

UNIVERSITY OF SARAJEVO FACULTY OF ELECTRICAL ENGINEERING

MODELING OF FUZZY SYSTEM FOR PROSTATE CANCER RISK ESTIMATION USING COEVOLUTIONARY ALGORITHM

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Project timetable:

- O 2004-2006/Jan-Dec ,Sarajevo
- Theoretical research :
- Field: Artificial Intelligence
- Field: Bioinformatics (Genomics&Proteomics)
- Field: Biomedical (Prostate&Cancer)

Practical implementation :

- MATLAB-Fuzzy Logic Toolbox
- MATLAB-Genetic Algorithm
 Toolbox

In this project, an attempt was made to explain modeling of fuzzy system for prostate cancer risk estimation using optimization based on genetic algorithm. Prostate cancer risk estimation is a real and complex problem that can be solved by building fuzzy system. Fuzzy modeling in this case is long and hard process, but it is possible to apply evolution algorithm to evolve fuzzy system and speed up fuzzy modeling process. The aim of this project is to present advantages and disadvantages of such modeling process and to discuss possibility of its practical application. As the result of this research, example of fuzzy system, which is obtained through evolution process, is presented.

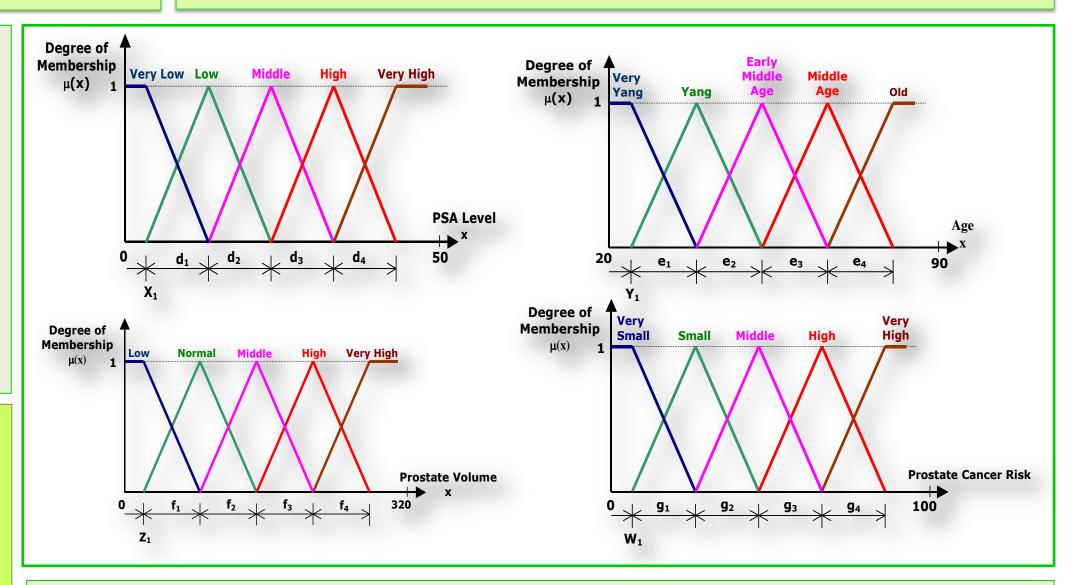
Key words — fuzzy, genetic algorithm, coevolution, prostate cancer, modeling.

Fuzzy systems – fuzziness

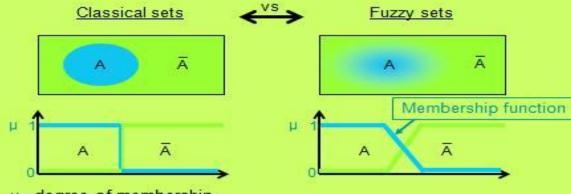
 Modeling of fuzzy systems is based on imprecise concepts and imprecise dependencies among them.

 Example: If Temperature is <u>High</u> and Sunshine is <u>Strong</u> then Vacation Conditions are <u>Perfect</u>.

 For every real value (e.g. T = 20°C) is possible to define our estimation that it fits into some concept (e.g. Temperature is <u>High</u>), and that is called *fuzziness*. So, *fuzziness* is a measure of our believe that some value (instance) fits into semantically ideal concept. Fuzzy system for prostate cancer risk estimation has 3 input fuzzy variables: PSA level, Age and Prostate Volume (PV), 1 output fuzzy variable: Prostate Cancer Risk (PCR) and collection of fuzzy rules. For presenting fuzzy variables, triangle, L and Γ types of fuzzy sets are used. They are choiced for simpler computations. Each variable is defined with 5 fuzzy sets, as shown on figures. Parameters X₁, d₁, d₂, d₃, d₄, Y₁, e₁, e₂, e₃, e₄, Z₁, f₁, f₂, f₃, f₄, W₁, g₁, g₂, g₃ and g₄ define the look of fuzzy variables.



