The Genetic code: Section III

Tranformations between number-base systems

Åsa Wohlin <u>www.u5d.net</u> 2015-03-23

CONTENT

Page

17.	Transformations beteen number-base syystems					79
18.	More on totals and other notable transformations					89
19.	P – phosphorous groups – Coenzymes – Nucleotides					S
	Met AUG	G - tRNA-en	d ACC		97	
20.	Additions	s to files 17-1	18	103		
21.	I. The tri	plet series. –	- II. An a	lterna	ative series	108
22	Other sul	bstances	113			
Disc	cussion	116				
Ref	erences	118				

17. Transformations between number-base systems (nb-x)

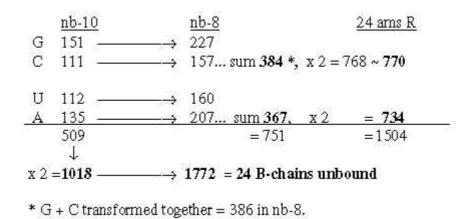
Bases - totals of ams - 5 x ES- numbers - Generative production of the 12-groups

1. Transformations of the codon-bases to the 12-groups of ams:

1.1 All geometrical dimensions should naturally be regarded as present in the cell simultaneously, on different levels, and interdependent through transformations into one another. One simple example is the geometries of proteins, forming linear threads (\sim D1), sheets (\sim D2) and globular forms (\sim D3).

The thought that different d-degrees could be associated with different number base systems (nb-x), as nb-10, nb-8, nb-6 for x = 5, 4, 3, led to a first test on mass of codon bases with remarkable results, figure 17-1 below. Further investigation showed also several connections with the ES-series. (Nb-x in text below often written as "-index figures. Figures in nb-8 and nb-6 are often rewritten with figures from nb-10.)

Fig 17-1: From mass of codon bases to the two 12-groups of ams:



Hence, 4 sets of the 4 bases give the total sum of 24 unbound ams.

We find also that 2 x G+C-bases in nb-8 as 768 gives total sum 3276 in nb-6:

 $\frac{\text{nb-10}}{768 \longrightarrow 3276} \quad 24 \text{ ams } R + B \text{, unbound (rewritten from 3320)}$ The sum of the 4 bases in nb-8 = 752 -/+1: $\frac{\text{nb-10}}{752 \longrightarrow 2848} \quad 24 \text{ ams } R + B \text{, bound (rewritten from 3050)}$

Fig 17-2. From 752 as sum of ES-numbers 5', 4' and 3' to 2848 in nb-6:

5' $292_{10} \rightarrow 1204_{6}$, 244 = 960 U+A 4' $252_{10} > 1100_{6}$, +244 = 1344, 24 B chains bound <u>3' $208_{10} \rightarrow 544_{6} = 544$. G C</u> **4** = 2848 1' $100_{10} \rightarrow 244_{6}$

2848 = 24 ams R + B bound

1.2 Some general annotations:

However strange the idea surely may seem for scientific "common sense", the many astonishing results here and below are rather difficult to dismiss as only haphazard. If they are not, if they reveal some connections on deep energy levels, they should represent one kind of references, one kind of guiding operators for potential growth - or just what is sometimes in the biochemical field is referred to as "affinities"?

All derived numbers shall naturally be regarded as nb-10-numbers, hence transformations as nb-10 \rightarrow > nb-8 may be repeated, illustrated for instance in the carbon-nitrogen cycle in the sun, from ¹²C to ¹⁴N to ¹⁶O, intermediate steps showing one way to perform such transformations.

It follows that all operations as multiplications are performed in nb-10. Indexes for x in nb-x are often used below to shorten the text. As mentioned above numbers in nb-8 and nb-6 are often rewritten with figures from nb-10.

A question is of course if such rewritings could be expressed in biochemical processes as for instance 20 equivalent with (\sim) 18 in nb-8 as -2H or 120 in nb-6 \sim 76 as - 44 (CO2)?

Another question is how to interpret nb-16 in many examples below If keeping to the thought of x in nb-x as first three numbers in the elementary chain $5' \rightarrow 4' \rightarrow 3'$ doubled, should nb-16 be regarded as 2 x 4 doubled or 2(5 + 3) doubled?

Fig Ti-1

A general feature may be noted: transformation of sums or whole units give larger numbers in lower nb-systems than their parts transformed and summed afterwards.

1.3 Halves of the 12-groups 770 and 734, -/+1 = 384 and 368:

Fatty acids, a first annotation here:

Cell membranes are an equally essential part of life as the genetic code. Two of the most common fatty acids give transformed to nb-6 three times these numbers 367 and 385, +/-1, a relation to R-chains of the 24 ams = 3/2 and simultaneously a relation d-degree 3 to 4(nb-6 to nb-8) with the assumed view above.

C16H32O2: 256-10 \rightarrow 1104-6 = 3 x 368 C18H36O2: 284-10 \rightarrow 1152-6 = 3 x 384, (Note: 1152 = 752 rewritten) Cf. the hexagonal pattern in Table 0: fatty acids as a third way to read such a pattern.

From the numbers **384** and **368** in nb-10 transformed in two steps to nb-8 we get 2 sets of bases G and A in nb-8, as in opposite direction to the figure above and without C and U:

 $384 \times \frac{1}{2} = 192-10 \rightarrow 300-8 / 300-10 \rightarrow 454-8 = 2 \times 227 = 2 \text{ G-8}$ $368 \times \frac{1}{2} = 184-10 \rightarrow 268-8 / 268-10 \rightarrow 414-8 = 2 \times 207 = 2 \text{ A-8}$

1.4 . Bases \rightarrow totals:

1.4.1 Four times G+U and A+C to ~ B- and R-chains of total 3276:

Sums of R+B-chains together in **nb-10**:

G1 + U1 = C2 + A2 = 1468C1 + A1 = G2 + U2 = 1808... Sums of coded amino acids (R + B)

With exchanged partners these sums are given from 4 times the bases:

Fig. 17-3

	10-bas	38	8-base
4 G-bases = $4 x$	151 = 604	\rightarrow	1134
4 C-bases = 4 x	1 1 1 = 4 44	\rightarrow	674sum 1808
1 A-base	= 135	\rightarrow	207, x 4= 828
<u>1 U-base</u>	= 112	\rightarrow	160, x 4= 640 sum 1468

In nb-10 we have groups of ams paired in keto-/amino types: Here G- and A-bases have exchanged partners and bases A and U must be multiplied with 4 after transformation.

Fig. 17-4

Rewriting 640 to 638 and 828 to 830 gives the right sums B 1772 and R 1504.

Fig. 17-5

		10-bas	se	8-ba:	se		
	"5";	292	\rightarrow	444	-92		
					>	818,	$\sim 1018 = 2 \ge 4 \text{ RNA-bases} 509$
	ⁿ 4 ⁿ	252	\rightarrow	374			in base-10 system
		16-base		10-base	100		
2 x	"3"	416 -	→>	1046	55		1046 = 2 x 4 DNA-bases 523 in base-10 system

Number 416 (2 x 3', 208) is the one which added to 544 gives the A-U-group of ams. Cf. that U-base gets replaced by T-base in DNA, a CH2-group added for inward direction to DNA. (It could perhaps be compared with the interpretation of nb-16 as 2 x (3' + 5'), a step backwards from 3' to 5', equivalent with inwards?

2. The bases in the ES-chain:

Fig. 17-6

<u>10-base</u>	8-base
G 151>	227 (denotation here G _g etc.)
U 112 ->	160
C 111 ->	157sum 544
	<u>10-base</u> <u>8-base</u>
A 135>	207 sum 208 ·1 $A_8 \frac{1}{207} \longrightarrow \frac{1}{317} = U_8 + C_8 (2 \times 158, 5)$

U 160 + C 157 in nb-8 approximate number 2' = 159 in the ES-series, together 317.

In nb-10 number 385 is the interval 544 to 159. Here G-8 becomes the same interval to both bases U-8 + C-8. Cf. that G-base can bind to both:

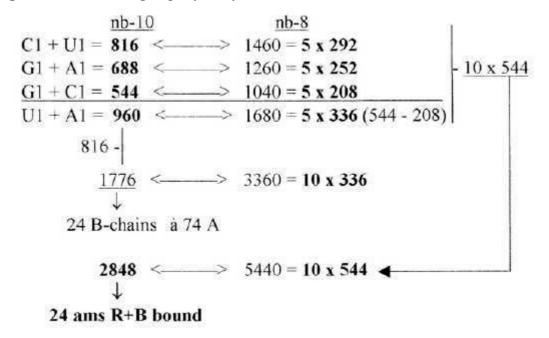
Fig 17-7: The bases in nb-8 in the ES-chain:

These relations could be a reason why G+C-bases get connected with the 12-group 770 of ams in spite of all bases equally represented in this group.

3. 5 times ES-numbers:

3.1 The transformations between nb-10 and nb-8 of main codon groups of ams and 5 times the ES-chain numbers 5' - 4' - 3' are among the most astonishing:

Fig 17-8: Main codon groups of ams from 5 times ES-numbers:



816 and 688 is the division of **R**-chains of total sum 1504 of 24 ams, a division between purine and pyrimidine codon groups, As a division in step 5 - 4 here it precedes the one between complementary pairs G-C and U-A, which are attained from the secondary division of 544 in 336 and 208, a division in step 4 - 3.

Note also about 1344, the 24 B-chains bound, included in sum 2848:

1344 in nb-10 = 2500 in nb-8 = ES-numbers 5(292 + 208)

These relations seem to support the relevance of both the ES-chain and the thought that nb-transformations could be part of the reference system.

