

GLOBAL JOURNAL OF MEDICAL RESEARCH PHARMA, DRUG DISCOVERY, TOXICOLOGY AND MEDICINE

Volume 13 Issue 2 Version 1.0 Year 2013

Type: Double Blind Peer Reviewed International Research Journal

Publisher: Global Journals Inc. (USA)

Online ISSN: 2249-4618 & Print ISSN: 0975-5888

The Insulin Bio Intervals

By Lutvo Kurić

Centro Universitário Feevale

Abstract - The modern science mainly treats the biochemical basis of sequencing in bio-macromolecules and processes in medicine and biochemistry. One can ask weather the language of biochemistry is the adequate scientific language to explain the phenomenon in that science. Is there maybe some other language, out of biochemistry, that determines how the biochemical processes will function and what the structure and organization of life systems will be? The research results provide some answers to these questions. They reveal to us that the process of sequencing in bio-macromolecules is conditioned and determined not only through biochemical, but also through cybernetic and information principles. Many studies have indicated that analysis of protein sequence codes and various sequence-based prediction approaches, such as predicting drug-target interaction networks (He et al., 2010), predicting functions of proteins (Hu et al., 2011; Kannan et al., 2008), analysis and prediction of the metabolic stability of proteins (Huang et al., 2010), predicting the network of substrate-enzyme-product triads (Chen et al., 2010), membrane protein type prediction (Cai and Chou, 2006; Cai et al., 2003; Cai et al., 2004), protein structural class prediction (Cai et al., 2006; Ding et al., 2007), protein secondary structure prediction (Chen et al., 2009; Ding et al., 2009b), enzyme family class prediction (Cai et al., 2005; Ding et al., 2009a; Wang et al., 2010), identifying cyclin proteins (Mohabatkar, 2010), protein subcellular location prediction (Chou and Shen, 2010a; Chou and Shen, 2010b; Kandaswamy et al., 2010; Liu et al., 2010), among many others as summarized in a recent review (Chou, 2011), can timely provide very useful information and insights for both basic research and drug design and hence are widely welcome by science community. The present study is attempted to develop a novel sequence-based method for studying insulin in hopes that it may become a useful tool in the relevant areas.

Keywords: human insulin, algorithm, insulin model, insulin code, bio interval.

GJMR-B Classification : NLMC Code: WK 820



Strictly as per the compliance and regulations of:



© 2013. Lutvo Kurić. This is a research/review paper, distributed under the terms of the Creative Commons Attribution-Noncommercial 3.0 Unported License http://creativecommons.org/licenses/by-nc/3.0/), permitting all non-commercial use, distribution, and reproduction inany medium, provided the original work is properly cited.

The Insulin Bio Intervals

Lutvo Kurić

Abstract - The modern science mainly treats the biochemical basis of sequencing in bio-macromolecules and processes in medicine and biochemistry. One can ask weather the language of biochemistry is the adequate scientific language to explain the phenomenon in that science. Is there maybe some other language, out of biochemistry, that determines how the biochemical processes will function and what the structure and organization of life systems will be? The research results provide some answers to these questions. They reveal to us that the process of sequencing in bio-macromolecules is conditioned and determined not only through biochemical, but also through cybernetic and information principles. Many studies have indicated that analysis of protein sequence codes and various sequence-based prediction approaches, such as predicting drug-target interaction networks (He et al., 2010), predicting functions of proteins (Hu et al., 2011; Kannan et al., 2008), analysis and prediction of the metabolic stability of proteins (Huang et al., 2010), predicting the network of substrate-enzyme-product triads (Chen et al., 2010), membrane protein type prediction (Cai and Chou, 2006; Cai et al., 2003; Cai et al., 2004), protein structural class prediction (Cai et al., 2006; Ding et al., 2007), protein secondary structure prediction (Chen et al., 2009; Ding et al., 2009b), enzyme family class prediction (Cai et al., 2005; Ding et al., 2009a; Wang et al., 2010), identifying cyclin proteins (Mohabatkar, 2010), protein subcellular location prediction (Chou and Shen, 2010a; Chou and Shen, 2010b; Kandaswamy et al., 2010; Liu et al., 2010), among many others as summarized in a recent review (Chou, 2011), can timely provide very useful information and insights for both basic research and drug design and hence are widely welcome by science community. The present study is attempted to develop a novel sequence-based method for studying insulin in hopes that it may become a useful tool in the relevant areas.

Keywords: human insulin, algorithm, insulin model, insulin code, bio interval.