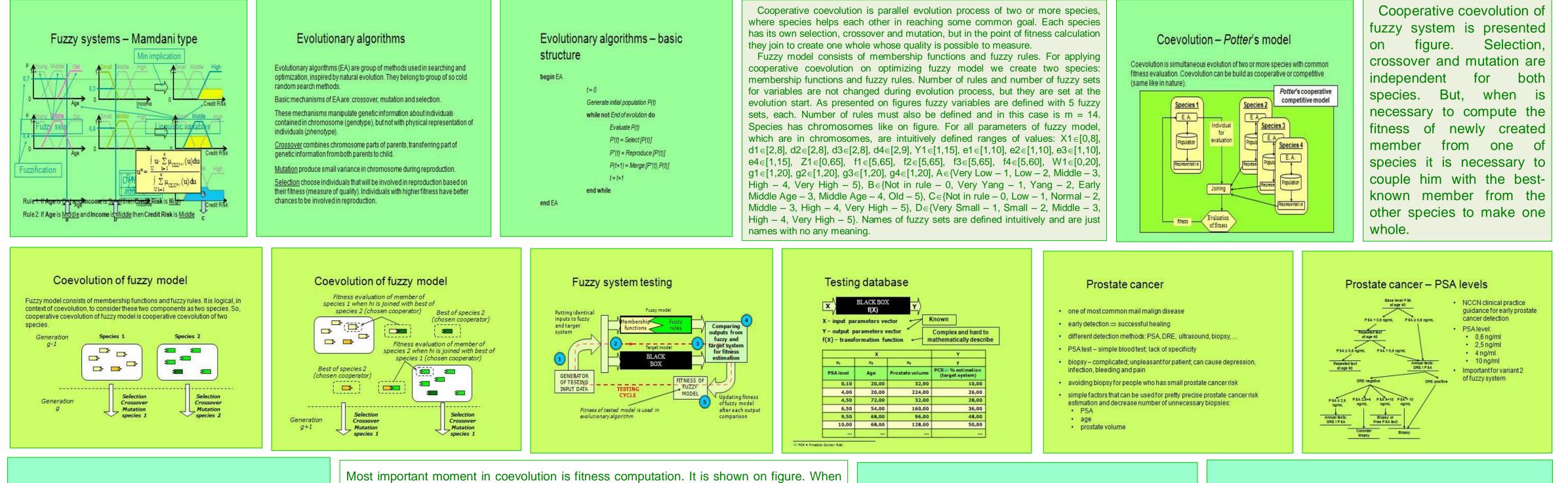
Fuzzy systems – fuzzy sets



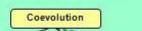
µ - degree of membership Fuzzy sets are generalized version of classical mathematical concept of sets.

Operators in fuzzy t membership gradati	heory are not uniquely defined because of on.
<u>∧ operator</u> ("AND" o following ways:	perator) is most commonly defined in one of the
Minimum	$\mu_A(x) \wedge \mu_B(x) = \min(\mu_A(x) + \mu_B(x))$
Product	$\mu_A(x) \land \mu_B(x) = \mu_A(x) \cdot \mu_B(x)$
Bounded sum	$\mu_A(x) \land \mu_B(x) = \max(0, \mu_A(x) + \mu_B(x) - 1)$
following ways:	perator) is most commonly defined in one of the
Maximum	$\mu_{A}(x) \lor \mu_{B}(x) = \max(\mu_{A}(x) + \mu_{B}(x))$
Product	$\mu_{A}(x) \lor \mu_{B}(x) = \mu_{A}(x) + \mu_{B}(x) - \mu_{A}(x) \cdot \mu_{B}(x)$

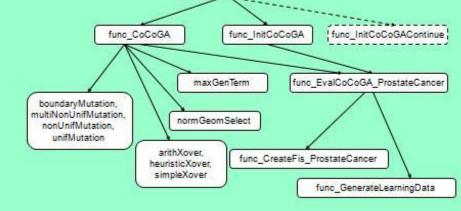
<u>~ operator</u> ("NOT" operator) is defined as: $\mu \sim A(x) = 1 - \mu A(x)$ Prostate cancer risk estimation can be based on three parameters: PSA (Prostate Specific Antigen) level, patient age and prostate volume (PV). PSA is a protein produced by the prostate that may be found in an increased amount in the blood of men who have prostate cancer, benign prostatic hyperplasia, or an infection of the prostate gland. A blood sample is measured in an assay and the amount of PSA is reported as [ng/ml]. Older patients have higher risk of prostate cancer, but also have higher prostate volume for normal PSA concentration. Prostate volume is determined by ultrasound imaging and is measured in [ml]. Many research projects are made to relate these parameters with prostate cancer risk estimation. Criteria for measuring the quality of one fuzzy system are based on the results of these projects.