Fig 17-9 *5 x half of 752, number 688 as an interval:*

	10-base		8-base
G+C, x 2 = 544x 2;	1088	<	$1880 = 3360 - 1460 = 5 \times 376$
G1+A1 R;	l 688 I	>	1260, 5 x 252
	1776	<	3360

There is also the feature that divisions stepwise as polarizations of numbers 816 in U1 + C1, separately transformed to nb-8 give 1260, next lower level, and this back to nb-10 and divided G1 and A1 gives 1040 in nb-8:

Fig 17-10. *Stepwise polarization giving next number x5 in Es-chain:*

Steps of "polarisations" "5" -> "4" -> "3":

Step	<u>s: Ams</u>	<u>nb-10</u>	<u>nb-8</u>	ES-numbers
"5"	$\underline{C1 + U1}$:	816 <	1460	= 5 x 292
	C1: U1:	$\sqrt{\frac{353}{463}}$ — 3	> 541 7171258	$\sim 1260 = 5 \times 252$
"4"	$\frac{G1+A1}{G1}$	688 <	1260 <	- 1260
	G1: A1:	\bigvee_{497}^{191}	$> \frac{277}{7611038}$	$\sim 1040 = 5 \ge 208$
"3"	G1+C1:	544 <	1040 <	- 1040

3.2 About the interval 84 = $292 \rightarrow 208$ in the ES-chain we have that n x 84 (n = 1, 2, 4) times 10 (1040 ~ 840, 1680 and 3360) in nb-8 gives the groups 544, 960 and 1776 in nb-10:.

Fig 17-11. *n x interval 84:*

84 = interval 292 - 208: ES-series nb-8 nb-10 <--- 336 x 10 24 B-chains à 74 A: 1776 =**4 x 84**, x 10 (292 - 208) <--- 168 x 10 A+U-coded ams R: 960 $= 2 \times 84$ " <--- 84 x 10 " G+C-coded ams R: 544 $= 1 \times 84$

3.3 5 times intervals in the exponent series in nb-8 give ams-groups -/+1:

Fig. 17-12 *5 x interval in the ES-chain:*

Ams	10-base	8-base	Intervals in the exponent series:
G1+1	192 <	-300 = 5 x interva	$1 \ 60 = 292 - 352 = "5" - ("4 + 1")$
A1 -1;	496	760 = 5 x interva	l152 = 252 - 100 = "4" "1"
U1+1	464	720 = 5 x interva	1144 = 352 - 208 = ("4 + 1") - "3"
C 1 - 1	352	540 = 5 x interva	1108 = 208 - 100 = "3 "1"

3.4 Nb-6: 5 times the ES-numbers 5', 4' 3' in nb-6:

It gives the sum of U- plus A-coded ams R and also all C-atoms in R-chains in nb-10, divided on G1 + A1 = 396 and U1 + C1 = 564:

5	J times		5 111 110 0 10 570 5	01.
	10-base		<u>6-base</u>	
396	396	<	$1460 = 5 \ge 292$	a)
	324	<	$1260 = 5 \ge 252$	
564 <			>	b)
	240	< <u> </u>	$1040 = 5 \times 208$	- 52
	960	×2	752 x 5	
↓ =	A + U			
396 =	292+10)4 396	+ 101 = 497 = A1	a)
564 =	460 +10	<u>)4</u> 564	-101 = 463 = U1	b)
Σ:	752 +20)8		:

Fig..17-13 5 times ES-numbers in nb-6 to 396-564.

3.5 The parts above of 960 in nb-16 gives the total mass of bound ams in nb-8:

Fig 17-14

		16-base	10-base	8-base	
G1,A1	- G2, A2	396	918	1626	
G1,A1 -	C2, U2	292	658 _	1222	
		688		2848 ->	2848 = 24 ams bound
But	2 x	344 -	>	1504 x 2 =	= 2 x 24 ams R

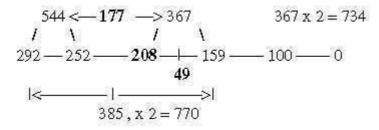
3.6 The numbers U1 + C1 = 816 and G1 + A1 = 688 read in nb-8 rewritten, give in two steps total 24 ams, R + B inbound in nb-8:

<u>nb-8</u> 816 ~ 1016 688 ~ 710 ... Sum 1726-8./ 1726-10 → **3276**-8

4. Generation of the two 12-groups of ams with mixed and non-mixed codons:

4.1 Generative production of sums within 12-groups of ams:

Fig. 17-15a. The ES-chain, numbers 177 and 208:



Cf Table 2,3 in file <u>0</u>2.

Numbers 770 and 734 generated from 177 and 208:

- From 177 we get 385 in two steps nb-10 to nb-8:

- From 208 we get 734 in three such steps:

Fig. 17-15b

 $\begin{array}{c} \underline{\text{nb-10} \rightarrow 8} & \underline{\text{nb-10} \rightarrow 8} \\ 177 \longrightarrow 261 \longrightarrow 385 & \rightarrow x \ 2 = 770 \\ |< \underline{\qquad 208 \longrightarrow }| \end{array}$

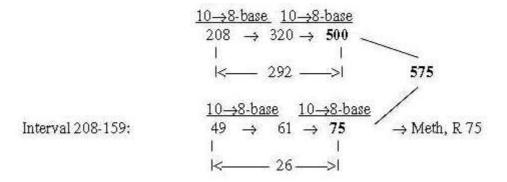
* 318 = 2 z 2', 159, from there only two steps:

Fig 17-15c:

$$\begin{array}{rcl} & \underline{10 \rightarrow 8\text{-base}} \\ 2 \ x \ 159; & \mathbf{318} \ \rightarrow \ 476 \ \rightarrow \ \mathbf{734} \ \rightarrow \ \mathbf{RNA+Pair-coded} \ \mathrm{ams} \end{array}$$

In group 734 U+A-coded ams = 575, a number given through two steps nb-10 to nb-8, either as sum of 500 + Meth 75 or from 208 + interval 49: Meth that starts the protein synthesis are attained from the middle interval in the ES-chain:

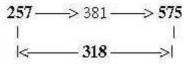
Fig 17-15-d:



Note too that Meth leaves its outer CH3-goup at start of synthesis, (= -15 + 1), which gives R-chain = 61, the intermediate number in the figure above.

575 directly from 208 + 49 = 257 in only two steps:

Fig 17-15e:



Number 75, R-chain of Meth:

In the ES-chain in nb-10 the number 75 = interval 292 - 367 (the sum in the middle of the chain). Transformed in two steps nb-10 to nb-8 it gives the number 159:

 $75 \rightarrow 113 \rightarrow 159$ (161 rewritten)

4.2 A- and T-bases give the sum 575 of ams with non-mixed codons:

Starting numbers 177 and 208 in transformations, minus 1 in each, are the <u>T</u>- and Abases in nb-8. With DNA-base T we get the sum **575** in two steps nb-10 \rightarrow >8: (*Cf. file* 02.)

Fig. 17-16: *A*+*T*

A: $\frac{10 \text{-base}}{135} \longrightarrow \frac{8 \text{-base}/10 \text{-base}}{207} \longrightarrow \frac{8 \text{-base}}{317} \xrightarrow{10 \text{-base}}{320} = \text{AA-AU-coded arms R}$ T: 126 176 $176 \longrightarrow \frac{258}{575} = \frac{255}{575} = \frac{10 \text{-base}}{175} = \frac{10 \text{-base}}{317} = \frac$

How explain the T-base here, a DNA-base giving A in RNA?

4.3 770-group from 4':

It can be added that 2 x 252 (= 4' in the ES-chain) in nb-10 leads directly to 770 in nb-8: $2 \times 4' (252) = 504-10 \rightarrow 770-8$

4.4 Parts of 12-group 770 from halved ES-chain:

The division of group 770 in Cross- and Form-coded ams, **418** and **352**, may be derived by dividing the whole ES-chain in step 4'-3' and halving these numbers:

Fig 17-17: From halved ES-parts to mixed codon groups

Cross-coded = $418 = 2 \times 209$; CA+CA+CU = 210, UG+UG+UC = 208, Form-coded = $352 = 2 \times 176$; GA+GA+GU = 175, AG+AG+AC = 177.

The exponent series:	<u> 292 - 252</u>	183	208 -	159	2	100
702	544			467		
<u> 10-base:</u>	x ½ = 272		X ½	2=234.	(r	ound number)
\downarrow	\downarrow			\downarrow		
<u>8-base:</u>	418			352		
inizimizment (2 x 209		4	2 x 176		

4.5 Derivation of N- and Z-numbers within the two 12-groups of ams:

Fig. 17-18:

G+C-group: $G_8 + C_8 = 384$, difference transformed in 1 step:

 $\frac{10 \text{-base}}{384 \text{-} \mathbf{G}_{10} \, 151 = 233} \xrightarrow{8 \text{-base}} 351 = \mathbf{N} \text{-number in 770-group}$ $\frac{384 \text{-} \mathbf{C}_{10} \, 111}{768} = \underline{273} \xrightarrow{} \frac{421 \text{-} 419}{772 \text{-} 770 \text{-} 768} = \mathbf{Z} \text{-number in 770-group}$

A+U-group: A₈ + U₈ = 367: difference transformed in 2 steps:

<u>10-base</u>	8-base	
$367 - U_{10} 112 = 255 \implies$	<u>377</u>	
377-112 = 265 →	411 ~ 409	= Z-number in 734-group
$367 \cdot A_{10} \ 135 = 232 \longrightarrow$	<u>350</u> ~348	
<u> </u>	325	= N-number in 734-group
734	734	

18. More on totals and other notable transformations

CCC – Why 24 ams? - H-atoms - N-numbers - C-atoms in R - B-chains - 1st to-2nd base

1. Total sum R+B-chains of 24 ams unbound = 3276:

3276 is about 1/10 of 2^{15} . In nb-16 it's CCC, which may be transcribed as 12.12.12 = 3072 (3 x $322 = 4 \times 768$) + 192 + 12:

Fig 18-1: *Total sum of 24 ams R*+*B*:

$$\frac{nb-16}{CCC} \longrightarrow \frac{nb-10}{3276} = 24 \text{ ams } R+B \text{ unbound}$$

$$\downarrow$$

$$12.12.12*$$

$$CC \longrightarrow \frac{nb-10}{204} \longrightarrow \frac{nb-8}{314} (-\pi \times 10^2)$$

$$= Trp, R + B,$$
the heaviest amino acid.

2 x 314 ------> 2848 = 24 ams R+B bound

 $(2 \pi x 100: \text{ the bound } 24 \text{ ams as a closed circle!})$

12.12.12: An association goes to carbon ${}^{12}C$ and the 3C-molecules from halved fructose in glycolysis from which first group of ams derives. Could we eventually read positions of the carbon atoms as decided and guided by oxygen ${}^{16}O$ in some way?! Much of the process in glycolysis seems to be about a stepwise displacement of oxygen along the C-C-chain.