I. Introduction

he biologic role of any given protein in essential life processes, eg, insulin, 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 distin 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 insulin. Numerical values are them assigned to these factors, enabling them to be measured. In this way it is possible to determine oif a connection really exists between the quantitative ratios in the process of transfer of genetic information and the qualitative appearance of the insulin 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 frenquencies. There is a arge numbers of these parameters, and each of their 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 insulin.

II. METHODS

The biologic role of any given protein in essential life processes, eg, insulin, 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 distin 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 insulin. Numerical values are them assigned to these factors, enabling them to be measured. In this way it is possible to determine oif a connection really exists between the quantitative ratios in the process of transfer of genetic information and the qualitative appearance of the insulin 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 frenquencies. There is a arge numbers of these parameters, and each of their 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 insulin.

Insulin can be represented by two different forms, ie, a discrete form and a sequential form. In the discrete form, a molecule of insulin is represented by a set of discrete codes or a multiple dimension vector. In the sequential form, an insulin molecule is represent by a series of amino acids according to the order of their position in the **chains 1Alo.**

Therefore, the sequential form can naturally reflect all the information about the sequence order and lenght of an insulin molecule. The key issue is whether we can develop a different discrete method of representing an insulin molecule that will allow accomodation 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?

Insulin can be represented by two different forms, ie, a discrete form and a sequential form.

Therefore, the sequential form can naturally reflect all the information about the sequence order and lenght of an insulin molecule. The key issue is whether we can develop a different discrete method of representing an insulin molecule that will allow accomodation 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?

III. Expression of Insulin Code Matrix- 1ai0

The matrix mechanism of Insulin, the evolution of biomacromolecules and, especially, the biochemical evolution of Insulin language, have been analyzed by the application of cybernetic methods, information theory and system theory, respectively. The primary structure of a molecule of Insulin is the exact specification of its atomic composition and the chemical bonds connecting those atoms.

R6 INSULIN HEXAMER (d1ai02)

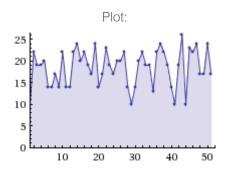
The structure 1Al0 has in total 12 chains. Out of these 2 are sequence-unique

>1AI0:A	GIVEQCCTSICSLYQLENYCN
>1AI0:B	FVNQHLCGSHLVEALYLVCGERGFFYTPKT
>1AI0:C	GIVEQCCTSICSLYQLENYCN
>1AI0:D	FVNQHLCGSHLVEALYLVCGERGFFYTPKT
>1AI0:E	GIVEQCCTSICSLYQLENYCN
>1AI0:F	FVNQHLCGSHLVEALYLVCGERGFFYTPKT
>1AI0:G	GIVEQCCTSICSLYQLENYCN
>1AI0:H	FVNQHLCGSHLVEALYLVCGERGFFYTPKT
>1AI0:I	GIVEQCCTSICSLYQLENYCN
>1AI0:J	FVNQHLCGSHLVEALYLVCGERGFFYTPKT
>1AI0:K	GIVEQCCTSICSLYQLENYCN
>1AI0:L	FVNQHLCGSHLVEALYLVCGERGFFYTPKT

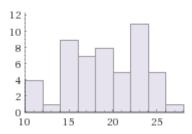
G I Ε Q C C Т S ı C S L Υ Q L Ε Ν Υ C Ν F Ν Q Н I C G S Н L Ε С Ε G F Ρ Т Α Υ G R Υ Τ K

Input:

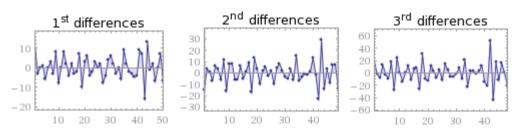
{10, 22, 19, 19, 20, 14, 14, 17, 14, 22, 14, 14, 22, 24, 20, 22, 19, 17, 24, 14, 17, 23, 19, 17, 20, 20, 22, 14, 10, 14, 20, 22, 19, 19, 13, 22, 24, 22, 19, 14, 10, 19, 26, 10, 23, 22, 24, 17, 17, 24, 17}



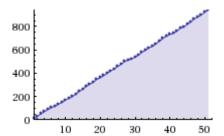
Histogram:



Differences:



Cumulative sums:



As wesee, the genetic information characterized only by biochemical, or also by cyberinformation principles

