

Atomic Hemoglobin Code

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Abstract—This paper discusses cyberinformation studies of the amino acid composition of *Hemoglobin protein sequences Q6WN27*, in particular the identification of scientific terminology that could describe this phenomenon, ie, the study of genetic information, as well as the relationship between the genetic language of hemoglobin and theoretical aspect of this system and cybernetics. The result of this research show that there is a matrix code for hemoglobin. It also shows that the coding system within the amino acidic language gives detailed information, not only on the amino acid „record“, but also on its structure, configuration and its various shapes. The issue of the existence of an hemoglobin code and coding of the individual structural elements of this protein are discussed. Answers to the following questions are sought. Does the matrix mechanism for biosynthesis of this protein function within the law of the general theory of information systems, and what is the significance of this for understanding the genetic language of hemoglobin? What is the essence of existence and functioning of this language? Is the genetic information characterized only by biochemical, or also by cyberinformation principles? The potential effects of physical and chemical, as well as cybernetic and information principles, on the biochemical basis of hemoglobin are also investigated. This paper discusses new methods for developing genetic technologies, in particular more advanced digital technology based on programming, cybernetics, and informational laws and systems, and how this new technology could be useful in medicine, bioinformatics, genetics, biochemistry, and other natural sciences.

I. INTRODUCTION

The biologic role of any given protein in essential life processes, eg, *Hemoglobin protein sequences Q6WN27*, depends on the positioning of its component amino acids, and is understood by the „positioning of letters forming words—Each of these words has its biochemical base. If this base is expressed by corresponding discrete numbers, it can be seen that any given base has its own program, along with its own unique cybernetics and information characteristics.

Indeed, the sequencing of the molecule is determined not only by distinct biochemical features, but also by cybernetic and information principles. For this reason, research in this field deals more with the quantitative rather than qualitative characteristics of genetic information and its biochemical basis. For the purposes of this paper, specific physical and chemical factors have been selected in order to express the genetic information for *Hemoglobin protein sequences Q6WN27*. Numerical values are then assigned to these factors, enabling them to be measured.

In this way it is possible to determine if connection really exists between the quantitative ratios in the process of transfer of genetic information and the qualitative appearance of the *Hemoglobin protein sequences Q6WN27* molecule. To select these factors, preference is given to classical physical and chemical parameters, including the number of atoms in the relevant amino acids, their analog values, the position in these amino acids in the peptide chain, and their frequencies. There is a large number of these parameters, and each of them gives important genetic information. Going through this process, it becomes clear that there is a mathematical relationship between quantitative ratios and the qualitative appearance of the biochemical „genetic processes—and that there is a measurement method that can be used to describe the biochemistry of *Hemoglobin protein sequences Q6WN27*.

II. METHODS

Hemoglobin protein sequences Q6WN27 can be represented by two different forms, ie, a discrete form and a sequential form. In the discrete form, a molecule of *Hemoglobin protein sequences Q6WN27* is represented by a set of discrete codes or a multiple dimension vector. In the sequential form, an *Hemoglobin protein sequences Q6WN27* molecule is represented by a series of amino acids according to the order of their position in the *Hemoglobin protein sequences Q6WN27*. Therefore, the sequential form can naturally reflect all the information about the sequence order and length of an *Hemoglobin protein sequences Q6WN27* molecule. The key issue is whether we can develop a different discrete method of representing an *Hemoglobin protein sequences Q6WN27* molecule that will allow accommodation of partial, if not all sequence order information? Because a protein sequence is usually represented by a series of amino acids should be assigned to these codes in order to optimally convert the sequence order information into a series of numbers for the discrete form representation? **Expression of Hemoglobin protein sequences Q6WN27** The matrix mechanism of *Hemoglobin protein sequences Q6WN27*, the evolution of biomacromolecules and, especially, the biochemical evolution of *Hemoglobin protein sequences Q6WN27* language, have been analyzed by the application of cybernetic methods, information theory and system theory, respectively. The primary structure of a molecule of *Hemoglobin protein sequences Q6WN27* is the exact specification of its atomic composition and the chemical bonds connecting those atoms.