Mechanisms developed in Matlab



two members, each from one of two species (the one whose fitness we compute and the chosen cooperator – best in other species) are coupled, they represent one fuzzy system. That fuzzy system we can test using data from the database of known diagnoses. At the



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first (step 1) we generate matrix of inputs (input parameters for PCR estimation). Sequentially (one by one) set of input values is put on input of fuzzy system (step 2) and fuzzy output that represents PCR estimation is computed (step 4). That is repeated for all data in input matrix and matrix of corresponding outputs is created. At the same time we put same inputs at the entrance to the target system (step 3) and take outputs (step 5) – this is not actually performed because at the start, in our database, we have these outputs. Then in step 6, corresponding outputs from both systems are compared and percentage of approximately equal answers (PCR estimations) is computed. That percentage is fitness of that fuzzy system. Actually it is the fitness of the member from one of species, since in this step fitness of chosen cooperator is not changed. All mechanisms for evolving fuzzy system including coevolution algorithm are implemented as Matlab functions.

	Fuzzy system type	Mamdani
Other parameters of fuzzy system	And operator	Min
011022y system	Or operator	Max
	Implication method	Min
	Aggregation method	Max
	Defuzzification method	Centroid

Model of fuzzy system - variant 1

Value

Parameter

	PS	ALe	vel				Age			P	rost	ate V	nulo	e	Cancer Risk Estim.				n.
Xs	d:	d2	d,	d4	Yi	e	e2	e,	e4	Zi	f ₁	f2	fa	f4	W:	9:	92	92	94
			e1			111		Rul	e2		100	·		Rule 6					
		- KUI	E 1																

	Fuzzy system type	Mamdani
ther parameters of fuzzy system	And operator	Min
offuzzy system	Or operator	Max
	Implication	Min
	Aggregation method	Max
	Defuzzification method	Centroid

Model of fuzzy system - variant 2

Value

Chron	nosomes	of species
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	PS	Alev	rel			Age	8	Prostate Volum			ne	Pros	state (Cancer	RiskE	stima	tion
8:	b,	C:	d,	e,	81	b2	C2	83	b ₂	C3	d,	84	b.	C4	d.	e4	f.
			20.—23 			Chro	moso	me of I	Fuzzy	Rules	specie	s _			a. 36		
		Pul	a 1			Chro				Rules	The second	9		Pul	- R		-
		Rul	e 1			Chro		me of I Rule 2		Rules	specie	8		Rul	e 8		

						I	Mem	bersł	hip fu	nction	s chro	mos	ome					
Coevolution result		PS	A Le	/el			1	Age				PV				I	PCR	
Chromosome of best known member of Membership Functions species - variant 1 PSA level Age Prostste Volume PCR estimation Xi di da da di di <thdi< th=""> di di</thdi<>	X ₁	d ₁	d ₂	d ₃	d ₄	Y ₁	e ₁	e ₂	e ₃ e	e ₄ Z ₁	f ₁	f ₂	f ₃	f ₄	W_1	g ₁	g ₂ g ₃	g 4
Chromosome of best known member of Fuzzy Rules species - variant 1. Rule 1 Rule 2 Rule 5 A1 B1 C1 D1 A2 B2 C2 D2 A4 B6 C6 D6 -5 4 -2 2 4 5 3 1 2 -2 0 2																		
Chromosome of best known member of Membership Functions species - variant 2 PSA level Age Prostate Volume PCR estimation								Fuzz	zy rule	es chro	moso	me						
as bs cs ds es as bs cs ds es Fa 3.9776/3920411399/S202041399/S2020/S9990.2429/S94274 - <th></th> <th>I</th> <th>Rule</th> <th>1</th> <th></th> <th colspan="4">Rule2</th> <th></th> <th></th> <th></th> <th></th> <th></th> <th colspan="4">Rule14</th>		I	Rule	1		Rule2									Rule14			
Rule1 Rule2 Rule8 A1 B1 C1 D1 A2 B2 C2 D2 A4 B2 C4 D4 -2 0 0 1 5 3 1 -2 3 3 0 2	A ₁	В	1	C ₁	D_1	A ₂	B ₂	С	2 C	D ₂					A ₁₄	B ₁	4 C ₁₄	D ₁₄