2. Why 24 ams?

One reason to suspect nb-transformations could be the 4 double-coded ams, If 20 ams have to be 24, then 4 ams must be repeated (!).

 $\mathbf{20\text{--}10} \rightarrow \mathbf{24\text{--}8}$

3. H-atoms, 152 in R-chains: and the total of R 1504:

Number of hydrogen atoms in R-chains was 152 = interval 4'- 1' in the ES-series.

This interval is divided 4'-3' = 44 and 3'-1' = 108: Transformed from nb-16 to nb-6 they give total Z-numbers of R-chains and N-numbers separately:

Fig 18-2: H-atoms:

 $44_{16} \longrightarrow 152_6 = H \text{ in R-chains}$ 152 < > 828 = total Z in R-chains

10816 -> 6766 - N in R-chains

Steps $44 \rightarrow 152 = +108$

Step $108 \rightarrow 676 = +568$...This sum is also = 676 = Z (or N) of atoms C, N. O, S. Cf. $676 = 26^2$ and the $2x^2$ -chain,file 13.

4. N-numbers in codon-groups of ams may lead to totals of ams:

Fig. 18-3: Neutron numbers to totals

	10-base	8-base		
G1: N	86>	126		
C1: N	158	236		
U1: N	213	325		
A1: N _	219	<u>333Σ</u> 1020, ~ 1018	$= 2 \times 4 \operatorname{co}$	de bases in 10-base system
\downarrow	1018 ->	1772	= 1772	= 24 B-chains
G2: N	187>	273		
C2: N	58	72		
U2; N	190	276		
<u>A2: N</u>	241	<u>361</u> Σ 982 = 2 x 491		
\downarrow	491 —>	↓ 753, x 2		= 24 R-chains +2H*
*2x491:	982 —>	1726 I		
$\downarrow \rightarrow$	1726 ->	3276	= 3276 =	= 24 ams R + B
0.0000000000000000000000000000000000000		written, sum of 4 RNA		5 1 4 A

- (982 re-written = 1202, +1018 = 2220 (nb-8), = 490 in nb-16,
- 490 in nb-16 = 1168 in nb-10 = 4 x 292 in the exponent series, = 4 x Inosine + 4 x Orotate, = the sum of ams with 3rd base A/G (A or G) or U/C +1.)

5. Number of C-atoms in R-chains as basis for divisions:

In file 04, para. 3, the ams were ordered after number of C-atoms in their R-chains and their mass summed. This division did not concern codon distribution but seemed related to the ES-series with certain assumptions. Here C for carbon. (8 ams with 4 C in R-chains got the sum 584 2 x 292.)

Phe and Tyr are synthesized as 3C- plus 4C-molecules, hence positioned between 4Cand 3C-.groups. Trp as 3C + 4C + 5C - 1C. Trp gets its B-chain from Ser, shares codon with Cys and can brake down to Ala, hence here regarded as "meeting the other way around", added to the 1C group. **Fig. 18-4a:** *Transformations along the ES-chain as a nxC-chain:*

C7			
C4	C3 + C0	C2	C1 + C9 = Trp
$\frac{584}{584}$ $\frac{198}{59}$	<u>306</u>)4	<u>162</u>	$\frac{124}{286}$ $\frac{+130}{}$
2 x 544 1088		2	2 x 208 416
<u>584 = 1088</u> 10-base> 8-ba	= 584 + 198 ase	+ 306	
<u>198 =</u> 10-base —	2 M 2 M		
	<u>306</u> 8-base	1000	286 (re-written) 8-base
			$\frac{286}{10-\text{base}} = \frac{436}{306} = \frac{306 + 130 \text{ (Trp)}}{10-\text{base}}$
		C2 C1	$ \begin{array}{rcl} 162 &=& 242 \\ \underline{124} &=& + & 174 &= & 416 \\ \hline 10\text{-base} &\longrightarrow 8\text{-base} & \\ &=& 286 + 130 \text{ (Trp)} \end{array} $
			<u>174> 256 = 124 + 130</u> + 2H 10-base> 8-base

Fig. 18-4b: Cf. triplet sums, file 15, numbers 714 and 792:

1	C7	1				
C4	1	C3 + C0	C2	C1	+ C9	
584	198	306	162	124	+130 (Trp)	
		790 ~ (C1+U2		1	(C here for ams groups)
0		714~3	71 + A2			

Fig. 18-4c: nxC-atoms - three more details:

The Exponent series:

Different intervals in transformations through re-writings:

<u>10-base</u>	8-base
584 -> 1088	3 ~1~ 890
	1
C7 =	198 out of re-writing
584 <	> 890
and the second se	5 = C3 + C0
200	

Trp:

Interval: 504
$$<$$
 \rightarrow 416
 $|$
 $\underline{88}$ $\xrightarrow{}$ $\underline{130}$ Trp
 $\underline{8}$ $\underline{88}$ $\underline{}$ $\underline{8}$ $\underline{8}$ $\underline{130}$ Trp
 $\underline{8}$ $\underline{8}$ $\underline{8}$ $\underline{1}$

6. B-chains:

6.1 Number 752, sum of first 3 numbers in the ES-chain:

752 from nb-16 to nb-10 gives the total 1772 of 24 B-chains unbound:

292-16 → 658-10 **252-16** → 594-10 **208-16** → 520-10... sum 1772, 24 B-chains unbound

Cf. that 752: nb-10 gave 2848 in nb-6, i.e., R+B-chains of 24 ams bound:

Fig. 18-5:

 $\frac{nb-16}{\frac{12}{2} \times 752}; \ 376 \longrightarrow 886, \qquad x \ 2 = 1772, 24 \text{ unbound B-chains}$

6.2 A single, unbound B-chain = 74:

Two sets of the 4 RNA-bases, sum 1018, gave in nb-8 the sum of 24 B-chains unbound = 1772. A single unbound B-chain à 74 gives the sum of 2 bound B-chains.

Fig 18-6: From one unbound B-chain to two bound ones:

$$\frac{\text{nb-10}}{74} \longrightarrow \frac{\text{nb-8}}{112} = 2 \text{ x } 56 = 2 \text{ B-chains bound}$$

Cf. U-base = 112 A and exchange T to U in mRNA for synthesis.

6.3 Halvings of 2 x 5' 584 transformed to unbound and bound B-chains:

Fig. 18-7: From number 5' in the ES-chain to B-chains in groups of 6:

6.4 Total B-chains unbound times 2 from the 4 bases:

Fig. 18-8:

	10-base	8-base			
4 G	604 —;	> 1134			
4 C	444	674sum 180	08~1810		
			>	$3544 = 2 \times 1772$,	B-chains
4 U	448	700			
4 A	540	1034sum	1734		

6.5 Total of bound B-chains = 1344 from the bases:

Fig. 18-9:

	10-base		8-bas	e		
	194	<>	302	2 x G 2	x RNA-ba	ses read as 8-base numbers
	184	<	270	2 x A		
	146	<	222	2 x C		
÷	148	<	224	2 x U		
	672 —			> 2	x 2 = 1344 ,	= 24 B-chains bound
	388	<	604	4 x G		
+_	352	<	540	<u>4 x A</u>		
=	740	~~>`	1344			= 24 B-chains bound
	<u>10-base</u>		<u>8-bas</u>	<u>e</u>	<u>6-base</u>	
	352 ↓	<	540	4 x A-base		
	352	2		>	1344	= 24 B-chains bound
		<u>16-base</u>		<u>10-base</u>		
	4 x A:	540	\rightarrow	1344		= 24 B-chains bound

6.6 Inosine 136 in repeated steps gives B-chains bound or unbound:

<u>Inosine or Hypoxanthine</u> 136 A (1/4 x 544) may give both B-chain numbers 1344 and 1772 bound and unbound through 4 steps of transformations:

Fig. 18-10:

*Note that without rewritings $530 \sim 528$ and $1020 \sim 1018$ we get 1776 (24 x 74 A).

7. Displacements between 1st and 2nd base order: Numbers 220 - 26:

7.1 Relations between displacements 220 and 26:

Fig. 18-11:

Arns groups R: $G1 + A1 \longrightarrow +246 \longrightarrow G2 + A2$ $C1 + U1 \longrightarrow -246 \longrightarrow C2 + U2$ $G1 \longrightarrow G2 = 220$, $C1 \longrightarrow C2 = -220$ $G1 \longrightarrow A2 = 26$, $U1 \longrightarrow U2 = -26$ $G1 \longrightarrow A2 = 26$, $U1 \longrightarrow U2 = -26$

220 • **26** = **194**, the difference +/-2 in the division of number **416** in the exponent series, (A+U) - (G+C): (A1-G1) - (U1 - C1) = 194 - 2 = 306 - 110. (A2 - G2) - (U2 - C2) = 194 - 2 = -112 + 304.

194 = 2 x 97; an H2PO4 -group, 194 also a charged ribose-P-group in nucleotides.

Fig 18-12:

Number 220 divided	N = 100 (N = 23 (in G-C-group in A-U-group.
G+C-group,	10-base	8-base	/10-base	8-base:
N 100 to 220:	100 —	->144	144 —	→ 220
	\mapsto displ	acement =	Z-number 1	20

The relations between displacement 220 in the G+C-group and 26 in the U+A-group could be explained through only a minus 1 in N- and Z parts and the results in nb-8 through transformations.

Regard number 144 in figure 18-11 above divided in 64 and 80:

Fig. 18-13: How the displacement 220 and 26 could be explained through -1:

10-base 8-base 64 100 N 100 N = N: C1 - C2, +1 = G2 - G1 (101)23 N = N; U1 - U2, -1 = A1 - A2 (22)1-1 – 23 N 63 77 80 120Z $120 \mathbb{Z} = \mathbb{Z}; \mathbb{C}1 - \mathbb{C}2; -1 = \mathbb{G}2 - \mathbb{G}1 (119)$ ⊢___3Z $3 \mathbf{Z} = \mathbb{Z}; \ \mathbf{U1} - \mathbf{U2}; +1 = \mathbb{A1} - \mathbb{A2} (4)$ 1-1 79 117 Z <u>8-base</u> -> 26 N+Z A1 - A2: N 22, Z 4: <u>10-base</u> N 22 <-

Fig. 18-14:

7.2 The number 220 in displacements in group G+C:

Fig. 18-15:

Number 220: = $G1 \rightarrow G2$, $C1 \leftarrow C2$, connected with the sum of ams G+C 544: In relation to numbers of the exponent series:

<u>220 in nb-16</u>; a transition version or reference for the G+C-guided groups 544 between 1^{st} and 2^{nd} base order?