Characteristics: Length : 147, molecular weight: 16043, CRC64 check sum: 1EC2AB4138BA7899

Sequence:>Q6WN27 Hemoglobin subunit beta GAFSDGLAHLNLDNLKGTFAQLSELHCDKLHVDPENFR
(Hemoglobin beta chain) (Beta-globin). LLGNVLVCVLAHHFGKEFTPQVQAAYQKVVAGVAN
MVHLTGEKAAVTALWGKVNVDVGGGEALGRLLV ALAHKYH.
VYPWTQRFESFGDLSSPDVMNNPKVKAHGKKVL

Fragment of sequence 147 AA;

Number of atoms															
M	V	H	L	T	G	E	E	K	A	.	.	H	K	Y	Sum
20	19	20	22	17	10	19	19	24	13	.	.	20	24	24	2699
1	2	3	4	5	6	7	8	9	10	.	.	144	145	146	10878

Amino acid 1 (AC1) = 20 atoms; AC2=19 atoms; AC3= 20 atoms, etc.

Figure 1. Group of amino acids from 1 to 147.

Notes: Aforementioned aminoacids are positioned from number 1 to 147. Numbers 1, 2, 3, n... present the position of a certain aminoacid. This positioning is of the key importance for understanding of programmatic, cybernetic and information principles in this protein. The scientific key for interpretation of bio chemical processes is the same for **Hemoglobin protein sequences Q6WN27** and as well as for the other proteins and other sequences in biochemistry.

The first aminoacid in this example has 20 atoms, the second one 19, the third one 20, etc. They have exactly these numbers of atoms because there are many codes in the **Hemoglobin protein sequences Q6WN27**, analog codes, and other coded features. In fact, there is a cybernetic algorithm which it is „recorded—that the first amino acid has to have 20 atoms, the second one 19, the third one 20, etc.

The first amino acid has its own biochemistry, as does the second and the third, etc. The obvious conclusion is that there is a concrete relationship between quantitative ratios in the process of transfer of genetic information and qualitative appearance, ie, the characteristics of the organism.

III. ALGORITHM

We shall now give some mathematical evidences that will prove that in the biochemistry of insulin there really is programmatic and cybernetic algorithm in which it is „recorded— in the language of mathematics, how the molecule will be built and what will be the quantitative characteristics of the given genetic information.

Arithmetic progression

Step 1 (Amino acids from 1 to 147)

$AC_1 = 20 \text{ atoms}; AC_2 = 19 \text{ atoms}; AC_3 = 20 \text{ atoms}; \dots AC_{147} = 20 \text{ atoms};$

$[AC_1 + (AC_1 + AC_2) + (AC_1 + AC_2 + AC_3) \dots + (AC_1 + AC_2 + AC_3 \dots + AC_{147})] = S_1;$

$AC_1 = APa_1 = 20;$

$(AC_1 + AC_2) = (20 + 19) = APa_2 = 39;$

$(AC_1 + AC_2 + AC_3) = (20 + 19 + 20) = APa_3 = 59;$

$(AC_1 + AC_2 + AC_3 \dots + AC_{147}) = APa_{147} = 2699 \text{ atoms};$

$APa_{1,2,3,n} = \text{Arithmetic progression of amino acids } 1,2,3,n$

$[APa_1 + APa_2 + APa_3 \dots + APa_{147}] = (20 + 39 + 59 \dots + 2699) = 199\,076;$

$S_1 = 199076;$

Example :

Arithmetic progression 1 (APa)															
M	V	H	L	T	G	E	E	.	.	K	Y	H	Sum		
20	19	20	22	17	10	19	19	.	.	24	24	20	2699		
1	2	3	4	5	6	7	8	.	.	145	146	147	10878		

↓

M	V	H	L	T	G	E	E	.	.	K	Y	H	Sum		
20	39	59	81	98	108	127	146	.	.	2631	2655	2679	199076		
1	2	3	4	5	6	7	8	.	.	145	146	147	10878		

$(0+20) = 20; (20+19)=39; (20+19+20) = 59; \text{ etc.}$

Figure 1. Arithmetic progression 1 (APa) of amino acids from 1 to 147.

Notes: By using chemical-information procedures, we calculated the arithmetic progression for the information content of aforementioned aminoacids.

