-GGTGACCAAGACTGTTGAGGCCCTTTGGC-AG</td>
<td>ATGGTGCACTGACTGCTGAGGAGAAGAC-TCATACCTACCACATCTGCTAAGGTGCA-GGTGACCAAGACTGTTGAGGCCCTTTGGC-AG</td>
<td>ATGGTGCACTGACTGCTGAGGAGAAGAC-TCATACCTACCACATCTGCTAAGGTGCA-GGTGACCAAGACTGTTGAGGCCCTTTGGC-AG</td>
</tr>
<tr>
<td>Lemur (92 bases)</td>
<td>ATGGTGCACTGACTGAGGAGAAGAC-TCATACCTACCACATCTGCTAAGGTGCA-GGTGACCAAGACTGTTGAGGCCCTTTGGC-AG</td>
<td>ATGGTGCACTGACTGAGGAGAAGAC-TCATACCTACCACATCTGCTAAGGTGCA-GGTGACCAAGACTGTTGAGGCCCTTTGGC-AG</td>
<td>ATGGTGCACTGACTGAGGAGAAGAC-TCATACCTACCACATCTGCTAAGGTGCA-GGTGACCAAGACTGTTGAGGCCCTTTGGC-AG</td>
</tr>
<tr>
<td>Mouse (92 bases)</td>
<td>ATGGTGCACTGACTGAGGAGAAGAC-TCATACCTACCACATCTGCTAAGGTGCA-GGTGACCAAGACTGTTGAGGCCCTTTGGC-AG</td>
<td>ATGGTGCACTGACTGAGGAGAAGAC-TCATACCTACCACATCTGCTAAGGTGCA-GGTGACCAAGACTGTTGAGGCCCTTTGGC-AG</td>
<td>ATGGTGCACTGACTGAGGAGAAGAC-TCATACCTACCACATCTGCTAAGGTGCA-GGTGACCAAGACTGTTGAGGCCCTTTGGC-AG</td>
</tr>
<tr>
<td>Rabbit (90 bases)</td>
<td>ATGGTGCACTGACTGAGGAGAAGAC-TCATACCTACCACATCTGCTAAGGTGCA-GGTGACCAAGACTGTTGAGGCCCTTTGGC-AG</td>
<td>ATGGTGCACTGACTGAGGAGAAGAC-TCATACCTACCACATCTGCTAAGGTGCA-GGTGACCAAGACTGTTGAGGCCCTTTGGC-AG</td>
<td>ATGGTGCACTGACTGAGGAGAAGAC-TCATACCTACCACATCTGCTAAGGTGCA-GGTGACCAAGACTGTTGAGGCCCTTTGGC-AG</td>
</tr>
<tr>
<td>Rat (92 bases)</td>
<td>ATGGTGCACTGACTGAGGAGAAGAC-TCATACCTACCACATCTGCTAAGGTGCA-GGTGACCAAGACTGTTGAGGCCCTTTGGC-AG</td>
<td>ATGGTGCACTGACTGAGGAGAAGAC-TCATACCTACCACATCTGCTAAGGTGCA-GGTGACCAAGACTGTTGAGGCCCTTTGGC-AG</td>
<td>ATGGTGCACTGACTGAGGAGAAGAC-TCATACCTACCACATCTGCTAAGGTGCA-GGTGACCAAGACTGTTGAGGCCCTTTGGC-AG</td>
</tr>
</tbody>
</table>

Fig. 1 Modified Hamori curve

Table 1 The coding sequences of the first exon of $\beta$-globin gene of 10 different species
For example, the first three entries of the first row are \( \frac{1}{\sqrt{2}} = 0.707 \), \( \frac{2}{\sqrt{2} + \sqrt{2}} = 0.707 \), and \( \frac{3}{\sqrt{2} + \sqrt{2} + 1} = 0.783 \).

<table>
<thead>
<tr>
<th>Base</th>
<th>A</th>
<th>T</th>
<th>G</th>
<th>G</th>
<th>T</th>
<th>G</th>
</tr>
</thead>
<tbody>
<tr>
<td>A</td>
<td>0.707</td>
<td>0.707</td>
<td>0.783</td>
<td>0.762</td>
<td>0.751</td>
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</tr>
<tr>
<td>T</td>
<td>0.707</td>
<td>0.828</td>
<td>0.783</td>
<td>0.762</td>
<td></td>
<td></td>
</tr>
<tr>
<td>G</td>
<td>1.00</td>
<td>0.828</td>
<td>0.783</td>
<td></td>
<td></td>
<td></td>
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<tr>
<td>G</td>
<td>0.707</td>
<td>0.707</td>
<td>0.783</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>T</td>
<td>0.707</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>G</td>
<td>0</td>
<td>0.707</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Formally, we assign the matrix \( LL \) to the sequence \( x \) with

\[
LL_x(i, j) = \frac{j - i}{d((x_i, i), (x_j, j))},
\]

where \( d((x_i, i), (x_j, j)) \) is the distance in the weighted graph \( K_i \otimes P_n \). More precisely,

\[
d((x_i, i), (x_j, j)) = \sum_{k=0}^{j-1} W((x_k, k)(x_{k+1}, k+1)) \quad \text{for} \quad j > i.
\]

(For \( i=j \) we put \( d((x_i, i), (x_j, j)) = 0 \) and for \( j < i \) we define \( d((x_i, i), (x_j, j)) = d((x_j, j), (x_i, i)) \).