544 + 220 = 764 = C1 + G2 = 353 + 411, difference 58 544 - 220 = 324 = G1 + C2 = 191 + 133, - " - 58

<u>220 in nb-6</u>; representing interval 84 (plus/minus) in the other context where number 544 is received in nb-6, from 208 in nb-10.

A note: Could different divisions of number 544 towards lower numbers in the exponent series be connected with different number base systems? For instance:

544 divided 292-252 = "5"-"4" 544 - " - 336 - 208 = ("5 + 4 - 3") - "3" 544 - " - 177 - 367 = ("5 + 4") - ("3 + 2") - ("3 + 2")10-base 8-base → 145 101 — 44 G1 = 292 - 101336 - 145 = 191 = G1208 + 145 = 353 = C1C1 = 252 + 101Number 220 as a nb-6 number: 6-base 220 ~ 176: G1 = 367 - 176 = 191 re-writing C1 = 177 + 176 = 353

Or: In 2^{nd} base order, using the interval 44 in the transformation nb-10 — nb-8 above? The 3^{nd} division of number 544 in the exponent series: 177 — 367:

8. The 4 double-coded ams, sum = 246

The sum of R-chains of the 4 ams with two different codons are "also" 246, i.e., the sum of displacements 220 and 26 above.

All 4 may become 37 in different nb-systems.

Fig. 18-16:

19- P- phosphorous groups - Coenzymes - Nucleotides - Met AUG

1. P-groups, the single, "inorganic" phosphorous groups:

Fig 19-1: P-groups:

a. $H_2PO_4^- \sim group, 97 A$, $PO3^2^- \sim group, 79 A$, $HPO_3^- \sim group = 80 A$ $HPO_3^- \sim group$ $80 \iff 98 \approx 120 = H_3PO_4 98 A$ $\downarrow = HPO_3^- \sim + H_2O = H_3PO_4$ H_3O $18 \iff 22$, difference

 $PO_{3}^{2} \sim \text{group, 79 A:} \begin{array}{c} \frac{10\text{-base}}{79} & \frac{8\text{-base}}{97 \sim 117} & (79 + 117 = 2 \times 98 = 2 \text{ H}_{3}\text{PO}_{4}) \\ \downarrow & \downarrow \\ \underline{\text{HPO}_{3}} \sim \text{group, 80 A:} & 80 & \longrightarrow \\ 120, \pm 117 & \downarrow \\ 2 \text{ P-groups } 79 + 80 = 159 & \longrightarrow \\ 237 & = 3 \text{ P-groups } 379 \\ (\text{Energy storing in the bindings.}) & \underline{8\text{-base}} \end{array}$

79 ~ 81 = + 2 H

b. Coenzyme groups:

	16-ba	se	6-base			
H ₃ PO ₄ =	98	->	372 = ribose-P-P-P	in coe	enzymes	of bases (-TP)
HP0,~ =	80	>	292 = ribose-P-P	3	19 (S)	(·DP)
2010		<u>10-base</u>				
HPO3		8 0 →>	212 = ribose-P	58		(-MP)

c. NAD (664 A) - NADP (744 A) from P-groups:

16-base 6-base HaPO4; 372 ~ 412 98 + < difference 80 ~ HPO3 ~ 1-18 + HPO.~: 80 292 ~ 332. 744 = 664 = NAD NADP in nb-10 system 10-base 16-base 8-base NADP 1860 3504 744 $= 2 \times 372$ $= 5 \times 372$ 12 x 292 $372 = P \sim P \sim P - ribose$, $292 = P \sim P - ribose$ d. The exponent series: 10-base 6-base 81 = H_PO_~ interval "3 - 2" = 49 -->

A form of life was found some years ago, said to use arsenic instead of phosphorus (P), i. e. next higher element in the phosphorus group of elements in the periodic system. If so, it could of course lead to the conclusion that all such transformations between

97

masses including phosphorus are irrelevant and in any case no necessary condition for life as an eventual part of a reference system.

Yet, phosphorus could have had a decisive role at the very creation of the genetic code, while this not excludes further evolution?

2. Coenzymes of the bases, -MP, -DP, -TP:

2.1 Tables of masses of the coenzymes

Fig. 19-2: Survey

Survey of mass numbers (A) in base-10 system:

4 - 5 code bases, mass numbers, including +1 for bond to ribose: G 151, A 135, U 112, C 111...Σ 509, +T 126...Σ 635

Sum of 2 x 24 bases, 1st and 2nd in the codons: 15 A + 13 U + 11 G + 9 C= 6141

Coenzymes of the code bases:

=	2495		2095		1695
TTP	498	TDP	418	TMP	338
	1997		1677		1357
CTP	483	CDP	403	CMP	323
UTP	484	UDP	404	UMP	324
ATP	507	ADP	427	AMP	347
GTP	523	GDP	443	GMP	363
<u>-TP</u>		<u>-DP</u>		<u>-MP</u>	

2.2 From 4 bases to their mass as coenzymes

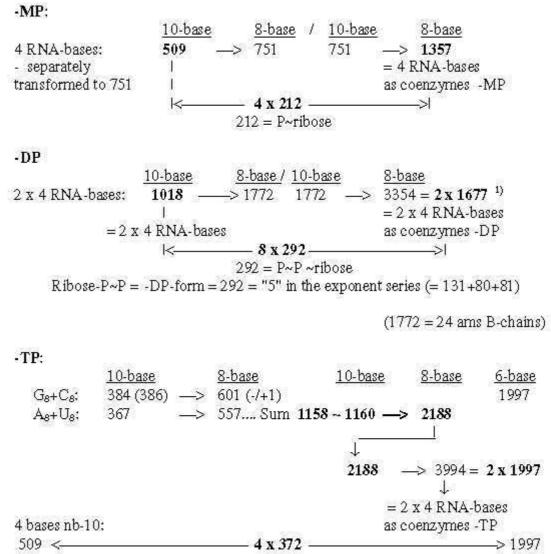
Fig 19-3: 509 - 1357, 4 coenzymes -MP:

4 RNA-bases	1	0-base	8-base	6-ba	se	
	G	151	227	411	411	
	C	111	157	303	~ 263	
	Α	135	207	343	343	
	U	112	160	304	~ 264	
		509	751	1361	1281	
		1	<u> </u>	1357	4	
		751 →>	1357		= 4 RN	A-nucleotides
			\downarrow			charged -1.
		(4 coe	nzymes H	RNA, -MP)		ann an thair a tha an tha an

2.3 Expansion of bases nb-10 to nb-8 adds the Px-ribose groups:

Some transformations from sums of the bases to sums of their appearance as coenzymes are shown in figures below. Note expansions where 212-292-372 correspond to the P(P(P)-ribose groups:

Fig. 19-4: From the bases to coenzymes -MP, -DP, -TP



$372 = P \sim P \sim P - ribose$

2.4 5 bases to 5 coenzymes -TP:

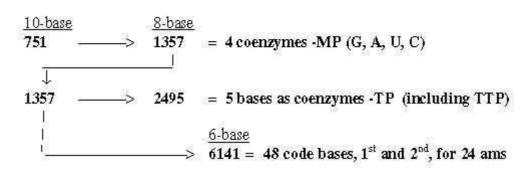
Fig. 19-5:

2.5 4 RNA-bases giving 5 coenzymes -DP-MP in nb-6:

Fig. 19-6:

 2.6 From 751, the sum of 4 bases in nb-8, to 5 bases as coenzymes -TP and to 6141, the sum of 48 codon bases:

Fig. 19-7:



6141 == 15 A + 13 U + 11 G + 9 C:

3. Nucleotides:

3.1 Survey of nucleotides in chain binding:

Fig 19-8:

Nucleotides in chain binding:

<u>RNA:</u> G 345, A 329, U 306, C 305...Σ 1285, ionized -1 in P-groups = 1281
 cGMP = 345, cAMP = 329
 <u>DNA:</u> G 329, A 313, T 304, C 289...Σ 1235, ionized -1 in P-groups = 1231

3,2 Two sets of the nucleotides from 2 sets of the bases (from file 17):

The four RNA-nucleotides in chain-binding and uncharged = 345, 329, 306 and 305 =1285.

The four DNA-nucleotides (= $1285 - 4 \times 16 + 14$ in T-base) = 1235.

Two sets of RNA-nucleotides are given from 2 "times G- and C-bases in three steps nb- $10 \rightarrow > 8$, as two sets of DNA-nucleotides from 2 times A- and U-base:

2G + 2C = 768 in nb-8: $768-10 \rightarrow > 1400-8/1400-10 \rightarrow > 2570-8 = 2 \times 1285 \sim \text{RNA-nucleotides}$ 2U + 2A = 734 in nb-8: $734-10 \rightarrow > 1336-8/1336-10 \rightarrow > 2470-8 = 2 \times 1235 \sim \text{DNA-nucleotides}$ **3.3 ES-number 752 gives in two steps the sum of 4 nucleotides in DNA and RNA:** Fig 19-9:

Nucleotides, 4 DNA 1235 + 4 RNA 1285 = 2520:

4 bses in nb-8: 751/753 $\begin{array}{rcl}
10-base & 8-base \\
752 & \longrightarrow & 1360 \\
& & & & \\
\hline & & & \\
1360 & \longrightarrow & 2520 & = 1285 + 1235
\end{array}$

3.4 The 4 bound RNA-bases in nb-16 gives the 4 RNA-nucleotides in nb-10:

Fig. 19-10:

4 RNA-bases <u>16-base</u> bound = $505 \iff 1285 = 4$ RNA-nucleotides, not ionized \downarrow $780 = 4 \times 195$: \downarrow Definese secure in chain hindinger effects 121 = UDO2 = 6

P-ribose-groups in chain bindings: ribose 131 + HPO2~ 64.

3.5 Bases read as nb-8-numbers, giving cGMP and cAMP in nb-10:

Fig. 19-11: *cGMP* - *cAMP*:

	10-base	8-base	
	167 <	247 A+U	J
	178	262 G+(2
	345	509	345 = cGMP = G-nucleotide
<674*			
	329 <	509	329 = cAMP = A-nucleotide
	10-base	6	-base
*Base pair:	262	$\longrightarrow 6$	$74 = 2/3 \times 1011$, sum of exponent series.
(674 also the	number of atoms		odon bases, 1 st and 2 nd , in the codons.)

