HOW TO KNOW IN WHICH LINE ONE CHILD CAN SHOW MAXIMUM ABILITY

SUNIRMAL ROY

ABSTRACT : Mainly nineteen cellular entities take place in mainly thirty chemical reactions in a cell. The cell is divided into two when the cellular entities are doubled at a time. In this way growth process takes place in a living body.

From the solution space of this cell generation it is possible to explain mathematically why the shapes of different species are different and why the shapes in the same species are same.

Relative growth rate of different parts of a body remains the same constant for the same species and these constants are different for different species. This is the cause of the same shape of all living things of the same species. This is verified for some castes of India. Similarities are also found in the ratios among the same type of experts (specially of sports). Observing these similarities in a child it can be predicted in which line the child will show his or her maximum ability.

INTRODUCTION, FORMULATION AND SOLUTION OF THE

PROBLEM

Growth is a basic property of biological species, and growth coupled with cell-divisions leads to an increase in population. In F. Heinmets1 we see that the rate constants K_1 , K_2 , K_n are taken to be constants throughout the whole generation time of cell. When the value of one rate constant is decreased then other equations will balance it and obviously the value of other rate constants will be changed. Therefore we cannot use the conception of rate constants. Hence the author has modified the model of F. Heinmets1 in a different way. The author has rearranged the Table as follows :-

Terminology and Symbols

Pools:-

into RNA

into amino acid.

P0015 :-		
	Pe	extra-cellular nutrient pool
	Pi	General intra cellular metabolic pool.
	Pa	Amino acid pool for protein synthesis.
	Pn	Nucleotide pool for RNA synthesis.
Enzymes	:	
	Е	Total protein.
	En	Enzymes which convert internal pool (P_i)
precurso	rs	
	Ea	Enzymes which convert internal pool (P_i)

Е_р RNA polymerase for messenger RNA (M) synthesis. E, Enzymes which convert external pool (P_a)

into internal

pool (P_i).

Rate constant Kn, Ka and Kt determine what fraction of total

protein represents respective enzymes.

E	Genes:-		
e s		G _е	Genes for messenger RNA (M) synthesis.
s		G _р	Gene for messenger RNA (MP) synthesis.
		Gb	Gene for the synthesis of RNA fraction
	ribosome.		
		G _c	Gene for transport RNA (C) synthesis.
	Messenge	ers:-	
		М	Messenger (RNA) for protein (E)
	synthesis.		
		Мр	Messenger (RNA) for Epsynthesis.
		B'	RNA fraction of ribosome.
		В	Ribosome.
		С	Transport RNA
)		Ν	Ribosome & messenger complex for
	protein (E) synthesi	s	
		Np	Ribosome and messenger complex for Ep
)	synthesis	(templa	ate).
		Ν	Inactive state of N.
		N'p	Inactive state of NP.

