

Coronavirus 2: Analysis of Regularity of Complete Genome (SARS-CoV-2/Hu/DP/Kng/19-020 RNA)

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Abstract

This paper shows a technique for the detection of regularities in the complete genome of Coronavirus 2. For this, the concept of intervals in the RNA chain and their comparison within the known sequence is used. The Maximum Regularity Index was applied. The results show that areas of high regularity are identified that coincide with ORF10, Gene N, nucleocapsid phosphoprotein and others. The interval around **3500** and the interval around **11000**, respectively, appear with high values of regularity and this do not correspond to particularly documented elements (both within Gene: **Orf1ab**). It might be in the interest of giving importance to that region. The detection of regularities can enhance the creation of biochemical or genetic techniques to deactivate certain functions of Coronavirus 2 or control its effects.

Keywords

Coronavirus 2; SARS-CoV-2; Complete Genome; RNA; index of maximum regularity; Covid-19; NCBI

Introduction

Coronavirus 2 has a sequence of 29902 nt (April 10th, 2020) [1]. Its full lineage is: Viruses; Riboviria; Nidovirales; Coronavirales; Coronaviridae; Orthocoronavirinae; Betacoronavirus; Sarbecovirus; Severe acute respiratory syndrome-related coronavirus and its genbank acronym is SARS-CoV2.

A method has been applied in which chains of RNA of a certain length are taken and compared with chains of the same length within the complete genome. In previous work this technique has been applied to other microorganisms [2],[3],[4].

Methodology

The index of maximum regularity $i_{max,r}$ was defined in [5] and in other papers such as [4] has been widely described.

In summary, a string or interval l_c is defined. It considers the new sequences that are generated $l_c - 1$ when a shift is made, one nucleotide at a time. Each ordered pair is part

of the set $B \times B$, where $B = \{A, G, T, C\}$. This procedure is repeated considering the original sequence together with the one generated by reversing its order.

Denoted by $n_{A,A,d}, n_{A,G,d}, n_{A,C,d}, n_{A,T,d}, n_{G,A,d}, n_{G,A,d}, n_{G,G,d}, n_{G,C,d}, n_{G,T,d}, n_{C,A,d}, n_{C,G,d}, n_{C,C,d}, n_{C,T,d}, n_{T,A,d}, n_{T,C,d}, n_{T,G,d}, n_{T,T,d}$ the number of ordered pairs actually counted. The $i_{max,r}$ is defined as:

$$i_{max,r} = \frac{\max\{D_1, \dots, D_{l_c-1}, D_0^*, \dots, D_{l_c-1}^*\}}{\left(l_c - \frac{l_c}{16}\right)^2 + 15\left(\frac{l_c}{16}\right)^2}$$

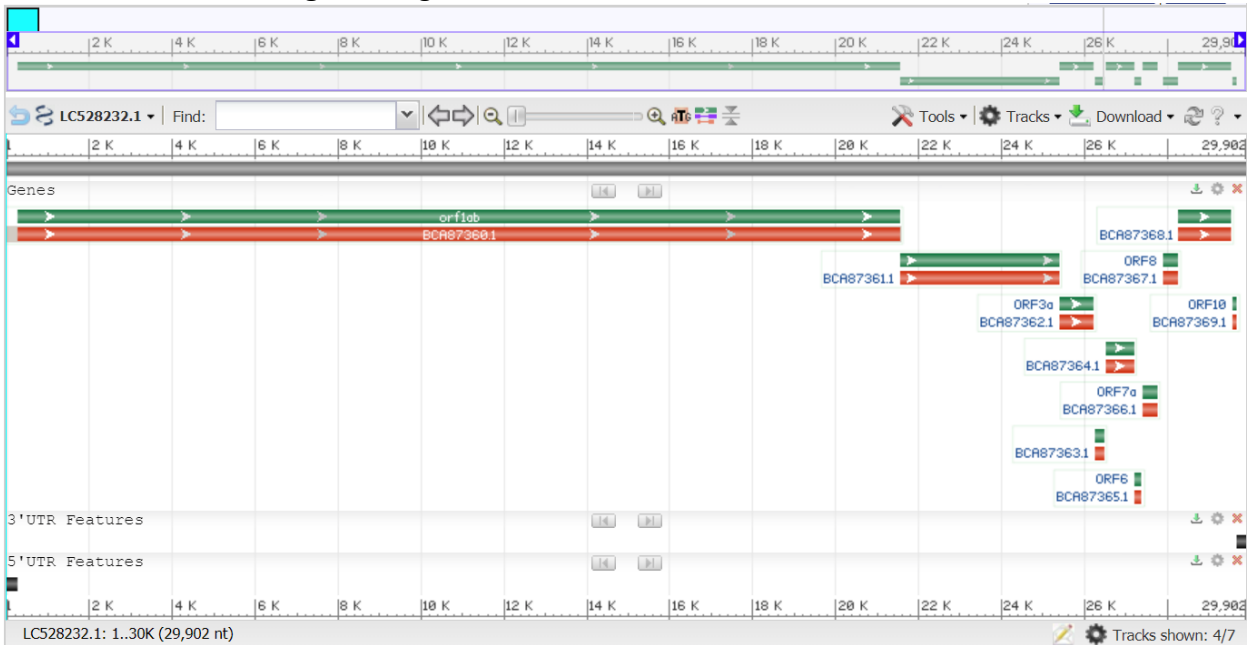
Where $D^*_d = \left(n_{A,A,d} - \frac{l_c}{16}\right)^2 + \left(n_{A,G,d} - \frac{l_c}{16}\right)^2 + \dots + \left(n_{T,T,d} - \frac{l_c}{16}\right)^2$