Step 2 (Amino acids from 147 to 1)

$$AC_{147} = 20 \text{ atoms}; AC_{146} = 24 \text{ atoms}; AC_{145} = 24 \text{ atoms}; \dots AC_1 = 20 \text{ atoms};$$

$$[AC_{147} + (AC_{147} + AC_{146}) + (AC_{147} + AC_{146} + AC_{145}) \dots + (AC_{147} + AC_{146} + AC_{145} \dots + AC_1)] = S_2;$$

$$AC_{147} = APb_{147} = 20;$$

$$(AC_{147} + AC_{146}) = (20 + 24) = APb_{146} = 44;$$

$$(AC_{147} + AC_{146} + AC_{145}) = (20 + 24 + 24) = APb_{145} = 68;$$

$$(AC_{147} + AC_{146} + AC_{145} \dots + AC_1) = APb_1 = 2699 \text{ atoms};$$

$$APb_{147,146,145, \dots, 1} = \text{Arithmetic progression of amino acids } 147, 146, 145, \dots, 1;$$

$$[APb_{147} + APb_{146} + APb_{145} \dots + APb_1] = (20 + 44 + 68 \dots + 2699) = 200\,376;$$

$$S_2 = 200\,376;$$

Example:

Arithmetic progression 2 (APb)													
Sum	M	V	H	.	.	N	A	L	A	H	K	Y	H
2699	20	19	20	.	.	17	13	22	13	20	24	24	20
10878	1	2	3	.	.	140	141	142	143	144	145	146	147
↓													
Sum	M	G	D	.	.	D	E	E	A	G	E	G	N
200 376	2699	2679	2660	.	.	153	136	123	101	88	68	44	20
10878	1	2	3	.	.	140	141	142	143	144	145	146	147

(0+20) = 20; (20+24)=44; (20+24+24)=68; etc.

Figure 2. Schematic representation of the arithmetic progression 2 from 147 to 1.

Within the digital pictures in biochemistry, the physical and chemical parameters are in a strict compliance with programmatic, cybernetic and information principles. Each bar in the protein chain attracts only the corresponding aminoacid, and only the relevant aminoacid can be positioned at certain place in the chain. Each peptide chain can have the exact number of aminoacids necessary to meet

the strictly determined mathematical conditioning. It can have as many atoms as necessary to meet the mathematical balance of the biochemical phenomenon at certain mathematical level, etc... the digital language of biochemistry has a countless number of codes and analogue codes, as well as other information content. These pictures enable us to realize the very essence of functioning of biochemical processes. There are some examples.

Table 1.Bio intervals APa and APb (sequence from 1 to 147 AA)

Sequence												
M	V	H	L	T	G	E	E	K	A	A	V	T
D	E	V	G	G	E	A	L	G	R	L	L	V
F	E	S	F	G	D	L	S	S	P	D	A	V
H	G	K	K	V	L	G	A	F	S	D	G	L
T	F	A	Q	L	S	E	L	H	C	D	K	L
L	L	G	N	V	L	V	C	V	L	A	H	H
V	Q	A	A	Y	Q	K	V	V	A	G	V	A
	M	D	H	G	K	L	K	G	P	Y	H	Sum
	20	16	20	10	24	22	24	10	17	24	20	207
	1	22	64	65	66	82	83	84	125	146	147	
	↓	↓	↓	↓	↓	↓	↓	↓	↓	↓	↓	
APa	20	399	1174	1184	1208	1491	1515	1525	2300	2679	2699	> 16194
	↓	↓	↓	↓	↓	↓	↓	↓	↓	↓	↓	
	-2679	-2280	-1126	-341	-307	0	+307	+341	+1126	+2280	+2679	
	↑	↑	↑	↑	↑	↑	↑	↑	↑	↑	↑	
APb	2699	2679	2300	1525	1515	1491	1208	1184	1174	399	20	> 16194
	↑	↑	↑	↑	↑	↑	↑	↑	↑	↑	↑	
	M	V	E	G	K	K	K	G	T	Q	H	Sum
	20	19	19	10	24	24	24	10	17	20	20	207
	1	2	23	65	66	67	83	84	85	126	147	

$(20-2699) = (-)2679$; $(399-2679) = (-)2280$; $(1174-2300) = (-)1126$; etc.
 $APa1=20$; $APa22=399$; $APa64=1174$; etc.
 $APb1=2699$; $APb2=2679$; $APb23=2300$; etc.
 $APa1= APb147$; $APa22= APb126$;
 $APa64= APb85$; $APa65= APb84$; $APa66 = APb83$
 $APa82 = APb67$; $APa83 = APb66$; $APa84 = APb65$; $APa125 = APb23$;
 etc.

Table 1. Schematic representation of the bio intervals APa and APb (sequence from 1 to 147 AA).