International Jour ISSN 2229-5518	nal of Scie	entific & Engineering Research, Volume 6, Issue 3, 1	March-2015		364				
1551 2227-5510	Si	Metabolite which converts templates N		15.	$r_{51}x_{15}$ of C = $r_{52}x_{15}$ of Pi.				
and Np into inactiv				16.	$r_{53}x_{16}$ of E = $r_{54}x_{16}$ of Pi.				
	Si'	Metabolite which converts inactive		17.	$r_{55}x_{17}$ of Ep = $r_{56}x_{17}$ of Pi.				
template N and NF				18.	r ₅₇ x ₁₈ of Pe+r ₅₈ x ₁₈ of Ei = r ₅₉ x ₁₈ of				
к ₁ кл	variousra	ate constants.	Pi+r ₆₀ x ₁₈ of Et.						
		TABLE I		19.	$r_{61}x_{19}$ of Pi+ $r_{62}x_{19}$ of En = $r_{63}x_{19}$ of				
	1.		Pn+r ₆₄ x ₁₉ of En.		01 13 02 13 03 13				
	2.	r_1x_1 of $E_p + r_2x_1$ of $P_n = r_3x_1$ of E_pP_n	64-19	20.	r_v ofPi∔r v ofFa−r v of				
		$r_4 x_2$ of B+ $r_5 x_2$ of E = r6x2 of B	Dava v of Ca	20.	$r_{65}x_{20}$ of Pi+ $r_{66}x_{20}$ of Ea = $r_{67}x_{20}$ of				
	3.	$r_7 x_8$ of GB+ $r_8 x_3$ of Pn = $r_9 x_3$ of	Pa+r ₆₈ x ₂₀ of Ea.						
GB+r ₁₀ x ₃ of B				21.	$r_{69}x_{21}$ of Ep+ $r_{70}x_{21}$ of C = $r_{71}x_{21}$ of				
	4.	$f_{11}x_4$ of GC+ $r_{12}x_4$ of P = $r_{13}x_4$ of	EpC.						
$GC+r_{14}x_4$ of C				22.	$r_{72}x_{22}$ of E+ $r_{73}x_{22}$ of Pi = $r_{74}x_{22}$ of Epi.				
	5.	$r_{15}x_5$ of GP+ $r_{16}x_5$ of P = $r_{17}x_5$ of		23.	$r_{75}r_{23}$ of Ep+ $r_{76}r_{23}$ of B = $r_{77}r_{23}$ of				
GP+r ₁₈ x ₅ of M p			EpB.						
	6.	$r_{19}x_6$ of Ge+ $r_{20}x_6$ of Ep P = $r_{21}x_6$ of		24.	$r_{78}x_{24}$ of Np+ $r_{79}x_{24}$ of Si = $r_{80}x_{24}$ of				
Ge+r ₂₂ x ₆			N'p.						
		of Ep+r ₂₃ x_6 of M.		25.	$r_{81}x_{25}$ of N'p+ $r_{82}x_{25}$ of Si = $r_{83}x_{25}$ of				
	7.	$r_{24}x_7$ of B+r $_{25}x_7$ of M = $r_{26}x_7$ of N.	Np.						
	8.	$r_{27}x_8$ of B+ $r_{28}x_8$ of Mp = $r_{29}x_8$ of Np.		26.	$r_{84}x_{26}$ of N+ $r_{85}x_{26}$ of Si = $r_{86}x_{26}$ of N.				
	9.	$r_{30}x_9$ of C+ $r_{31}x_9$ of P = $r_{32}x_9$ of C Pa.		27.	$r_{87}r_{27}$ of N'+ $r_{88}r_{27}$ of Si = $r_{89}r_{27}$ of N.				
				28.	$r_{90}x_{28}$ of N = $r_{91}x_{28}$ of Pi.				
	10.	$r_{33}x_{10}$ of N+ $r_{34}x_{10}$ of CPa = $r_{35}x_{10}$ of		29.	$r_{92}x_{29}$ of Np = $r_{93}x_{29}$ of Pi.				
^{B+r} 36 ^x 10				30.	$r_{94}x_{30}$ of Pi = $r_{95}x_{30}$ of X.				
		of $C+r_{37}x_{10}$ of	Where	^r 1 ^{, r} 2	r_{95} are constants. The equations of Table 1				
M+r ₃₈ x ₁₀ of B			are written on th	e understa	anding that the quantitative measure of the				
	11.	$r_{39}x_{11}$ of NP = $r_{40}x_{11}$ of CPa = $r_{41}x_{11}$ of	entities in a rea	ction are	one to another in constant ratios. These				
^{B+r} 42 ^x 11			constants are r	1, r2et	c. which can be calculated. These are				
		of	independent of th	nerate co	nstants which are not fixed. Therefore the				
Mp+r ₄₃ x ₁₁ of C+i	r ₄₄ x ₁₁ of I	Ep	defect of F Heinm	ets ¹ iselim	inated in this model.				
	12.	$r_{45}x_{12}$ of M = $r_{46}x_{12}$ of Pn.			f the functional entities.				
	13.	$r_{47}x_{13}$ of M p = $r_{48}x_{13}$ of Pa	-	Ep, Pn, Ep, B', B, C, M, Mp, N, N'p, Pa, CPa, E, Pi, EpC, EpB,					
	14.	$r_{49}x_{14}$ of B = $r_{50}x_{14}$ of Pi	Epi, respectively in	nitially be					

ISSN 2229-5518 X₃₁, X₃₂, X₃₃, X₃₄, X₃₅, X₃₆, X₃₇, X₃₈, X₃₉, X₄₀, X₄₁, X₄₂,

X₄₃, X₄₄, X₄₅, X₄₆, X₄₇, X₄₈, X₄₉.