Fragment 1

From 1 to 102

G	ı	V	Е	Q	С	С	Т
10	22	19	19	20	14	14	 17
		3					

Fragment 2

From 103 to 204

G	ı	٧	Е	Q	С	С	Т
10	22	19	19	20	14	14	 17
103	104	105	106	107	108	109	204

Fragment 3

From 205 to 306

G	1	٧	Е	Q	С	С	Т
10	22	19	19	20	14	14	 17
205	206	207	208	209	210	211	306

Aforementioned aminoacids are positioned from number 1 to 306. 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 insulin and as well as for the other proteins and other sequences in biochemistry. The first aminoacid in this example has 10 atoms, the second one 22, the third one 19, etc... Why do they have exactly this many atoms? It is because there are many codes in the molecule of insulin, analogue codes and other coded features. In fact, there is a programcybernetic algorithm in which it is "recorded" that the firs amino acid has to have 10 atoms, the second one 22, the third one 19, etc... The first amino acid has its own

biochemistry, the second and the third one also. The conclusion here has to be that there is a concrete relationship between quantitative ratios in the process of qualitative transfer of genetic information and appearance, i.e. the characteristics of organisms.

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.

Step 1 (From 1 to 306)

$$R1 = 10; R2 = 22; R3 = 19; ... R306 = 17; \\ [R1 + (R1+R2) + (R1+R2+R3)..., + (R1+R2+R3..., + R306)] = S; \\ R1 = 10 \text{ atoms}; \\ (R1+R2) = (10+22) = 32; \\ (R1+R2+R3) = (10+22+19) = 51; \\ (R1+R2+R3..., + R306) = 5640 \text{ atoms}; \\ R1,2,3,n = \text{Number of atoms in amino acids } 1,2,3,n \\ [R1 + (R1+R2) + (R1+R2+R3)..., + (R1+R2+R3..., + R306)] = \\ = (10+32+51+70..., + 5640) = S; \\ S = 863 \ 208; \\ Step 2 \text{ (From 306 to 1)} \\ R306 = 17; R305 = 24; R304 = 17; ... R1 = 10 \text{ atoms}; \\ [R306 + (R306+R305) + (R306+R305+R304)..., + (R306+R305+R304..., + R1)] = S1; \\ R306 = 17;$$

(R306+R305) = (17+24) = 41;(R306+R305+R304) = (17+24+17) = 58;(R306+R305+R304..., +R1) = (17+24+17..., +10) = 5640;

R1,2,3,n = Number of atoms in amino acids 1,2,3,n

$$[R306 + (R306 + R305) + (R306 + R305 + R304)..., + (R306 + R305 + R304..., + R1)] =$$

$$= (17 + 41 + 58 + 75..., + 5640) = S1;$$

$$S1 = 868 \ 272;$$

From 1 to 102 and from 102 to 1

	G	I	٧	Ε	Q	С	С			Т	SUM
	10	22	19	19	20	14	14	•	•	17	1880
	1	2	3	4	5	6	7			102	5253
Step 1	10	32	51	70	90	104	118			1880	95976
Step 2	5640	5630	5608	5589	5570	5550	5536			3777	481184
											577160

$$(0+10) = 10$$
; $(10+22)=32$; $(10+22+19)=51$; etc.
 $(17+24+17..., +10) = 5640$; $(17+24+17..., +22) = 5630$; etc.
 $(95976 + 481184) = 577160$:

From 103 to 204 and from 204 to 103

	G	Ī	٧	Ε	Q	С	С		Т	SUM
	10	22	19	19	20	14	14		17	1880
	103	104	105	106	107	108	109		204	15657
Step 1	1890	1912	1931	1950	1970	1984	1998		3760	287736
Step 2	3760	3750	3728	3709	3690	3670	3656	•	1897	289424
										577160

(287736 + 289424) = 577160;

From 205 to 306 and from 306 to 205

	G	- 1	٧	Е	Q	С	С		Т	SUM
	10	22	19	19	20	14	14		17	1880
	205	206	207	208	209	210	211		306	26061
Step 1	3770	3792	3811	3830	3850	3864	3878		5640	479496
Step 2	1880	1870	1848	1829	1810	1790	1776		17	97664
										577160

$$(479496 + 97664) = 577160;$$

From 1 to 102	F	From 103 to 204	From 205 to 306
•		4	\
577160		577160	577160
Ä		+	Ľ
		1.731.480	

$$1.731.480 = (5640 + 5640 + 5640... + 5640);$$

Number of atoms in Insulin = 5640;

Schematic representation of the bio intervals from 1 to 306 we will show in the fig.1 and 2.

In that group of chains there are three groups with 102 amino acids. Each of these three groups of amino acids has an identical number of atoms.

It can be concluded that there is a connection between quantitative characteristics in the process of

transfer of genetic information and the qualitative appearance of given genetic processes.