4. Met - codon AUG and tRNA-ends ACC:

AUG, the codon for Meth, leads the string at transcriptions from DNA. Chain-bound nucleotides AUG, transformed from nb-10 to nb-8 give the whole sum of 24 ams R, 1504. There is also the equivalence between the 4 bases 509 in nb-8, the A-nucleotide 329 in nb-10 and Meth 149 (R+B) in nb-16,

a. Meth as a kind of reference - or the opposite, the bases a reference to Meth?

 $\frac{16\text{-base}}{149} = \text{Meth, R+B} \implies \frac{8\text{-base}}{509} = 4 \text{ RNA-bases in base-10 system}$

 $\frac{16\text{-base}}{149} \longrightarrow \frac{10\text{-base}}{329} = cAMP, \text{ also} = A\text{-nucleotide}$

The exponent series:

 $\frac{16\text{-base}}{5^{\circ}=292} \longrightarrow 658 = 2 \times 329 (\sim \text{cAMP})$

The "triplet series"

b. A-U-G-nucleotides separately transformed:

	10-base	8-base		543-432-321-210:
Nucleotides:			9 (4 RNA-bases)	343-432-321-210,
	306 U		Σ. 973	= 543 + 432 - 2
	<u>345 G</u>	<u>531</u>	531	= 321 + 210
		= 1504	= 24 ams R	

A+U+G: the mass numbers of the bases interpreted as base-8 numbers:

10-b:	ase:	8 <u>-base;</u>		
93	<	135: A		
74	<	112: U		
+ 105	<	151: G		
= 272	< [398	$272 = \frac{1}{2} \times 544$,	"5" + "4" in the exponent series

5. A-C-C - ends of tRNA:

A-C-C make up the common ends of tRNAs and one may ask why? The three bases (as unbound) give the sum 544 - 1, the sum 5' + 4', 292 + 252 in the ES-series, when transformed in nb-8.

Fig 19-13: tRNA-ends ACC:

c. ACC-ends of tRNAs:

Cf. mass numbers for A and C from Triplets, file 21;

012 + 123 = 135 (A-base), + 234 = 357. Two of the intervals in the steps = 2 x 111 (2 x C-base).

20. Additions to files 17 - 18

1. Rewritings

1.1 Rewriting G - C:

G- and C-bases transformed further to nb-6 becomes sums in later steps of the ES-chain, through rewritings, implying -44:

Fig. 20-1:

544 460 367 259 1 1 11 λ. 1 1 292 252 208 159/158 100 0 10-base 6-base $= 411^* \sim 367 = "3 + 2"$ re-written $(367 = A_8 + U_8)$ G-base 151 1 - 108C-base $= 303^* \sim 259 = "2 + 1"$ re-written 111

*411 = sum of G2-coded ams.

Cf. 44 = the interval 252 - 208 = 4' - 3'. G1 + C1 = 544 divided $\frac{177 + 367}{62}$: C2 = 177 - 44 = 133 G2 = 367 + 44 = 411

1.2 Number 65 - 101 - 81, bases and codon-grouped ams:

Fig. 20-2:

292 - 101 = 191 = G1-coded ams R 252 + 101 = 353 = C1-coded ams R 16-base 10-base 8-base 6-base \rightarrow 101 65 65 \longrightarrow 101 ~ 81 re-written* \rightarrow 145 1 **292** - $65 = 227 = G_g$ **272**, - $65 = 207 = A_g$. (207 + 2 x 145 = A1) $+ 252 + 65 = 317 = C_8 + U_8 + 272 + 65 = 337 = (U+C)_8 (317 + 145 = U1 - 1)_8$ = 544 544 $317 = U_8 + C_8$ **272** - 81 = 191 = G1. **544** - 81 = 463 = U1. *Cf. ams-groups: 272 + 81 = 353 = C1. 416 + 81 = 497 = A1.

[U 112 and C 111 = 223, transformed together = 337-8. Further transformed to nb-6 = 1011 = total sum of the ES-chain in nb-10..]

1.3 Simple rewriting of 2 x 5', 4', 3' in the ES-chain, taken as nb-8 numbers:

This rewriting gives closely the two sets of ams, sums of G1+G2, C+C2 etc.

 $2 \times 292-10 = 584$. $584-8 \sim 604 = G1 + G2 + 2$; $\rightarrow 604 + 416 = 1020 = A1 + A2$ $2 \times 252-10 = 504$, $504-8 \sim 484 = C1 + C2 - 2$; $\rightarrow 484 + 416 = 900 = U1 + U2$

Fig. 20-3:

	8-base	->	8-base	Ams-groups R-chain	s in base-10 system
2 x 292:	584	~	604 =	G1 + G2 +2	(191 + 411)
2 x 252;	504	N	484 =	C1 + C2 - 2	(353 + 133)
2 x 208:	416	÷	604 = 1020	0 = A1 + A2	(497 + 523)
	<u>416</u>	+	484 = 900	0 = U1 + U2	(463 + 437)

1.4 From A-base to 273, mean value of 2 ams R+B:

Fig. 20-4:

10-base8-base10-base8-baseA-base:135
$$\rightarrow$$
207, ~ 187 re-written187 \rightarrow 273x 12 = 3276 \downarrow Mean value of 2 arns R+B in base-10 system

2. Parents of the codon bases, Inosine 136 and Orotate 156:

It was found (file 03) that the sum 292 of the parens to the base-types, when distributed to following numbers in the ES-chain, x 2, gave the codon-groups of ams C1 + U1 and G1 + A1:

Fig. 20-5:

292 —	2.52	— 208	"5-4-3" in the exponent series
┝──>	+ <u>156</u>	—> + <u>136</u>	Orotate and Inosine added
Sums:	408	344	
x 2 =	816	688	
	C1+U1	G1+A1	

Fig. 20-6: The nb-10 and nb-8 numbers added (!), a curious operation:

3. Number 888 in different appearances:

Fig. 20-7:

888 in nb-10 = 543 + 345, numbers of the triplet series = $12 \times B$ -chains à 74 A

888 in nb-8 =
$$1110_8 = 584$$
 in nb-10 = 2 x 292 in the exponent series.

888 in nb-6 = 344 in nb-10 = 888 • 544. 344 x 2 = 688 = ams-groups G1 + A1. 344 in nb-6 = 136 (= Inosine) in nb-10 (1/4 x 544).

888 in nb-16 = 2184-10 = 4 x 546, 8 x 273 (the mean value of 2 ams R+B = 273)

4. Difference of bases in nb-10 and nb-8, read in nb-16, gives 2 x 272 = 544;

Fig. 20-8:

<u>nb-8</u>	<u>nb-10</u>	16-base	10-base	8-base	
G:227 -	151 = 76	76 -	—> 118	166	
C:157 -	111 = 46	46	70	106sum	272
U;160 -	112 = 48	48	72	110	
A:207 -	135 = 72	72	114	162sum	272
			374		$544 = \operatorname{ams} G + C(R)$

5. DNA-bases transformed giver as intervals the G+C- and T+A-pairs and 752:

Fig. 20-9:

	<u>10-base</u>	8-base	
4 DNA-bases:	523 < 4	90> 1013	
	1 .		$\rightarrow 261 = base pair A_{10} + T_{10}$
	490 <26	i2 ——> 752	
		L.	
	base pair	$G_{10}+C_{10}$	

6. Sum of the whole ES-chain 1011:

6.1 N +3 and Z +3 from the ES-chain transformed separately and whole:

Fig. 20-10:

Cf. sum 3282 and sum of triplet series in

6.2 DNA-bases as nb-6 numbers give the sum of the ES-chain:

Fig. 20-11:

	10-base		6-base	
	67	<	151	G-base
	43	<	111	C-base
	59	<	135	A-base
+	54	<	126	T-base
-	223		523	
	223 —	>	1011 =	the sum of the whole exponent series.
22	.3 = C11	1 + U 112.	elenander de Se	and characterized and the constraint of the constraints of the constraint of the constraint of the constraints

7. Totals, two mere operations

7.1 From ES-number 5' to 1/3 of the total 3276:

Fig. 20-12:

"5": $\frac{16 \text{-base}}{292} \xrightarrow{8 \text{-base}}{444} 444 \text{ x } 4 = 24 \text{ B-chains à 74 A.}$ \downarrow $444 \longrightarrow 1092 \longrightarrow 1092 \text{ x } 3 = 3276 = 24 \text{ arms } \text{R+B}$ $888 \longrightarrow 2184 = 4 \text{ x } 546, \text{ x } 3/2 = 3276$

7.2 G+C-bases transformed two times give 2 times total R 1504:

Fig. 20-13:

Base pair G+C
$$\begin{array}{cccc} 10\text{-base} & 8\text{-base} \\ \hline 262 & \longrightarrow & 386 \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & & \\ & & & & & \\ &$$

8. Individual R-chains of ams related through transformations ?

Transformations often imply additional numbers equivalent with molecules, as e. g. plus CH2. There are formally of course a lot of transformations possible between individual ams, only some of which may correspond to biochemical relations. Some examples are shown in the figure below, here regarding R-chains:

It could be added that all four ams with double codons may transformed get the number 37: Ser AG 31-10 = 37-8, Arg AG 101-6 = 37-10, Ile and Leu 57-6 = 101-6 = 37-10, (file 18, para. 8).