Since it is assumed that all functional entities are approximately doubled after one generation time of the cell, therefore we get the equations of Table 2.

- TABLE 2
- 1. $x_{31} = {}^{-r}1^{x}1^{+r}22^{x}6^{+r}44^{x}11^{-r}55^{x}17^{-r}69^{x}21^{-r}75^{x}22$
- 2. $x_{32} = -r_2 x_1 r_8 x_3 r_{12} x_4 + r_{16} x_5 + r_{46} x_{12} + r_{48} x_{13} + r_{66} x_{19}$
- 3. $x_{33} = r_3 x_1 r_{20} x_6$
- 4. $x_{34} = -r_4 x_2 + r_{10} x_3$
- 5. $x_{35} = {}^{r}6^{x}2^{-r}24^{x}7^{-r}27^{x}8^{+r}35^{x}10^{+r}41^{x}11^{+r}49^{x}14^{-r}76^{x}23^{x}10^{+r}41^{x}11^{+r}49^{x}14^{-r}76^{x}23^{x}10^{+r}10^{-$
- 6. x₃₆ = r14x4-r30x9+r36x1+r43x11-r51x15-r70x21
- 7. $x_{37} = r_{23}x_6 r_{25}x_7 r_{45}x_{12} + r_{37}x_{10}$
- 8. $x_{38} = r_{18}x_5 r_{28}x_8 + r_{42}x_{11} r_{47}x_{13}$
- 9. $x_{39} = r_{26}x_7 r_{33}x_{10} r_{84}x_{26} + r_{89}x_{27} r_{90}x_{28}$
- 10. $x_{40} = r_{29}x_8 + r_{39}x_{11} + r_{78}x_{24} + r_{83}x_{25} + r_{92}x_{29}$
- 11. $x_{41} = r_{86}x_{26} r_{87}x_{27}$
- 12. $x_{42} = r_{80}x_{24} r_{81}x_{25}$
- 13. $x_{43} = r_{31}x_9 + r_{67}x_{20}$
- 14. $x_{44} = r_{32}x_9 r_{34}x_{10} r_{40}x_{11}$
- 15. $x_{45} = -r_5 x_2 + r_{38} x_{10} r_{53} x_{10} r_{72} x_{22}$

16. $x_{46} = r_{50}x_{14} + r_{2x_{15}} + r_{54}x_{16} + r_{56}x_{17} + r_{59}x_{13} - r_{61}x_{19} + r_{91}x_{28} - r_{65}x_{20} + r_{93}x_{29} - r_{73}x_{22} - r_{94}x_{30}$ 17. $x_{47} = r_{71}x_{21}$

- 18. $x_{48} = r_{77} x_{23}$
- 19. $x_{49} = r_{74} x_{22}$

Since x_{31} , x_{32} , ..., x_{49} are fixed in an individual body and the ratios r_1 , r_2 r_{94} can be calculated therefore there are 30 variables x_1 , x_2 x_{30} , and 19 equations.

There are 30 variables and 19 equations and hence 11 variables can be given arbitrary values satisfying the corresponding equations. Let the solutions make V vector space. Let the solutions of x_1 , x_2 x_{30} during the whole life of an individual body make a vector space

S. Then S is a subspace of V. Let the solutions of $x_1, x_2, ..., x_{30}$ (during the whole life) of different organs, tissues, different parts of the body make solution vector spaces $S_1, S_2, ..., etc.$ Then $S_1, S_2, ..., etc.$ are subspaces of S and hence subspaces of V. Considering $X_{31}, X_{32}, ..., X_{49}$ these subspaces $S_1, S_2, ..., S_n$ and hence S can be studied and the relations of different organs, tissues with each other and with the whole body can be known.