BIO INTERVALS - AMINO ACIDS FROM 1 TO 306

Within the digital pictures in biochemistry, the physical and chemical parameters are in a strict compliance with programmatic, cybernetic information principles. As an example, we will here give you the mathematical gravity forces. These forces determine the positioning of aminoacids in their molecules. 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.

Insulin Bio Intervals(1)

Step 1

Number of	Bio		Number of	Bio		Number of atoms	Bio	
atoms	intervals	Rank	atoms	intervals	Rank	or alorno	intervals	Rank
199			1139			2079		
From 1			From 1			From 1		
to 12	199	78	to 63	199	2016	to 114	199	6555
323			1263			2203		
From 1			From 1			From 1		
to 18	124	171	to 69	124	2415	to 120	124	7260
361			1301			2241		
From 1			From 1			From 1		
to 20	38	210	to 71	38	2556	to 122	38	7503
579			1519			2459		
From 1			From 1			From 1		
to 32	218	528	to 83	218	3486	to 134	218	9045
617			1557			2497		
From 1			From 1			From 1		
to 34	38	595	to 85	38	3655	to 136	38	9316
741			1681			2621		
From 1			From 1			From 1		
to 41	124	861	to 92	124	4278	to 143	124	10296
940			1880			2820		
From 1			From 1			From 1		
to 51	199	1326	to 102	199	5253	to 153	199	11781
3760	940		10340	940		16920	940	

```
(199-0) = 199; (323-199) = 124; (361-323) = 38; (741-617) = 124;
     (940-741) = 199; (1139-940) = 199; (1263-1139) = 124;
                               etc.
               (Chain A + Chain B) = 940 atoms,
               (Chain C + Chain D) = 940 atoms,
                               etc.
               (3760 + 16920) = (10340 + 10340);
               10340 = (940 + 940 + 940... + 940);
                  3760 = (940 + 940 + 940 + 940)
               10340 = (940 + 940 + 940..., + 940)
               16920 = (940 + 940 + 940..., + 940)
                   (2079-1139) = (1139-199);
                   (2203-1263) = (1263-323);
```

(2241-1301) = (1031-361);etc.

Correlation of bio intervals

Number		Number		Number	
of	Bio	of	Bio	of atoms	Bio
atoms	intervals	atoms	Intervals		Intervals
199		1139		2079	
	199		199		199
	Γ/	100 0070	N . O1 44	00.	

[(199 + 2079) : 2] = 1139;

Number		Number		Number	
of	Bio	of	Bio	of atoms	Bio
atoms	intervals	atoms	Intervals		Intervals
323		1263		2203	
	124		124		124

[(323 + 2203) : 2] = 1263;

Number		Number		Number	
of	Bio	of	Bio	of atoms	Bio
atoms	intervals	atoms	Intervals		Intervals
361		1301		2241	
	38		38		38
		061 004	1). 0] 10/	11.	

[(361 + 2241): 2] = 1301;

Number		Number		Number	
of	Bio	of	Bio	of atoms	Bio
atoms	intervals	atoms	Intervals		Intervals
579		1519		2459	
	218		218		218

[(579 + 2459) : 2] = 1519;

Number of	Bio	Number of	Bio	Number of atoms	Bio
atoms	intervals	atoms	Intervals		Intervals
617		1557		2497	
	38		38		38

[(617 + 2497) : 2] = 1557;

Number of atoms	Bio intervals	Number of atoms	Bio Intervals	Number of atoms	Bio Intervals
741	124	1681	124	2621	124

 $[(\overline{741} + 2621) : 2] = 1681;$

Number of atoms	Bio intervals	Number of atoms	Bio Intervals	Number of atoms	Bio Intervals
940	199	1880	199	2820	199

[(940 + 2820) : 2] = 1880;

Insulin Bio Intervals(2)

	Number			Number			Number		
	of	Bio		of	Bio		of atoms	Bio	
	atoms	intervals	Rank	atoms	intervals	Rank		intervals	Rank
	3019			3959			4899		
	From 1			From 1			From 1		
	to 165	199	13695	to 216	199	23436	to 267	199	35778
	3143			4083			5023		
	From 1			From 1			From 1		
	to 171	124	14706	to 222	124	24753	to 273	124	37401
	3181			4121			5061		
	From 1			From 1			From 1		
	to 173	38	15051	to 224	38	25200	to 275	38	37950
	3399			4339			5279		
	From 1			From 1			From 1		
	to 185	218	17205	to 236	218	27966	to 287	218	41328
	3437			4377			5317		
	From 1			From 1			From 1		
	to 187	38	17578	to 238	38	28441	to 289	38	41905
	3561			4501			5441		
	From 1			From 1			From 1		
	to 194	124	18915	to 245	124	30135	to 296	124	43956
	3760			4700			5640		
	From 1			From 1			From 1		
	to 204	199	20910	to 255	199	32640	to 306	199	46971
ı	23500	940		30080	940	3—3. 0	36660	940	
		3 7 0			3 7 0			9 7 0	

(3019-2820) = 199; (3143-3019) = 124; (3181-3143) = 38; etc.