Notes: Namely, having mathematically analyzed the Bio intervals model of *Hemoglobin protein sequences Q6WN27* (Table 1) we have found out that the protein code is based on a periodic law. This being the only to „read—the picture, the solution of the main problem (concerning an

arrangement where each amino acid takes only one, precisely determined position in the code), is quite manifest: Bio intervals model of *Hemoglobin protein sequences Q6WN27* should, in fact, be „remodelled—into a periodic system.

Table 2. Bio intervals Apa and APb (sequence from 2 to 146 AA)

M	V	H	L	T	G	E	E	K	A	A	V	T	A	L	W	G	K	V	N	V
D	E	V	G	G	E	A	L	G	R	L	L	V	V	Y	P	W	T	Q	R	F
F	E	S	F	G	D	L	S	S	P	D	A	V	M	N	N	P	K	V	K	A
H	G	K	K	V	L	G	A	F	S	D	G	L	A	H	L	D	N	L	K	G
T	F	A	Q	L	S	E	L	H	C	D	K	L	H	V	D	P	E	N	F	R
L	L	G	N	V	L	V	C	V	L	A	H	H	F	G	K	E	F	T	P	Q
V	Q	A	A	Y	Q	K	V	V	A	G	V	A	N	A	L	A	H	K	Y	H

	D	H	G	K	L	K	G	P	Y	Sum	
	16	20	10	24	22	24	10	17	24	167	
	22	64	65	66	82	83	84	125	146		
	↓	↓	↓	↓	↓	↓	↓	↓	↓		
APa	379	1154	1164	1188	1471	1495	1505	2280	2659	>	13295
	↓	↓	↓	↓	↓	↓	↓	↓	↓		
	-2280	-1126	-341	-307	0	+307	+341	+1126	+2280		
	↑	↑	↑	↑	↑	↑	↑	↑	↑		
APb	2659	2280	1505	1495	1471	1188	1164	1154	379	>	13295
	↑	↑	↑	↑	↑	↑	↑	↑	↑		
	V	E	G	K	K	K	G	T	Q	Sum	
	19	19	10	24	24	24	10	17	20	167	
	2	23	65	66	67	83	84	85	126		

Table 2. Schematic representation of the bio intervals APa and APb (sequence from 2 to 146 AA).**Table 3. Bio intervals Apa and APb (sequence from 3 to 145 AA)**

M	V	H	L	T	G	E	E	K	A	A	V	T	A	L	W	G	K	V	N	V
D	E	V	G	G	E	A	L	G	R	L	L	V	V	Y	P	W	T	Q	R	F
F	E	S	F	G	D	L	S	S	P	D	A	V	M	N	N	P	K	V	K	A
H	G	K	K	V	L	G	A	F	S	D	G	L	A	H	L	D	N	L	K	G
T	F	A	Q	L	S	E	L	H	C	D	K	L	H	V	D	P	E	N	F	R
L	L	G	N	V	L	V	C	V	L	A	H	H	F	G	K	E	F	T	P	Q
V	Q	A	A	Y	Q	K	V	V	A	G	V	A	N	A	L	A	H	K	Y	H

	R	P	L	H	K	
	26	17	22	20	24	
	31	59	89	117	145	
	↓	↓	↓	↓	↓	
APa	508	1035	1581	2108	2616	> 7848
	↓	↓	↓	↓	↓	
	-2108	-1073	0	+1073	+2108	
	↑	↑	↑	↑	↑	
APb	2616	2108	1581	1035	508	> 7848
	↑	↑	↑	↑	↑	
	H	L	K	S	H	
	20	22	24	14	20	
	3	32	60	90	118	

Table 3. Schematic representation of the bio intervals APa and APb (sequence from 3 to 145 AA).**Table 4. Bio intervals Apa and APb (sequence from 4 to 144 AA)**

M	V	H	L	T	G	E	E	K	A	A	V	T	A	L	W	G	K	V	N	V
D	E	V	G	G	E	A	L	G	R	L	L	V	V	Y	P	W	T	Q	R	F
F	E	S	F	G	D	L	S	S	P	D	A	V	M	N	N	P	K	V	K	A
H	G	K	K	V	L	G	A	F	S	D	G	L	A	H	L	D	N	L	K	G
T	F	A	Q	L	S	E	L	H	C	D	K	L	H	V	D	P	E	N	F	R
L	L	G	N	V	L	V	C	V	L	A	H	H	F	G	K	E	F	T	P	Q
V	Q	A	A	Y	Q	K	V	V	A	G	V	A	N	A	L	A	H	K	Y	H