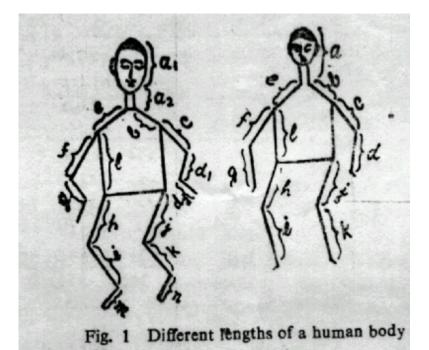
The relative growth of different organs, tissues can be shown as constant for an individual body. For this reason, the individual body remains it's shape unchanged. For another individual body of the same species similarly the solution vector-spaces can be studied which is a subspace of V and similarity of growth and shape of the two individuals of the same species can be explained.

The solution spaces of other individual bodies of other species are also subspaces of V and hence similarities and dissimilarities of different individuals of different species can be studied in this respect.

Growth process remains unstopped throughout the whole life through the cellular division. But when different organs, different parts of the body gets stable size then and when the time of cell division is too much greater than the remaining life time then also the growth of that organ is continued, but the growth of the body is not marked and it seems that growth of that part is stopped.

The equations can be discussed in different way also. The activities of different genes in different organs, different parts of a body are not the same. In one organ one gene can be inactive and in another organ the same can be active. Thus we will get different set of equations for different parts of body and obviously we will get different solution spaces, different nature of different parts. But the cell generation time for all cells of all organs are same though solutions may be different. For this, growth rate of one organ is same as the growth rate of another organ and relative growth rate of another organs remains the same. This is understood from the experimental results also.

Verification of the constants of relative growth of different parts of 400 human body of different castes and experts of different field, where the parts are shown in Fig 1 (Denote first part by X and second part by Y, lengths are in cm.)



					ij	44.43	47.2	9.46	1.55				
						.976 =-24.24+1.60x							
VI	ERIFICA	TION FO	OR SOME O	CASTES	jk	47.2	44.43	1.55	9.46				
		.976		=1	6.42+5.93x	Ĩ							
Kayastha-	1				k—I 44.43	53.6	9.46	1.23		.491			
Miean of	Miean of	Std. dev	. Std. c	lev. Correlation			=82.0	4-6.40x					
Regression	า				4.2								
х	Y	Х	Y	co-efficient	Kayastha	- 2							
equation													
bc	13.3	34	9.89	8.16	bc	11.8	27.15	2.05	8.12	.9428			
.6185	Y =27.214	4+5.102x			-16.848 +3.728x								
cd	34	44.2	8.16	1.26	cd	27.15	37.5	8.12	9.37				
0.63		=78.2+(-′	1)x		.990		6.26 +1.15	ĸ					
de	44.13	13.3	1.37	9.89	de	37.5	11.5	9.37	2.05				
.188		=7.2	28+1.36x		.924	4.20	+2.02x						
ef	13.3	34	9.89	8.16	fg	27.15	37.5	8.12	9.27				
.618		=27	.21+5.10x		.936		6.26 +1.15	x					
fg	34	44.13	8.16	1.37	hi	40.32	43.25	10.47	,				
.645		=8	1.53-1.10x		6.10		.93	21.41	+5.41x				
hi	47.2	44.43	1.55	9.46	ij	43.25	40.32	6.10	10	0.47			
.976		=1	6.42+5.93x				.93	-28.75	+1.59x				