Schematic representation of the bio intervals from 199 to 5640 we will show in the fig.3.

Correlation of bio intervals

Number of	Number of	Number of atoms
atoms	atoms	
3019	3959	4899

 $(3019 + 4899) = (3959 \times 2);$

Number of	Number of	Number of atoms
atoms	atoms	
3143	4083	5023

 $(3143 + 5023) = (4083 \times 2);$

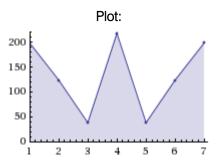
Number	Number	Number of atoms
atoms	atoms	or atoms
3181	4121	5061

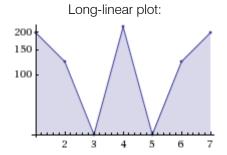
 $(3181 + 5061) = (4121 \times 2)$; etc.

The molecule of insulin we can understand as words built from letters, i.e. aminoacids. The meaning of words is determined by positioning of letters. Each of these words has its biochemical base. If this base is expressed by corresponding discrete numbers, we find out that the base has its own program, cybernetic and information characteristics. In fact, we will find out that the sequencing of the molecule is conditioned and determined not only by biochemical, but also by cybernetic and information principles.

BIO INTERVALS(3)

Mathematica plaintext input:





ListLogPlot[{199, 124, 38, 218, 38, 124, 199}, Joined -> True, Mesh -> All, Filling -> Axis]

BIO INTERVAL - AMINO ACIDS-FROM 306 TO 1.

Step 2

Number of	Bio		Numbe of	er	Bio		Numb of aton		Bio	
atoms	intervals	Rank	atoms		intervals	Rank			Intervals	Rank
199 From 306 to 297	199	3015	1139 From 306 to 246	0	199	16836	2079 From 306 195	to	199	28056
323 From 306 to 290	124	5066	239	0	124	18530	2203 From 306 188	to	124	29393
361 From 306 to 288	38	5643	237	0	38	19005	2241 From 306 186	to	38	29766
579 From 306 to 276	218	9021	1519 From 306 to 225	0	218	21771	2459 From 306 174	to	218	31920
617 From 306 to 274	38	9570	1557 From 306 to 223	0	38	22218	2497 From 306 172	to	38	32265
741 From 306 to 268	124	11193	1681 From 306 to 217	0	124	23535	2621 From 306 166	to	124	33276
940 From 306 to 256	199	14331	205	0	199	26061	2820 From 306 154	to	199	35190
3760	940		10340		940		16920		940	

[(199 + 2079) : 2] = 1139; [(323 + 2203) : 2] = 1263; etc.

Schematic representation of the bio intervals from 306 to 1 we will show in the fig.4.

Step 2 (323-199) = 124; (361-323) = 38; (579-361) = 218; etc.

Connection:

(36660 and 3760) > 36660 3760 = (940 x Y1); (3760 and 36660) > 3760 36660 = (940 x Y2); (30080 and 10340) > 30080 10340 = (940 x Y3); (10340 30080) > 10340 30080 = (940 x Y4); (23500 and 16920) > 23500 16920 = (940 x Y5); (16920 and 23500) > 16920 23500 = (940 x Y5);

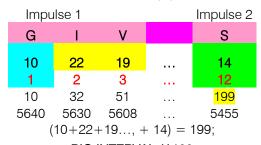
 $(36660 + 3760) + (30080 + 10340) + (23500 + 16920) = (940 \times Y);$ Y = 43;

In those examples there is an exact mathematical balance of groups of aminoacids from 1 to 102, 103 to 204, 205 to 306 and 199 to 5640.. This balance is one of important quantitative characteristics of all processes in biochemistry. How functioning of biochemistry is determined through cybernetic information principles, will be discussed further in Table 1.

BIO INTERVALS(4)

The result of the research that we have carried out clearly shows that there is a matrix code in insulin. It also shows that the coding system within the amino acidic language gives a full information, not only for the amino acid "record", but also for its structure, configuration and its various shapes. In the following text we shall discuss the issue of the existence of the insulin code, and also the issue of coding of individual structural levels in this protein.