292 -- 252 -- 208 -- 159/158 --- 100 -- 0 40 44 49 59/58 K------133------>I 100 - intervals as starting K----->l numbers: |<------108----->|| R-chains: 10--->8 59--> 73 Asp --> Glu / Lys 58--> 72 Asn --> Gln <u>6->10</u> <u>6</u> -->10 <u>10->8</u> <u>10->8</u> <u>6->10</u> <u>6->10</u> 49---> 59, ~ 61 ---> 75<---> 47 <---> 31 <---> 19<---> 15 Meth Cys Ser Asd Ala |- 10-->8 | 10->8 10->6 Т 31 ---> 37 ---> 101 (~ 61, ~ 57) 73 Glu / Lys E Ł | 10->6 Ser 31 Arg 49 ---> 81 11 | <u>10->8</u> <u>10->8</u> <u>10->6</u> His Т 31 ---> 37 ---> 45 ---> 73 Thr Ser Glu 31 18-base Lys | <u>10->8</u> <u>10->8</u> <u>10->8</u> 47---> 57---> 71 ---> 107 81~101 Arg Ileu / Leu | Tyr 110->16 49---> 31 = Ser 1 6->10 71 ---> 43 = Val <u>8 --> 10</u> <u>8->10</u> 101 (Arg) ---> 65 ---> 53 ---> 43 Pro before ring binding. 100 -----> 64 ---> 52 ---> 42 Pro 8-base 6 ---> 10 6-->10 8-->10 108 ~ 110 110 ---> 42 84 ---> 52 ---> 42 Pro Pro 8-base 10 --> 8 108~ 88 ----> 130 10->8 10->8 8-->8 6-->6 59 --> 73 --> 111 ~ 91 Trp 111~107 Phe Tyr <u>8->10</u> <u>8->10</u> <u>8->10</u> 133 ---> 91 ---> 73 ---> 59 Phe Glu/Lys Asp 8->10 8->10 8->10 93* --> 75 159 --->113 --> 75 <u>6->10 10->8 1</u>0->8 Meth Meth 133 ---> 57 ---> 71 ---> 107 10--->8 Leu/Ileu * 93 --> 135 A-base Tyr <u>10-->6 6-base</u> <u>8-base 10->8</u> Only some of these steps *44*---> 112 ~ 108 108 ~ 88 ---> 130 would possibly have Trp chemical correlations.

21. I. Triplet series — II. An alternative series 151-111

I. The triplet series

I. Triplet series; intervals outwards - inwards:

1.1 Triplet chains in nb-8, transformed to nb-10:

The triplets as 4 numbers in two series, outwards and inwards (as 543-345, 432-234 etc., treated as nb-8-numbers, give in pairs in nb-10 sums 4 x 146, 3 x 146, 2 x 146, 1 x 146, the total 5 times 292 = 5' in the ES-chain.

Intervals in nb-10 "outwards - inwards" = 126, $\frac{1}{2} \times 252$ (4').

Fig. 21-1:

8-base	10-base	Sums	10-base	8-base
345>	229	\rightarrow 4 x 146 \leftarrow	355 <	543
234	156	\rightarrow 3 x 146 \leftarrow	282	432
123	<u>83</u>	$ ightarrow$ 2 x 146 \leftarrow	<u>209</u>	321
012	10	\rightarrow 1 x 146 \leftarrow	136	210
714	478		982	1506

982 = 2 x 491: 491-10→<u>753</u>-8 But 478-10 →<u>736</u>-8.

Triplets read "inwards" approximate the 734-group of ams in middle of the ES-chain, hypothetically representing an inward direction in relation to the 770-group as outward directed.

Cf. for 982 file 18, figure 18-3 and for directions file 14, para 3, figure 14-2.

Fig. 21-2: Number 982:

10-base	8-base
355	543
282	432
209	321
136982	210
982>	1726 = 1506 + 220
4	¥ :
1726>	3276, total sum 24 ams R+B, unbound

1.2 Codon bases read as nb-8-numbers give sums triplets in nb-10:

Fig. 21-3:

	4 DNA	-bases		4 R.N.	A-bases:
	10-base	<u>8-base</u>		10-base	8-base:
G	105 <	<i>⊱</i> 151	G	105	← 151
С	73	111	С	73	111
Т	86	126	U	74	112
A	93	135	A	93	135
Sum:	357	523		345	509
		Tripl	et series "inv	vards"	
4 RNA-base	s →	345 =	345, + 0	1 <u>2</u> = 357	$\rightarrow DNA$
4 DNA-base	s →	357 =	1 234	Í	
			123	Í.	
			012	Ì.	

2. Sums 1506 - 714 and intervals 792:

Fig. 21-4:

The Triplet chain "outwards" - "inwards":

543	345	
432975	234	
321	123	
210531	012	
1506 <── 792	-> 714	
10-base	8-base	
792 —>	1428	= 2 x 714

Fig. 21-5: *Total sum of R for 24 ams, sum 1506 -2 from 2 x 4 bases:*

	10-base	8-base
2 G	302 —>	456
2 C	222	336 792
2 x U	112 x 2 →	• 160 x 2 = 320, ~318
2 x A	135 x 2	207 x 2 = 414, ~394sum 714, 712

3. Number n x 273 from codon bases;, two other transformations:

273, the mean value of 2 ams R+B unbound:

<u>nb-16</u> <u>nb-10</u>

C-base: 111 -> 273

The triplet chain with intervals **111**: 543 - 432 - 321 - <u>210</u>:

$210-10 \rightarrow 546-6 = 2 \ge 273$.

From file 20: Number n x 111, the intervals in the triplet steps:

Fig. 21-6:

<u>1</u>	<u>6-base</u> 975>	<u>10-base</u> 2421	
1506 <	- 444 531>	↓ 1329	1092, x 3 = 3276 = 24 ams R+B
Compare:	666>	1638	x = 3276, = 24 ams R+B

4. The triplet series and number 1875:

Pairs of the triplets = 753 transformed as a number in nb-16 gives 1875 in nb-10. All 4 triplets separately transformed, see figure below, give n x 273 as the differences.

Fig 21-7: Number 1875:

$$543_{16} \rightarrow 1347_{10} \\ 432_{16} \rightarrow 1074_{10} \\ 753 < >1875 \\ 321_{16} \rightarrow 801_{10} \\ 210_{16} \rightarrow 528_{10}$$

Intervals 1347 - 528 = 3 x 273 = **819**, x **4** = **3276**, total R+B of 24 ams.

The sums (pair wise added) reminds of the second spectral line of hydrogen from Balmer series, mentioned in *Introduction:* Formula $1/2^2 - 1/4^2 = 0,1875$. Cf. 210 and spectral line 0,21 (!).

Two other operations give relations between sums and intervals:

 10 **log 1,875** \approx 0,<u>273</u> 00...

 $187,5^{2/3} \ge 100 = 3275,93 \approx 3276$, total of 24 ams R+B

 $[1/4 \times \text{ES-chain numbers} = 73 - 63 - 52 - 39.75 - 25,$ with exponent 3/2 = 623.7. - 375. - 500 - 250.6. - 125: sum ~1875 (1874.32.)

Note: $63 \times 52 = 3276$, total sum of 24 ams R+B. Cf "quark numbers" (in "17 short files")

 $15/8 = 5 \times 3 \times 1 / 4 \times 2 = 1.875$

24 ams R+B = 3276. = $\underline{409} \times \mathbf{8.01}$.

48 codon bases $(1^{\text{st}} \text{ nd } 2^{\text{nd}}) = 6141 = 409 \times 15.01.$

II. An alternative numeral series

Another series, from G- to C-base:

Such a series, not treated above, shows some interesting features:

151 - 141 - 131 - 121 - **111**

First and last numbers = mass of G- and C-bases. The DNA-bases (+1 in A-base) are shown in figure below: $272 = 2 \times 136$ (~ Hypoxanthine), $252 = 2 \times 126 =$ T-base:

Fig 21-8: *An alternative series G - to C:*

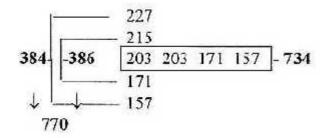
With last three numbers doubled the sum in $nb-10 = 2 \times RNA$ -bases = 1018, in nb-8 = 1772, the 24 unbound B-chains.

All these numbers transformed to nb-8 give the triplet sums 975(543 + 432) - 2 and 531(321 + 210), sum 1504, 24 ams R:

Fig. 21-9:

The 12-groups 770 and 734 of ams are shown in the figure below. Here it may be noted that we get the 734-group in the middle of the chain as in the ES-series, with 2 times 208 in that chain included, corresponding to both 203-groups here.

Fig 21-10:



The ams groups 816 and 688 from -/+ last number 157:

973 - 157 = **816** = U1 + C1 531 + 157 = **688** = G1 + A1. Some other paired groups of ams R from this alternative series:

Fig. 21-11:

With a last step in the chain: 101, plus/minus:

 $\begin{array}{rl} \underline{10\text{-base}} & \underline{8\text{-base}} \\ 101 & \longrightarrow & 145 \end{array}$ 973 - 145 = 828 = Z total 24 ams R 531 + 145 = 676 = N total 24 ams R 973 - 157, + 145 = 961 = A+U +1 531 +157. - 145 = 543 = G+C -1 \end{array}

Fig. 21-12:

848 -- 656 division:

	10-base		8-base
	151		227
	141		215
2 x	131	2 x	203 = 406Sum 848 = G2 + U2
2 x	121	2 x	171 = 342
2 x	111	2 x	157 = 314Sum 656 = C2 + A2

792-712:

Exponent series:

215 + 171 = 386	1993 8 19969999999999999
203 + 203 = 406sum 792	= 2 x 292 + 208
227 + 171 = 398	
157 + 157 = 314sum 712	$= 2 \times 252 + 208$

The doubled last steps re-written:

203~183 =	-20
171~169 =	- 2
157	
107	

22. Other substances

Fats — Sugar — Na-Cl, Na-K-pump

Some annotations about other substances:

1. Fatty acids

Two common fatty acids C18H36O2 = $284-10 \rightarrow 1152-6 (\sim 752 \text{ rewritten}) = 3 \times 384$ and C16H32O2 = $256-10 \rightarrow 1104-6 = 3 \times 368$ are already mentioned in file 17-1:

Fig. 22-1: Two comon fatty acids

	10-base	6-base	Cf. codon type groups of ams:
C18	284	1152	$= 3 \times 384$
			$> 3 \ge 752 = 3/2 \ge 24 \text{ ams R}$
C16	256	1104	$= 3 \times 368$
384 +	1 x 2 = Cross	s- plus Forr	n-coded ams R. $384 = G_8 + C_8$
			coded ams R. $367 = A_8 + U_8$
6-bas€	e 1152 ~ 75	52 re-writte	$n = \frac{1}{2} \ge 24$ ams R.

2. Carbohydrates:

Carbohydrates, some examples, transformations nb-16 \rightarrow 10 \rightarrow 8 or \rightarrow 6: - ${}^{12}C \rightarrow H2O \rightarrow HCOH = \underline{12-16} \rightarrow \underline{18-10} \rightarrow \underline{30-6}$, the building stone of sugar.