Results

The $i_{max,r}$ was applied to the complete genome of Coronavirus 2. Below are the results of coincidence with the genes already discovered by other techniques (NCBI, 2020) and possible areas of study for possible deepening.

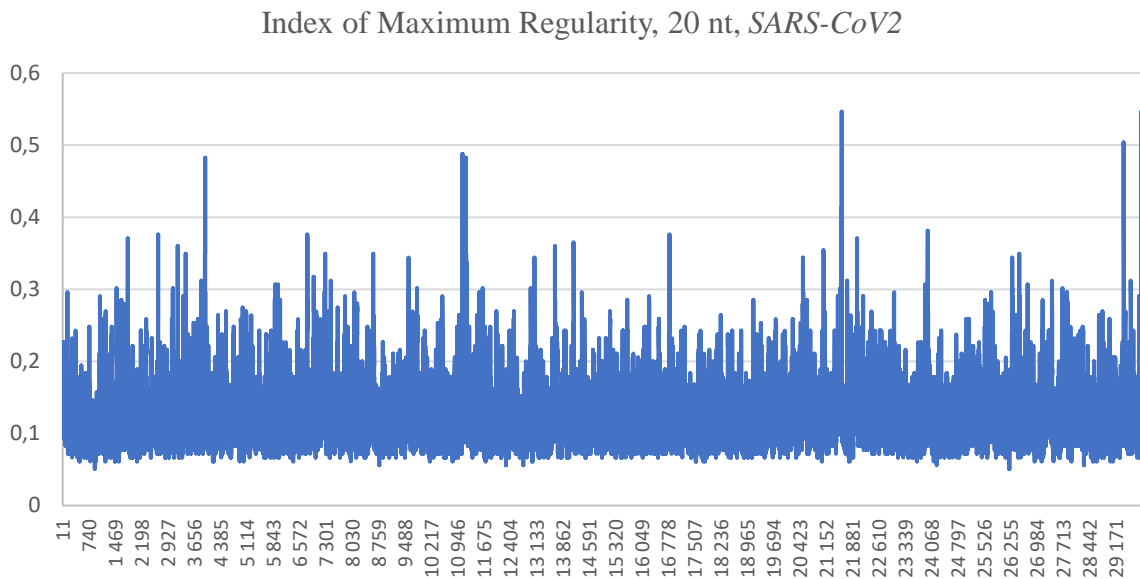
For a visualization on the components of Coronavirus 2, the configuration of the NCBI (2020) is shown.

Image 1: Diagram on the structure of Coronavirus 2.



Note: From [6].

Graph 1 presents the analysis made for 20nt intervals



Graph 1: Index of Maximum Regularity, 16 nt, SARS-CoV2

In the indicated regions, there are intervals that take higher values (Table 1).