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jk	40.32	43.25	10.4	7 6.10		a1,a2	15.72	6.42	.810925	.355106	
.93	21.	41 +5.41x				.645914	y= 6.874	64			
kl	43.25	44.72	6.10	10.94	Ļ						
	.859	-21.99 +1.	54x							- 2.828	3447x
4.3						a2,b	6.43	20	0.27	3.5510	06
Brahmin	- 1					0.26096	.980918	s y= 156	3489		
ab	33		12.47	1.58	1.42						
0.95		-15.90+0	0.86x							+ .072	0856x
cd	31.5	41.	.77	4.12	8.07	b,c	20.27	12.43	0.26096 3.	55106	
. 803		-7.79+1.	57x			.9810077	y= -14.62	2543			
e-f	12.47	31.5	1.42	4.12							
.964		28.02+2	.78x							+13.3	47x
gh	41.77	44.57	8.07	1.52		c,d	12.43	14.96	.355106	.867409	.666172
877		-24.75+′	1.65x				y= -5.267	723			
ij	43.42	44.57	1.13	1.52	.961						
		-11.49+ ⁻	1.29x			_				+16.2	729x
<u>4.4</u>						d,e	14.96	28.63	.86	7409	5.55230
Brahmin	- 2						.960979	y= -63.	38807		
		н.									
ab	32.6	12.3	5 1.50	3.04						+6.15	125x
0.99		5.78+2.0)1x			e,f	28.63	15.44	5.5523	0 .534217	,
cd	31.775	40.8	1.323	7.87		.534217	y= 14611	75			
0.89		23.93+5	.30x								
e-f	12.35	31.775	3.04	1.32	. 978					+.289	294x
		-20.79+4	4.25x			f,g	15.44	20.49	.300666		.353412
gh	40.3	43.27	7.87	1.27			.973095	y= 2.82	7608		
.731		- 5.09+1	.18x								
ij	42.55	43.27	0.05	1.27						+1.14	39x
.126		28.15+3	.55x			g,h	20.49	27.53	.353412 .8	74128	
kl	42.55	50.8	0.455	7.87		.65485	y= -5.656	610			
.306		28.24+5	.30x								
										+1.61	962x
Bengali:-						h,i	27.53	25.65	.874128 .4	73814	
S-1, (Sports 1-short r un)					.657933 y= 15.83231						

					a2,b	a2,b 6.64		20.69 .4152107		4.03608
				+.356617x	.9929359	y= 14.281	132			
i.j	25.65	27.68	.473814	.942337						
.7233121	y= -9.46	6422							+.9651	632x
					b,c	20.69	11.7	.403597 .4	19524	.998090
				+1.448118x		y= 9.765	252			
j,k	27.7	25.52	.942337	.396988						
.7019981	y= 17.38	695						+1.03147x	c	
					c,d	11.7	15.5	9 .419524 .37	75324	
				+0.293824x	.997955	y= 5.153	485			
k,l	25.52	15.9	.396988	1.31453						
.802761	y= -51.9	3197							+.8720	15x
					d,e	15.59	11.62	.375326	.4	4118159
				+.265799x		.997507	y= -5.44′	153		
l,m	15.9	13.23	1.31453	6.94319						
.494287	y= 17.40	752	1.1						+1.094	39x
					e,f	11.62	15.68	4.11825		4.16654
				262737x		.997476	y= 5.442	599		
m,n	13.23	15.24	.694319	.3747069						
.454186	y= 15.2 ⁻	1572							+1.094	454x
					f,g	15.68	20.75	.4166341	.4	4694786
				+.245113x		.9966539	y= 3.140	326		
n,h	15.54	154.31 .3	374699 1.′	157179 .995475						
	y= 159	.0874							+1.123	2066x
					g,h	20.75	25.69	.4694786		4805291
				.3074248x		0.99042	y= 4.654	995		
h,w	154.31	52.4	1.15515 4	4.34051						
.2044297	y= 170.9	93320							+1.013	755x
								n the previou	-	
768157x								softwoorga		
	rts2-High						-	that, the ra		
a1,a2 14.65 6.64 .427200 .415210 .9978941 or gans for different castes are almost same. From th										
	y= -7.56	805			can conclude that the cell generation time of cells of any					
					organ ar	e same, oth	erwise it i	s not possible		
				+.9698328x						

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We see from the tables of sports there are close relations between the lengths of different parts of the bodies of the same type of experts in sports. This is applicable to other type of experts also.

Seeing the relations of the lengths of different parts of one child's body it can be predicted in which line the child will show his or her maximum capacity.

REFERENCE

1. Heinmets, F. – Mathematical Modeling in Simulation Process(1970)

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