- O2 16 A \rightarrow H2CO3 62 A (built into ribose): 32-16 \rightarrow 50-10 \rightarrow 62-8 = + 18, H2O, + 12, C.

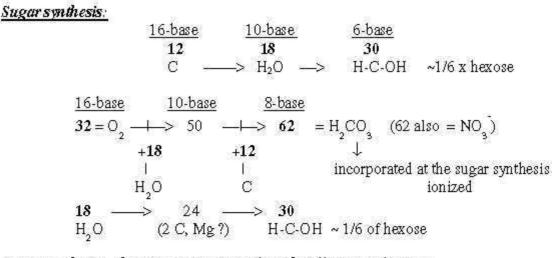
- Hexoses 180 in nb-10: In nb-16 $\underline{180 = 384}$ -10 (= 2 citrate à 192 or e.g. G-8 + C-8).

- A fructose in P-P-bonds = 178: $178-16 = 376-10 = \frac{1}{2} \times 752$ in the ES-chain.

- Ribose 150 as a number in nb-16 = 336 in nb-10, 544 - 208 in ES-chain.

- A disaccharide 342 or two hexoses 180 from ES-numbers as *intervals* in transformation steps:

 $252-16 \rightarrow 594-10 = \pm 342$, a disaccharide. 146-16 $\rightarrow 326-10 \rightarrow 506-8$ (ATP charged -1) = $\pm 180, \pm 180$.



Sugar synthesis - the summation formula with following relations:

	372	372 -	\rightarrow Cf. P-group 98 nb-16 to nb-6.
	<u>16-base</u>	<u>10-base</u>	<u>8-base</u>
6 H	120: 108		x CO2
		180 = 6	$H \cdot C \cdot OH \iff 264 = 6 CO2$
	60	02: 192	— > 300
		<u>10-base</u> 46	<u>8-base 6-base</u>
a)	6 H2O	$108 \rightarrow \vee -$	>154> 300 (-108=192)
	372	< 146	Sum of intervals 46 + 146 = 192
a)	6 CO2	264 — V —	- 410
351-10		= 192	\rightarrow 192 = 6 O2 , in nb-10
		1	> 300
		▶ 108	$\rightarrow 108 = 6 \text{ H2O}$
b)	6 O2	192—	- 300
N. 8 55	372	< 84	Sum of intervals 108 + 84 = 192
b)	6 HCOH	180 — V —	$-264 \rightarrow 264 = 6 \text{ CO2}$ in base-10
0)			

A simultaneous fixation of nitrogen occurs during which Molybdenum take part: Mo 42 Z, 96 A.

If presuming 2 Molybdenum atoms = 84 Z, 108 N = 192 A, sme number as ~ 6 O2, numbers of the transformation intervals above:

 $NO3^{-} = 62 A, x 6 = 372 = 108 + 264$ or 180 + 192.

62 = 108 - 46, 146 - 84, intervals above. **NO3⁻** + **NO2⁻** = 62 + 46 = 108. α -ketoglutarate, aminating amino acids = 146.

Hexoses as intervals - ? - in transformations within the exponent series, i. e.:

<u>16-base</u>	<u>10-base</u>	<u>16-b</u> :	<u>16-base</u>		ase	8-base
252 —⊢>	594	146	>	326	_ >	506 (~ ATP ⁻)
342 ~ disaccharide			180		180 ~	fructose/glucose

3. Na-Cl and the Na-K-pump:

Na-Cl and Na-K-pump in the nervous system:

Na 11 Z \rightarrow Cl 17 Z \rightarrow K 19 Z: <u>Na 11-16 \rightarrow Cl 17-10 \rightarrow K 19-8 Na 23 A, Cl 35 A (most common isotope): <u>Na 23-16 \rightarrow >Cl 35-10</u></u>

Cf. Na, Cl, K ionized, 10 e, 18 e: in nb-10 to nb-8 = +2, number for the transport of 2H through membranes.

Fig. 22-3:

11 Z, 23 A Na Cl 17 Z, 35 A (or. 37 A, mean value 35,4 A in nature) K 19 Z, 39 A (or. 41 A, 0,0018%) Z-numbers -> Z-numbers <--> A-numbers: 16-base 10-base 8-base Z 11 17 Cl, Z Na Z Cl17 19 KΖ > Κ Z 19 -> 23 NaA $A \longrightarrow Z$: 10-base 8-base 16-base K 39~41 -21 A -33 1 1 Ζ 17 2111 C1, Z ~19, K: Z Na Z <u>A:</u> 10-base 8-base 16-base $(27_{10} = 39_6)$ Κ 39 A 27 4 27 Na 23 A < ClA 35 23 Na: A e-numbers: 8-base 10-base 12 = +2Na 10 8 K 22 = + 4 ~ H-wanderings - ? e 18 Cl 22 Ħ through cell membrane 18 e ы H₀ A 18 22

Discussion

The amount of correlations between the genetic code and numeral series is difficult to regard as only random ones.

A general problem is of course that it still doesn't seem to exist any known biochemically accepted mechanisms that could "explain" construction along such numeral series, however established facts in the other mentioned examples. It could however be questioned in which sense the 2x2-series behind the periodic system is "explained", or the formula for spectral lines of hydrogen.) Facts are there. Science has only its models, as far as possible congruent with the facts.

With the hypothesis here that they really reveal features in how Nature organized the genetic code, what should it imply? About the elementary series $5 \rightarrow > 0$, the series of valences for atoms in the genetic code could be remembered: P - C - N - O,S - H = valences 5 - 4 - 3 - 2 - 1. A-dimensional interpretation seems inevitable, with regard to exponents and to transformations between nb-systems.

How should the exponent 2/3 be explained? We have squares in the $2x^2$ -chain behind the periodic system and intervals between inverted squares behind the spectral lines of hydrogen. These formulas concern electron shells of atoms, i. e. the property charge. With mass and charge most elementary assumed as a mutual relation D3 to D2, cubes become natural. We have mass as the energy form concentrated in atomic nuclei, charge expressed in the atomic shell with released energy in kinetic form. Why then inverted cubes? They lead inwards to a deeper level, as does the inward direction toward nucleus in an atom.

It may be remembered too that there are a similar inverted relation between radii and mass in neutron stars.

The many relations of disparate kinds to the $2x^2$ -chain and other simpler chains support the interpretation of the genetic code as built on an elementary chain x = 5 - 0with exponents of different degrees. With a dimensional view on the exponents, it could imply, either that such chains preceded the more elaborated ES-chain when the coding system emerged or could be regarded as simultaneously existing on underlying levels. It's possible to imagine a dimensional development from both ends of the chain towards step 3 - 2 in the middle with increasing agreement of mass distribution in the genetic code:

$$x^4 \rightarrow x^3 \rightarrow [x^{3/2} \rightarrow \leftarrow x^{2/3}] \leftarrow x^2 \leftarrow x^1.$$

The mass distribution as described in section I often implied minus/plus lower numbers in the ES-series, correlating with features in the background model. It points to a twoway direction in he chain of both disintegration and synthesis. This could seem to conflict with the common view on evolution as a stepwise synthesis towards more complex and bigger units. Yet, a double-direction is natural in Nature, if we think of macrocosm, Big Bang and both processes in celestial Hx-clouds. It could be mentioned that even among physicists this opposite view of disintegration, starting from a whole, has been proposed. (There is a similar pattern of two-way direction in the protein synthesis, where tRNAs as from opposite strands of DNA meet mRNA "the other way around" at ribosomes in the "middle" of the process.) See figure 1 in section I, with dimensional interpretation of the forms from double direction (D4) in DNA to singlestrnded RNA as vector (pole 4b) outwards to ribosomes (D3) - meeting tRNAs (as "clover leaves" D2) and ams.

It's shown too that not only mass distribution on codon groups of ams correlates with the ES-chain but also other bases for mass division, for instance with main groups of atom kinds and the not codon-dependant B-chains as well as with several features in the origin of ams from stations in glycolysis - citrate cycle. This suggests an interpretation where the same principle scheme is developed on different levels or as representing different axes in a coordinate system when the genetic code emerged.

The single fact that the mass division on C-skeleton and other atoms (960 and 544) is the same as between main codon groups (U+A, 960 and G+C, 544) supports in itself the general suggestion that the code is built on a numeral series.

In several ways the results seems to agree with the coevolution theory [6, 7]. There is the relation with biochemical origins of ams from glycolysis and citrate cycle. There is the view of codon domains as totals, differentiated in following steps, even if the "codon domains" here is related to mass sums of ams. There is also the fact that G1-coded ams "arrive first" in the number chain as 5 out of about 7 ams assumed first in that theory: GG-GC-GU-GA-GA besides Ser UC and Phe UU..

Then about mass again, rejected as irrelevant for codon assignments: In addition to arguments in the *Introduction* it's reasonable to ask for instance why precisely these ams have been selected for coding, not other ones? The selection seems rather random. Why just this number of ams with oxygen as end groups, that number of ams with nitrogen? (Besides that both types and polar and non-polar ams surely have been necessary.)

Further, when much research in this field has been focusing on the "most stable" configuration of the coding system, one could naturally ask what the background is for this stability? One aspect is of course that the most common isotopes have shown up to be most stable. (When calculating with common mix of isotopes today, atomic weights should change the sum of R- plus B-chains of ams from $3276 \rightarrow > 3280$ abbreviated, R-chains from $1504 \rightarrow > 1506$, no more than the deviations of single units (u) in this analysis.) In addition, the analysis here mostly concerns groups of ams, i. e. sums were an individual deviation in mass might have a rather small influence.

The fact that Ileu sometimes gets mixed with Leu by tRNAs could also be mentioned, differing in structure but having the same mass and atoms.

Does the proposal for a guiding numeral series exclude such an individual invention among certain organisms as Pyl, called the 22nd ams, occupying a stop codon? Pyl adds 108 to R-chain of Lys, i. e. the interval 3' to 1' in the ES-chain and could eventually be suspected as a "misreading" of the chain, leading to a compound, a new "word"?

The examples of transformations between nb-systems are astonishing and certainly provocative. They support however a general dimensional view in the creation of the code and actually too the relevance of the ES-chain. They seem to reveal a deep level in the reference system of a hitherto unknown kind, representing the very steps between dimensional degrees. In physical and biochemical terms they should imply something like mutual resonances between "mass fields" in different dimensional degrees, relations and fragmentation guided by geometrical and arithmetical rules. A problem is naturally the superfluity of such possible transformational relations.