BIO INTERVAL (+)199

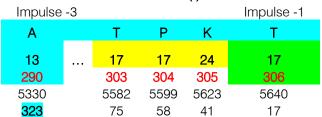


BIO INTERVAL (-)199



BIO INTERVAL (+)323 Impulse 1 Impulse 3 Ν G ٧ 10 22 19 17 2 1 3 18 10 32 323 51 5640 5630 5608 5334

BIO INTERVAL (-)323



BIO INTERVAL (+)361

Impu	ılse 1		Impulse 4
G	- 1	٧	С
10	22	19	 14 20
1	2	3	20
10	32	51	361
5640	5630	5608	5293

BIO INTERVAL (-)361

Impulse	:-4				Impulse –1
V		Т	Р	K	Т
19 288		<mark>17</mark> 303	17 304	24 305	17 306
5298		5582	5599	5623	5640
361		75	58	41	17
			etc.		

The molecule of insulin we can understand as words built from letters, i.e. aminoacids. The meaning of words is determined by positioning of letters. Each of these words has its biochemical base. If this base is expressed by corresponding discrete numbers, we find out that the base has its own program, cybernetic and information characteristics. In fact, we will find out that the sequencing of the molecule is conditioned and determined not only by biochemical, but also by cybernetic and information principles.

BIO INTERVAL – AMINO ACIDS-FROM 1 to 306 and 306 TO 1.

Bio intervals of insulin we will show in the table 1 and table 2

Step 1

Table 1: Bio intervals (from 1 to 306)

199	323	361	579	617	741	940
1139	1263	1301	1519	1557	1681	1880
2079	2203	2241	2459	2497	2621	2820
3019	3143	3181	3399	3437	3561	3760
3959	4083	4121	4339	4377	4501	4700
4899	5023	5061	5279	5317	5441	5640
(4.40)		/00-		(004	0 00 70	

(1139 - 199) = (2079 - 1139) = (3019 - 2079); etc.

Step 2

Table 2: Bio intervals (from 306 to 1)

940	741	617	579	361	323	199
1880	1681	1557	1519	1301	1263	1139
2820	2621	2497	2459	2241	2203	2079
3760	3561	3437	3399	3181	3143	3019
4700	4501	4377	4339	4121	4083	3959
5640	5441	5317	5279	5061	5023	4899
				\		

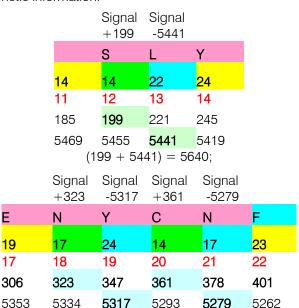
(1880-940) = 2820-1880; etc.

These tables contains an overview of all bio intervals. The bio intervals show some of the quantitative characteristics of the molecule of insulin.

Schematic representation of the bio intervals we will show in the fig.5, 6 and 7.

The Insulin Bio Signal Code

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.



557

579

4741

4717

5103 5083 5061 5042 5023 5010 (579+5061) = (617+5023)(617-579) = (5061-5023)

Signal Signal

617

630

652

598

	+741	-4899				
С	G	Е	R			
14	10	19	26			
40	41	42	43			
731	741	760	786			
4923	4909	4899	4880			
(7	(741 + 4899) = 5640;					
	Signal	Signal				
	+940	-4700				
K	+940 T	-4700 G	I			
K 24			I 22			
	T	G	I <mark>22</mark> 53			

Aforementioned bio-codes were calculated using of corresponding groups of aminoacids. These are unions with different number of aminoacids. There are different ways and methods of selecting these unions of amino acids. We hope that science will determine which method is most efficient for this selection. Some signals have a positive, and some have negative numeric value.

(4700 + 940) = 5640; etc.

4700

4690

DISCUSSION IV.

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 factor ³, some used the values derived from the cellular automata 4-7, some used hydrophobic and/or hydrophilic values 8-16, some were through Fourier transform ^{17,18}, and some used the physicochemical distance ¹⁹. The author [34-40] is devoted to provide a digital code for each of 20 native amino acids. These digital codes should more complete and better reflect the essence of each of the 20 amino acids. Therefore, it might stimulate a series of future work by using the author's digital codes to formulate the pseudo amino acid composition for predicting protein structure class [20-22], subcellular location [23, 24], membrane protein type [9, 25], enzyme family class [26, 27], GPCR type [28, 29], protease type [30], protein-protein interaction [31], metabolic pathways [32], protein quaternary structure [33], and other protein attributes. 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), stereo-chemical and digital structure of proteins.