	E	G	A	A	E	N	H	
	19	10	13	13	19	17	20	
	7	25	71	77	122	140	144	
	↓	↓	↓	↓	↓	↓	↓	
APa	68	388	1237	1335	2184	2504	2572	> 10288
	↓	↓	↓	↓	↓	↓	↓	
	-2504	-2116	-947	0	+947	+2116	+2504	
	↑	↑	↑	↑	↑	↑	↑	
APb	2572	2504	2184	1335	1237	388	68	> 10288
	↑	↑	↑	↑	↑	↑	↑	
	L	E	G	F	H	F	A	
	22	19	10	23	20	23	13	
	4	8	26	72	78	123	141	

Table 4. Schematic representation of the bio intervals APa and APb (sequence from 4 to 144 AA).

Table 5. Bio intervals APa and APb (sequence from 5 to 143 AA)

M	V	H	L	T	G	E	E	K	A	A	V	T	A	L	W	G	K	V	N	V
D	E	V	G	G	E	A	L	G	R	L	L	V	V	Y	P	W	T	Q	R	F
F	E	S	F	G	D	L	S	S	P	D	A	V	M	N	N	P	K	V	K	A
H	G	K	K	V	L	G	A	F	S	D	G	L	A	H	L	D	N	L	K	G
T	F	A	Q	L	S	E	L	H	C	D	K	L	H	V	D	P	E	N	F	R
L	L	G	N	V	L	V	C	V	L	A	H	H	F	G	K	E	F	T	P	Q
V	Q	A	A	Y	Q	K	V	V	A	G	V	A	N	A	L	A	H	K	Y	H

	E	S	G	A	A		
	19	14	10	13	13		
	8	73	75	139	143		
	↓	↓	↓	↓	↓		
APa	65	1252	1278	2465	2530	>	7590
	↓	↓	↓	↓	↓		
	-2465	-1213	0	+1213	+2465		
	↑	↑	↑	↑	↑		
APb	2530	2465	1278	1252	65	>	7590
	↑	↑	↑	↑	↑		
	T	K	D	L	N		
	17	24	16	22	17		
	5	9	74	76	140		

Table 5. Schematic representation of the bio intervals APa and APb (sequence from 5 to 143 AA).

Table 6. Bio intervals APa and APb (sequence from 6 to 142 AA)

M	V	H	L	T	G	E	E	K	A	A	V	T	A	L	W	G	K	V	N	V
D	E	V	G	G	E	A	L	G	R	L	L	V	V	Y	P	W	T	Q	R	F
F	E	S	F	G	D	L	S	S	P	D	A	V	M	N	N	P	K	V	K	A
H	G	K	K	V	L	G	A	F	S	D	G	L	A	H	L	D	N	L	K	G
T	F	A	Q	L	S	E	L	H	C	D	K	L	H	V	D	P	E	N	F	R
L	L	G	N	V	L	V	C	V	L	A	H	H	F	G	K	E	F	T	P	Q
V	Q	A	A	Y	Q	K	V	V	A	G	V	A	N	A	L	A	H	K	Y	H

	L	L	W	T	D	K	A
	22	22	27	17	16	24	13
	15	32	38	39	53	60	63
	↓	↓	↓	↓	↓	↓	↓
APa	169	471	599	616	873	1000	1056
	↓	↓	↓	↓	↓	↓	↓
	-2331	-1860	-1430	-1285	-1011	-627	-444
	↑	↑	↑	↑	↑	↑	↑
APb	2500	2331	2029	1901	1884	1627	1500
	↑	↑	↑	↑	↑	↑	↑
	G	W	L	T	Q	A	V
	10	27	22	17	20	13	19
	6	16	33	39	40	54	61

T
17
65
↓
1444
↓
Q
↓
1444
↑
T
H
20
64

Q	D	G	N	A	Q	L
20	16	10	17	13	20	22
88	95	108	109	116	132	142
↓	↓	↓	↓	↓	↓	↓
1500	1627	1884	1901	2029	2331	2500
↓	↓	↓	↓	↓	↓	↓
+444	+627	+1011	+1285	+1430	+1860	+2331
↑	↑	↑	↑	↑	↑	↑
1056	1000	873	616	599	471	169
↑	↑	↑	↑	↑	↑	↑
F	L	K	N	V	H	K
23	22	24	17	19	20	24
86	89	96	109	110	117	133

Table 6. Schematic representation of the bio intervals APa and APb (sequence from 6 to 142 AA).