If proposals in this paper are accepted as hypotheses, they will naturally raise many new

questions and lead to secondary hypotheses, which in their turn could be possible to test. The dimensional aspects, mostly omitted here, should reasonably, if elaborated further, have implications for protein structures and their different functions in cells.

Whatever to believe about the arithmetic, something of that kind resembles life - in being very simple and very productive - and naturally multidimensional.

*

References

References, referred to in the text within brackets []:

1. Chou KC: "Prediction of Protein Cellular Attributes Using Pseudo-Amino Acid Composition." PROTEINS: Structure, Function and Genetics, 43, 246-255 (2001).

2. shCherbak VI, "Arithmetic inside the universal genetic code," [Abstract],

Biosystems 70 (3), 187-209 (2003).

- 3, Rakocevic MM, "A harmonic structure of the genetic code." J theor. Biol. 229, 221-234 (2004).
- 4.Downes AM, Richardson J, "Relationships Between Genomic Base Content and Distribution of Mass in Coded Proteins." J Mol Evol 55, 476-490 (2002).
- 5. Perez JC, "Codon populations in single-stranded whole human genome DNA are fractal and fine-tuned by the Golden Ratio 1.618." Interdiscip Sci 2(3), 228-240 (2010).
- 6. Wong TF, "The evolution of a universal genetic code." Department of Biochemistry, University of Toronto, Canada. Communicated by J. Tuzo Wilson. (1976)

7. Wong JT, "Coevolution theory of the genetic code at age thirty." [Abstract], Bioessays, 27, 416-425 (2005).

Since this research started in the beginning of 1980's, the main source used was:

Karlson P., 1974. Biokemi (*Biochemisttry*). Liber Läromedel, Lund, Swedish version of Karlson, P. Introduction to Modern Biochemistry, forth edition, 1975, Academic Press Inc. with later, new editions.

A few data taken from

a. Lindahl P E, Kihlström J E, Kiesling K-H, Sundell L-E: Zoofysiology (*Zoophysiology*) 1967, Almqvist & Wiksell, Uppsala.

b. Nicholson D. F 1976. Metabolic Pathways. Koch-Light Laboratories, UK.[A map over main processes in biochemistry.]

c. Wikipedia, the free encyclopedia, i.e. http://en.wikipedia.org/wiki/Histone#Classes.

A selection of other articles dealing with the same topic:

Arquès DG, Michel CJ: A Circular Code in the Protein Coding Genes of Mitochondria. J. theor. Biol. 1997, 189:273-290.

Arquès DG, Michel CJ: A Complementary Circular Code in the Protein Coding Genes. J. theor Biol. 1996, 182:45-58.

Balázs A: Some introductory formalizations on the affine Hilbert spaces model of the origin of life. I. On quantum mechanical measurement and the origin of the genetic code: A general physical framework theory. BioSystems 2006, 85:114-125.

Bedian V: Self-description and the origin of the genetic code. BioSystems 2001, 60:39-47. **Berger G**: Deterministic hypotheses on the origin of life and of its reproduction. Med Hypotheses 2003, 61(5-6):586-592.

Biro J C, Benyó B, Sansom C, Szlávecs A, Fördös G, Micsik T, Benyò Z: A common periodic table of codons and amino acids. BBRC 2003, 306:408-415.

Chechetkin V R: Block structure and stability of the genetic code. J theor Biol. 2003, 222:177-188.

Chou, K-C: Using amphiphilic pseudo amino acid composition to predict enzyme subfamily classes. Bioinformatics 2005, 21.1:10-19.

Copley S D, Smith E, Morowitz H J: A mechanism for the association of amino acids with their codons and the origin of the genetic code. Proc. Natl. Acad. Sci. 2005, 102:4442-4447.

Damjanovic Z M, Rakocevic M M: Genetic Code. An Alternative Model of Translation. Ann. N. Y. Acad. Sci. 2005, 1048:517-523.

Delarue M: An asymmetric underlying rule in the assignment of codons: Possible clue to a quick early evolution of the genetic code via successive binary choices. RNA 2007, 13:161-169. **Di Giulio M**: The origin of the genetic code: theories and their relationships, a review. BioSystems 2005, 80:175-184.

Hartman H: Speculations on the Evolution of the Genetic Code. KIV. The Evolution of the Aminoacyl-tRNA Synthetases. Orig Life Evol Biosph. 1995, 25:265-269.

Heal J R, Roberts G W, Raynes J G, Bhakoo A, Miller A D: Specific Interactions Between Sense and Complementary Peptides: The Bases for the Proteomic Code. ChemBioChem. 2002, 3:136-151.

Hobish M K, Wickramasinghe N S M D, Ponnamperuma C: Direct Interaction between Amino Acids and Nucleotides as a possible physicochemical Basis for the Origin of the Genetic Code. Adv Space Res. 1995, 15:(3)365-(3)382.

Hoffmeyer J: Life and reference. BioSystems 2001, 60:123-130.

Hornos J E M, Braggion L, Magini M, Forger M: Symmetry Preservation in the Evolution of the Genetic Code. (Hypothesis Paper). IUBMB Life 2004, 56(3):125-130.

Ikehara K: Origins of gene, genetic code, protein and life: comprehensive view of life systems from GNC-SNS primitive genetic code hypothesis. J. Biosci. 2002, 27:165-186.

Jiménez-Montaño M A: Protein evolution drives the evolution of the genetic code and vice versa. BioSystems 1990, 54:47-64.

Johnson F Yan, Alexander K Yan, Benjamin C Yan: Prime Numbers and the Amino Acid Code: Analogy in Coding Properties. J. theor. Biol. 1991, 151:333-341.

Judson O P, Haydon D: The Genetic Code: What Is It Good For? An Analysis of the Effects of Selection Pressures on Genetic Codes. J Mol Evol. 1999, 49:539-550.

Karasev V A, Stefanov V E: Topological Nature of the Genetic Code. J theor Biol. 2001, 209:303-317.

Knight R D, Landweber L F: Rhyme or reason: RNA-arginine interactions and the genetic code. Chem Biol. 1998, 5:R215-R220.

Kohler H, Murali R, Kieber-Emmons T: The hidden code in genomics: a tool for gene discovery. Review. J. Mol. Recognit. 2001, 14:269-272.

Lu Y, Freeland S: On the evolution of the standard amino-acid alphabet. Genome Biol. 2006, 7:102:102.1-102.6.

Michel C J: An Analytical Model of Gene Evolution with 9 Mutation Parameters: An Application to the Amino Acids Coded by the Common Circular Code. Bull Math Biol. 2007, 69:677-698.

Mussat M, Bégin M E, Bureau J P: A constructionist model predicting the emergence, complementarity and classification of the nucleotide bases. Med Hypotheses 1998, 51:511-523. Osawa S, Jukes TH: Codon reassignment (codon capture) in evolution [abstract. J Mol Evol. 1989, Apr; 28(4):271-8.

Patel A: The triplet genetic code had a doublet predecessor. J. theor. Biol. 2005, 233:527-532. **Ronneberg T A, Landweber L F, Freeland S J**: Testing a biosynthetic theory of the genetic code: Fact or artifact? Proc. Nat. Acad. Sci. 2000, 97:13690-13695.

Sanchez R, Grau R: A genetic code Boolean structure. II. The genetic information system as a Boolean information system. Bull Math Biol. 2005, 67:1017-1029.

Schimmel P, Giegé R, Moras D, Yokoyama S: An operational RNA code for amino acids and possible relationship to genetic code. Proc Natl Acad Sci. 1993, 90:8763-8768.

Sciarrino A: A mathematical model accounting for the organization in multiplets of the genetic code. BioSystems 2003, 69:1-13.

Seligmann H, Amzallag G N: Chemical interactions between amino acid and RNA: multiplicity of the levels of specificity explains origin of the genetic code. Naturwissenschaften 2002, 89:542-551.

shCherbak V I: Twenty Canonical Amino Acids of the Genetic Code: The Arithmetical Regularities. Part I. J. theor. Biol. 1993, 162:399-401.

shCherbak V I: Arithmetic inside the universal genetic code. BioSystems 2003, 70:187-209.

Shen N, Guo L, Yang B, Jin Y, Ding J: Structure of human tryptophanyl-tRNA synthetase in complex with tRNATrp reveals the molecular basis of tRNA recognition and specificity. Nucleic Acid Res. 2006, 34:3246-3258.

Sitaramam V: Genetic code preferentially conserves long-range interactions among the amino acids. FEBS Lett. 1989, 247:46-50.

Sowerby S J, Petersen G B, Holm N G: Primordial Coding of Amino Acids by Adsorbed Purine Bases. Orig Life Evol Biosph 2002, 32:35-46.

Sukhodolets V. V: The Genetic Code as a Clue to Understanding of Molecular Evolution. J. theor. Biol. 1989, 141:379-389.

Sungchul J: The Linguistics of DNA: Words, Sentences, Grammar, Phonetics, and Semantics. Ann N Y Acad Sci 1999, 870:411-417.

Trevors J T, Abel D L: Chance and necessity do not explain the origin of life. Cell Biol Int. 2004, 28:729-739.

Wilhelm T, Nikolajewa S: A New Classification Scheme of the Genetic Code. J Mol Evol 2004, 59:598-605.Woese C R: A Proposal Concerning the Origin of Life on the Planet Earth. J Mol Evol. 1979, 13:95-101.

Wong J T-F: Question 6: Coevolution Theory of the Genetic Code: A Proven Theory. Orig life Evol Biosph 2007, 37:403-408.

Wu H-L, Bagby S, van den Elsen J M H: Evolution of the Genetic Triplet Code via Two Types of Doublet Codons. J Mol Evol. 2005, 61:54-64.

Yarus M: Amino Acids as RNA Ligands: A DirectRNA-Template Theory for the Code's Origin. J Mol Evol. 1998, 47:109-117.

Yarus M, Caporaso J G, Knight R: Origins of the Genetic Code: The Escaped Triplet Theory. Annu. Rev. Biochem. 2005, 74:179-198.

Yu J: A Content-Centric Organization of the Genetic Code. Geno. Prot. Bioinfo. 2007, 5:No. 1 (1-6).

END