References Références Referencias

- Gene Cloning & K.C. Chou, Expression Technologies, Chapter 4 (Weinrer, P.W., and Lu, Q., Eds.), Eaton Publishing, Westborough, MA (2002), pp. 57-70.
- K.C. Chou, Prediction of protein cellular attributes using pseudo amino acid composition PROTEINS: Structure, Function, and Genetics (Erratum: ibid., 2001, Vol.44,60) 43 (2001) 246-255.
- X. Xiao, S. Shao, Y. Ding, Z. Huang, Y. Huang, K. C. Chou, Using complexity measure factor to predict protein subcellular location, Amino Acids 28 (2005) 57-61.
- X. Xiao, S. Shao, Y. Ding, Z. Huang, X. Chen, K. C. Chou, Using cellular automata to generate Image representation for biological sequences, Amino Acids 28 (2005) 29-35.
- X. Xiao, S. Shao, Y. Ding, Z. Huang, X. Chen, K. C. Chou, An Application of Gene Comparative Image for Predicting the Effect on Replication Ratio by HBV Virus Gene Missense Mutation, Journal of Theoretical Biology 235 (2005) 555-565.
- X. Xiao, S. H. Shao, Z. D. Huang, K. C. Chou, Using pseudo amino acid composition to predict protein structural classes: approached with complexity measure factor, Journal of Computational Chemistry 27 (2006) 478-482.

- 7. X. Xiao, S. H. Shao, Y. S. Ding, Z. D. Huang, K. C. Chou, Using cellular automata images and pseudo amino acid composition to predict protein sub-cellular location, Amino Acids 30 (2006) 49-54.
- 8. K. C. Chou, Using amphiphilic pseudo amino acid composition to predict enzyme subfamily classes, Bioinformatics 21 (2005) 10-19.
- 9. K. C. Chou, Y. D. Cai, Prediction of membrane protein types by incorporating amphipathic effects, Journal of Chemical Information and Modeling 45 (2005) 407-413.
- Z. P. Feng, Prediction of the subcellular location of prokaryotic proteins based on a new representation of the amino acid composition, Biopolymers 58 (2001) 491-499.
- 11. Z. P. Feng, An overview on predicting the subcellular location of a protein, In Silico Biol 2 (2002) 291-303.
- 12. M. Wang, J. Yang, Z. J. Xu, K. C. Chou, SLLE for predicting membrane protein types, Journal of Theoretical Biology 232 (2005) 7-15.
- S. Q. Wang, J. Yang, K. C. Chou, Using stacked generalization to predict membrane protein types based on pseudo amino acid composition, Journal of Theoretical Biology, in press (2006) doi:10.1016/j.jtbi.2006.1005.1006.
- M. Wang, J. Yang, G. P. Liu, Z. J. Xu, K. C. Chou, Weighted-support vector machines for predicting membrane protein types based on pseudo amino acid composition, Protein Engineering, Design and Selection 17 (2004) 509-516.
- 15. S. W. Zhang, Q. Pan, H. C. Zhang, Z. C. Shao, J. Y. Shi, Prediction protein homo-oligomer types by pseudo amino acid composition: Approached with an improved feature extraction and naive Bayes feature fusion, Amino Acids 30 (2006) 461-468.
- Y. Gao, S. H. Shao, X. Xiao, Y. S. Ding, Y. S. Huang, Z. D. Huang, K. C. Chou, Using pseudo amino acid composition to predict protein subcellular location: approached with Lyapunov index, Bessel function and Chebyshev filter, Amino Acids 28 (2005) 373-376.
- Y. Z. Guo, M. Li, M. Lu, Z. Wen, K. Wang, G. Li, J. Wu, Classifying G protein-coupled receptors and nuclear receptors based on protein power spectrum from fast Fourier transform, Amino Acids 30 (2006) 397-402.
- 18. H. Liu, M. Wang, K. C. Chou, Low-frequency Fourier spectrum for predicting membrane protein types, Biochem Biophys Res Commun 336 (2005) 737-739.
- K. C. Chou, Prediction of protein subcellular locations by incorporating quasi- sequence-order effect, Biochemical & Biophysical Research Communications 278 (2000) 477-483.
- 20. K. C. Chou, A novel approach to predicting protein structural classes in a (20-1)-D amino acid