We will characterize the coding sequences of the first exon of \( \beta \)-globin gene of 10 species (including human), shown in the Table 1, by means of the leading eigenvalues, \( \lambda \), of the \( LL \) matrix. Eigenvalues of a matrix are one of the best known matrix invariants. If a matrix is symmetric, as is the case with all the matrices considered here, the eigenvalues are real. A set of eigenvalues can be viewed as a characterization of a structure, but as is well known such characterization is not unique. In other words, different graphs and different structures may have the same set of eigenvalues. Such graphs are known as isospectral and have received considerable attention in mathematics [14, 7] and chemistry [8], of which we only indicated some earlier contributions. While it was initially thought that the complete coincidence of all eigenvalues may be an exception rather than a rule, the subsequent research revealed that isospectral graphs are more a rule than exception. That, however, does not diminish their utility, although they would fail to discriminate structures in testing for isomorphism [19]. On other hand, if two structures are similar they are likely to have similar eigenvalues and consequently similar product of leading eigenvalues. In a recent study in which the DNA sequence was characterized by average distances between various nucleic acid bases was shown that is very sensitive already when a single nucleic base has been changed [22].

Our characterizations are based eigenvalues of the matrix \( LL \). In [14] the product of the 10 largest and 10 smallest eigenvalues was taken. Here we will compare this numerical characterization with the second which is the product of the five largest and five smallest eigenvalues. Species have different lengths of DNA sequence, shortest is DNA sequence of the bovine (86 bases) and longest of the Chimpanzee (105 bases).

It may be reasonable to consider ways to cancel out from comparison the influence of different lengths of sequences as much as possible. Therefore we also consider a normalized characterization, where we take the \( n \)-th root of the product of the eigenvalues.

### 3 Similarities/dissimilarities among the coding sequences of the first exon of \( \beta \)-globin gene of different species

We will illustrate a natural method for the characterization of the DNA sequences with the examination of the similarities/dissimilarities among the 10 coding sequences shown in Table 1. The analysis of similarity/dissimilarity is based on the assumption that two DNA sequences are similar if the corresponding difference between the value of the numerical characterization is small.

The values of the numerical characterizations are as follows:
Formally we can define similarity relations as:

\[ S_x(y) = | \Lambda_x(x) - \Lambda_y(y) |, \]

where \( x, y \) are sequences of the species.

In this way we obtain a matrix of mutual similarities among species. First we present a matrix with similarities (because of the size, we present them at the end of paper) and then we draw two graphs based on the values in the matrix. First graph represents nearest similarities relation and second graph represents widest dissimilarities relation. Similarities/dissimilarities matrix for \( \Lambda_n(x) \) is in Table 3 and corresponding graphs are on Fig 2 and 3. Similarities/dissimilarities matrix for \( \Lambda_s(x) \) is in Table 4 and corresponding graphs are on Fig 4 and 5 and finally, similarities/dissimilarities matrix for \( \Lambda_s(x) \) is in Table 5 and corresponding graphs are on Fig 6 and 7.

While of course not surprisingly the three similarity measures give different numerical values, the overall results are not very much different. In particular, the smallest differences are associated with the pairs (human, chimpanzee), (human, gorilla) and (gorilla, chimpanzee) which is in accordance with our intuitive expectations and, not surprisingly, also in accordance with other studies [[10], [16]]. On the other hand, the largest entries in the similarity/dissimilarity matrix appear in rows belonging to bovine and opossum.

We may conclude that all presented numerical characterizations have captured some important features of the DNA sequences considered.

### 4 Conclusion

Our objective in [14] was to arrive at a numerical characterization of DNA sequences. This may be accomplished in a relatively simple algebraic manner and as such makes the proposed approach very attractive for the characterization of DNA sequences having 1,000 or more bases.

In this follow-up report we add results on related numerical characterizations showing that the approach is robust, hence the somewhat arbitrary choice of 5 or 10 eigenvalues taken does not severely influence the results of the method. The preliminary results presented here support the intuition that some important structural information of the sequences is encoded in the spectrum, and in particular in the largest and smallest eigenvalues. We have provided a method that is computationally more efficient than some earlier approaches. Needles to say that the outlined approach may be suitable for characterization of local fragments of DNA, which is precisely how one may look on the truncated DNA fragment considered in this work. Conceptually and computationally the approach is simple and therefore can be very useful in the field of bioinformatics.

### References:


[29] http://www.ebi.ac.uk/
Table 3 Matrix of the $\Lambda_1(x)$ similarities

<table>
<thead>
<tr>
<th></th>
<th>Human</th>
<th>Chimpanzee</th>
<th>Gorilla</th>
<th>Opossum</th>
<th>Gallus</th>
<th>Lemur</th>
<th>Mouse</th>
<th>Rabbit</th>
<th>Rat</th>
<th>Bovine</th>
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Fig. 2 Largest $\Lambda_1(x)$ similarities

Fig. 3 Largest $\Lambda_1(x)$ dissimilarities
### Table 4 Matrix of the $\Lambda_2(x)$ similarities

<table>
<thead>
<tr>
<th></th>
<th>Human</th>
<th>Chimpanzee</th>
<th>Gorilla</th>
<th>Opossum</th>
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**Fig. 4** Largest $\Lambda_2(x)$ similarities

**Fig. 5** Largest $\Lambda_2(x)$ dissimilarities
### Table 5 Matrix of the $\Lambda_n(x)$ similarities

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<th>Chimpanzee</th>
<th>Gorilla</th>
<th>Opossum</th>
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**Fig. 6** Largest $\Lambda_n(x)$ similarities.

**Fig. 7** Largest $\Lambda_n(x)$ dissimilarities.