Table 1: Index of Maximum Regularity, 20 nt, SARS-CoV2 vs NCBI

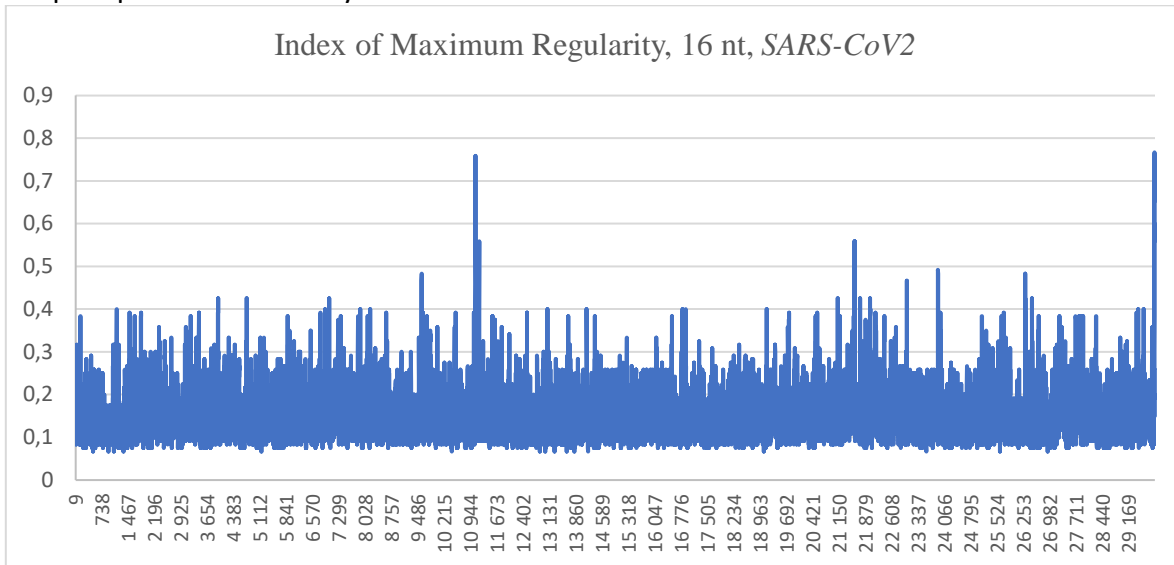
Interval (middle point)	$i_{max,r}$	Already classified by NCBI
21578,5	0,546666667	Gene S (start)
29875,5	0,546666667	Gene ORF10
29876,5	0,546666667	Gene ORF10
29877,5	0,546666667	Gene ORF10
29390,5	0,504	Gene N
29882,5	0,504	Gene ORF10
29883,5	0,504	Gene ORF10
11077,5	0,488	Undefined
3952,5	0,482666667	Undefined
11075,5	0,482666667	Undefined
11076,5	0,482666667	Undefined
11078,5	0,482666667	Undefined
11079,5	0,482666667	Undefined
11080,5	0,482666667	Undefined
11180,5	0,482666667	Undefined
29880,5	0,482666667	Gene ORF10
29881,5	0,482666667	Gene ORF10

Thus, it is evident that in the areas near the intervals around 21578, 29876, 29390, 3952, 11076 have high values. As consulted in the database, this is positive because **Gene S**, **Gene N** and **ORF10** were identified. It is important to research in more detail what happens around **11078** interval (20 nt). It should be clarified that **Gene: Orf1ab** is in the range [269,

21 558] (71% of the entire genome) which is extremely wide and the $i_{max,r}$ indicates something of importance around 11000

The following is the complete genome analysis with 16 nt. (Graph 2 and Table 2)

Graph 2 presents the analysis made for 16 nt intervals



Graph 2: Index of Maximum Regularity, 16 nt, SARS-CoV2

Table 2: Index of Maximum Regularity, 16 nt, SARS-CoV2 vs NCBI

Interval (middle point)	$i_{max,r}$	Already classified by NCBI
29881,5	0,766666667	Gene ORF10
29882,5	0,766666667	Gene ORF10
11078,5	0,758333333	Undefined
11079,5	0,758333333	Undefined
29878,5	0,758333333	Gene ORF10
29879,5	0,758333333	Gene ORF10
29880,5	0,65	Gene ORF10
29884,5	0,6	Gene ORF10
11076,5	0,558333333	Undefined
11077,5	0,558333333	Undefined
11182,5	0,558333333	Undefined
21576,5	0,558333333	Gene S (start)
21577,5	0,558333333	Gene S (start)
21581,5	0,558333333	Gene S (start)
29883,5	0,558333333	Gene ORF10
11080,5	0,55	Undefined
11081,5	0,55	Undefined
11082,5	0,55	Undefined
11083,5	0,55	Undefined
21580,5	0,55	Gene S (start)
29875,5	0,55	Gene ORF10
29876,5	0,55	Gene ORF10
29877,5	0,55	Gene ORF10

In this string size, **Gene S (start)** and **Gene ORF10** are identified. And again, the $i_{max,r}$ points to an important zone around the **11000** interval.

When calculating with a length interval of 28 nt, the sectors around **3500** and **11000** appear with high values. Longer chains are not considered because the probability of occurrence decreases with the number of combinations of the four possibilities.

Conclusions

During the analysis **Gene S**, **Gene N** and **ORF10** were identified. The interval around **3500** and the interval around **11000**, respectively, appear with high values of regularity and this do not correspond to documented elements (both within Gene: **Orf1ab**). It might be in the interest of giving importance to that region.

The strategy developed here may offer insights into Coronavirus 2 and eventually, with much more research and sophistication, a treatment may be found.

Acknowledgements

To the memory of Dra. Tatiana Láscaris-Comneno-Slepuin.

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