Table 7. Bio intervals APa and APb (sequence from 7 to 141 AA)

M	V	H	L	T	G	E	E	K	A	A	V	T	A	L	W	G	K	V	N	V
D	E	V	G	G	E	A	L	G	R	L	L	V	V	Y	P	W	T	Q	R	F
F	E	S	F	G	D	L	S	S	P	D	A	V	M	N	N	P	K	V	K	A
H	G	K	K	V	L	G	A	F	S	D	G	L	A	H	L	D	N	L	K	G
T	F	A	Q	L	S	E	L	H	C	D	K	L	H	V	D	P	E	N	F	R
L	L	G	N	V	L	V	C	V	L	A	H	H	F	G	K	E	F	T	P	Q
V	Q	A	A	Y	Q	K	V	V	A	G	V	A	N	A	L	A	H	K	Y	H

	K	N	E	Q	K	D	L	F	Q	G	A
	24	17	19	20	24	16	22	23	20	10	13
	9	20	23	40	67	80	106	123	126	137	141
	↓	↓	↓	↓	↓	↓	↓	↓	↓	↓	↓
APa	62	286	310	626	1124	1344	1842	2188	2212	2406	2468
	↓	↓	↓	↓	↓	↓	↓	↓	↓	↓	↓
	-2406	-2150	-1902	-1532	-718	0	718	1532	1902	2150	2406
	↑	↑	↑	↑	↑	↑	↑	↑	↑	↑	↑
APb	2468	2406	2212	2158	1842	1344	1124	626	310	256	62
	↑	↑	↑	↑	↑	↑	↑	↑	↑	↑	↑
	E	A	V	V	R	V	N	L	T	V	V
	19	13	19	19	26	19	17	22	17	19	19
	7	10	21	24	41	68	81	107	124	127	138

Table 7. Schematic representation of the bio intervals APa and APb (sequence from 4 to 144 AA).

Notes:Consequently, the periodic law allows for only one form of the amino acid bio interval (Table 1,2,3,4,5,6 and 4) which can show both the relationships among the amino acids, and their position in peptide chain. As we see, protein code can be developed into the periodic system (and the table), in which each of 20 natural amino acids has an exactly established place.etc.Biological particularity of proteins depends on the order of amino acids in their molecules. Change of that order will lead to the change of their biological particularity. The base parameters that determine the change of status of biosynthesis matrix of macromolecules are the system entropy, volume of information transferred and the level of probability that the genetic information will be transferred as a whole.Each peptide chain can have as many atoms as necessary to meet the mathematical balance of the biochemical phenomenon at certain mathematical level. By using the system and information procedures and methods in studying biochemistry, we can analyze the effects of the classical and information parameters in the protein evolution process. We would particularly like to stress here that the genetic, as well as biochemical information in a broader sense of the word, is determined and characterized by very complex cybernetic and information principles. The constantans in those principles are: the number of atoms and molecules, atomic numbers, atomic weight, physical and chemical parameters, even and odd values, codes and analogue codes, standard deviations, frequencies, primary and secondary values, and many other things.

IV. CONCLUSION

The results of our research show that the processes of sequencing the molecules are conditioned and arranged not only with chemical and biochemical lawfulness, but also with program, cybernetic and informational lawfulness too. At the first stage of our research we replaced nucleotides from the Amino Acid Code Matrix with numbers of the atoms in those nucleotides. Translation of the biochemical language of these amino acids into a digital language may be very useful for developing new methods of predicting protein sub-cellular localization, membrane protein type, protein structure secondary prediction or any other protein attributes. Since the concept of Chou's pseudo amino acid composition was proposed^{1,2}, there have been many efforts to try to use various digital numbers to represent the 20 native amino acids in order to better reflect the sequence-order effects through the vehicle of pseudo amino acid composition. Some investigators used complexity measure factor³, some used the values derived from the cellular automata⁴⁻⁷, some used hydrophobic and/or hydrophilic values⁸⁻¹⁶, some were through Fourier transform^{17,18}, and some used the physicochemical distance¹⁹. It is going to be possible to use a completely new strategy of research in genetics in the future. However, close observation of all these relationships, which are the outcomes of periodic laws (more specifically the law of binary coding), stereochemical and digital structure of proteins.

V. DISCLOSURE

The author reports no conflict of interest in this research.

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