- composition space, Proteins: Structure, Function & Genetics 21 (1995) 319-344.
- 21. K. C. Chou, C. T. Zhang, Predicting protein folding types by distance functions that make allowances for amino acid interactions, Journal of Biological Chemistry 269 (1994) 22014-22020.
- 22. K. C. Chou, C. T. Zhang, Review: Prediction of protein structural classes, Critical Reviews in Biochemistry and Molecular Biology 30 (1995) 275-349.
- 23. K. C. Chou, D. W. Elrod, Protein subcellular location prediction, Protein Engineering 12 (1999) 107-118.
- 24. K. C. Chou, Review: Prediction of protein structural classes and subcellular locations, Current Protein and Peptide Science 1 (2000) 171-208.
- 25. K. C. Chou, D. W. Elrod, Prediction of membrane protein types and subcellular locations, PROTEINS: Structure, Function and Genetics 34 (1999) 137-153.
- 26. K. C. Chou, D. W. Elrod, Prediction of enzyme family classes, Journal of Proteome Research 2 (2003) 183-190.
- 27. K. C. Chou, Y. D. Cai, Predicting enzyme family class in a hybridization space, Protein Science 13 (2004) 2857-2863.
- 28. K. C. Chou, D. W. Elrod, Bioinformatical analysis of G-protein-coupled receptors, Journal of Proteome Research 1 (2002) 429-433.
- 29. K. C. Chou, Prediction of G-protein-coupled receptor classes, Journal of Proteome Research 4 (2005) 1413-1418.
- 30. K. C. Chou, Y. D. Cai, Prediction of protease types in a hybridization space, Biochem. Biophys. Res. Comm. 339 (2006) 1015-1020.
- 31. K. C. Chou, Y. D. Cai, Predicting protein-protein interactions from sequences in a hybridization space, Journal of Proteome Research 5 (2006) 316-322.
- 32. K. C. Chou, Y. D. Cai, W. Z. Zhong, Predicting networking couples for metabolic pathways of Arabidopsis, EXCLI Journal 5 (2006) 55-65.
- 33. K. C. Chou, Y. D. Cai, Predicting protein quaternary structure by pseudo amino acid composition, PROTEINS: Structure, Function, and Genetics 53 (2003) 282-289.
- 34. L.Kurić, The digital language of amino acids. Amino Acids (2007) 653-661.
- 35. L.Kurić, The Atomic Genetic Code. J. Comput Sci Biol 2 (2009) 101-116.
- 36. L.Kurić, Mesure complexe des caracteristiques dynamiques de series temporelles "Journal de la Societe de statistique de Paris" tome 127, No. 2.1986.
- 37. L.Kurić, The Insulin Bio Code Zero Frenquencies, GJMR Vol. 10 Issue 1: 15 May 2010.
- 38. L.Kurić, Molecular biocoding of insulin, Advances

- and Applications in Bioinformatics and Chemistry, Jul. 2010.p.45 - 58.
- 39. L.Kurić, The Insulin Bio Code Prima sequences, GJMR Vol. 1 Issue 1: 15 June 2010.
- 40. L.Kurić, ATOMIC HEMOGLOBIN CODE, GJMR Volume 10 Issue 2, October 2010.
- 41. L.Kurić, Language of Insulin Decoded:Discret code 1128, IJPBS JOURNAL, October 2010.
- 42. L.Kurić, "Measures of Bio Insulin Frequencies", IJCSET (Volume 1. Issue 4. December, 2010)
- 43. L.Kurić, The Insulin Bio Code Standard Deviation, International Journal of Scientific and Engineering Research (IJSER) Nov 25, 2010 under ISSN 2229-5518.
- 44. L.Kurić, Molecular biocoding of insulin amino acid Gly, International Journal of Scientific and Engineering Research (IJSER) - March 2011 issue.
- 45. L.Kurić, Algorithm and computational complexity of International Journal of Computer Technology and Application (IJCTA) - Feb 2011 issue.
- 46. L.Kurić, Algorithm and computational complexity of Insulin - amino acid Asn, International Journal of Computer Technology and Application (IJCTA).

Awards

Gold medal for Bosnia and Herzegovina, in the name of your people and Bosnia and Herzegovina. Such individuals deserve recognition not only within their country of origin, but also worldwide. As such, the American Biographical Institute - a highly esteemed leader in the research of upstanding individuals around the globe - has selected you to receive one of its most internationally prominent honors, the GOLD MEDAL FOR BOSNIA AND HERZEGOVINA. -International health professional of the year 2010., -Man of the year in medicine and healthare designation for 2010.,-The international Hipocrates awards for medical achievement, -Cambridge certificate for outstanding medical achievement for 2011. - International health professional of the year 2012.-Cambridge certificate for Outstanding Medical Achievement, 2012., -ABI-Man of the year 2012. - Nomination for Great Minds of the 21st Century, a major reference directory including just 1.000 of the world's top thinkers and intellectuals. My contributions to the field of medicine and healthcare have waranted the high regard of nomination for Great Minds of the 21st Century.