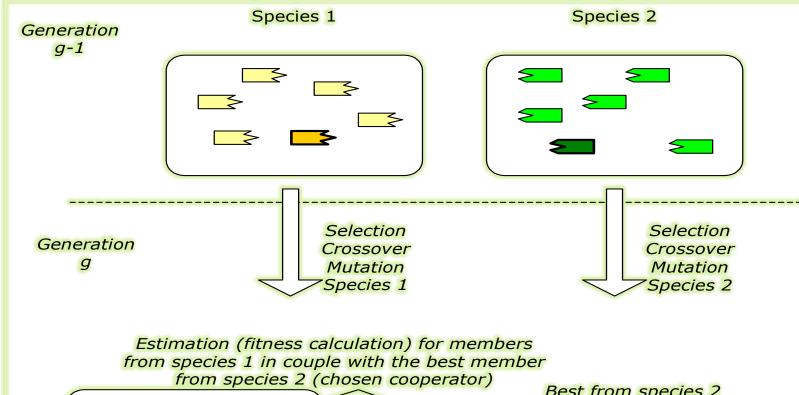
Fuzz	Final fuzzy system – variant 1	Fuzzy	Fii
No	Rule	No	
		1.	If (PSAL
1.	If (<u>PSALevel</u> is not VeryHigh) and (<u>Age</u> is MiddleAge) and (<u>ProstateVolume</u> is not Normal) then (<u>ProstateCancerRisk</u> is Small)	2.	If (PSA
2.	If (<u>PSALevel</u> is High) and (<u>Age</u> is Old) and (<u>ProstateVolume</u> is Middle) then (<u>ProstateCancerRisk</u> is Small)	3.	If (PSA
3.	If (<u>PSALevel</u> is VeryLow) and (<u>Age</u> is not Young) and (<u>ProstateVolume</u> is not Small) then (ProstateCancerRiskis VerySmall)	4.	(Prostat
4.	If (PSALevel is High) then (ProstateCancerRisk is not Middle)	12	(Prostat
5.	If (<u>PSALevel</u> is VeryLow) and (<u>Age</u> is VeryYoung) and (<u>ProstateVolume</u> is High) then (ProstateCancerRiskis Small)	5.	If (<u>PSAL</u> not Nor
6.	If (<u>PSALevel</u> is Low) and (<u>Age</u> is not Young) then (<u>ProstateCancerRisk</u> is <u>Small</u>)	6.	If (PSAL Decrea
7.	If (<u>PSALevel</u> is Middle) then (<u>ProstateCancerRisk</u> is Middle)	7.	If (PSAL
8.	If (PSALevel is High) then (ProstateCancerRisk is High)	8.	If (PSAL
9.	If (PSALevel is VeryHigh) then (ProstateCancerRiskis VeryHigh)	d.	Middle

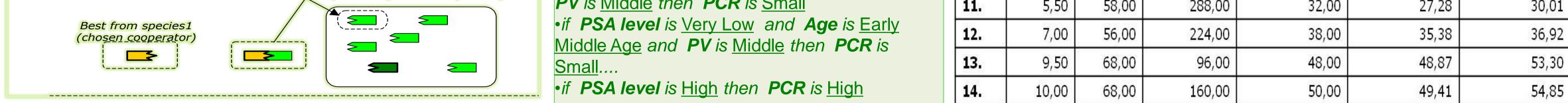
t 2	Fuzz	Final fuzzy system – variant 2
	No	Rule
<u>k</u> is Mali)	9,	If (PSALevel is About"0") then (ProstateCancerRisk is Small)
nd (<u>ProstateVolume</u> is	10.	If (PSALevel is About"0.6") and (Age is About"60") then (ProstateCancerRisk is Small)
not Decreased) then	11.	If (P <u>SALevel</u> is About"2.5") and (<u>Age</u> is About"60") and (<u>ProstateVolume</u> is Normal) then (<u>ProstateCancerRisk</u> is Middle)
s not Normal) then	12.	If (P <u>SALevel</u> is About"2.5") and (<u>Age</u> is UpTo"40") and (<u>ProstateVolume</u> is Normal) then (<u>ProstateCancerRisk</u> is Middle)
and (<u>ProstateVolume</u> is	13.	If (P <u>5ALevel</u> is About"4") and (<u>Age</u> is About"60") and (<u>ProstateVolume</u> is Increased) then (<u>ProstateCancerRisk</u> is Middle)
(ProstateVolume is not	14.	If (PSALevel is About"4") and (Age is UpTo"40") then (ProstateCancerRisk is High)
and (ProstateVolume is	15.	If (PSALevel is Over"10") then (ProstateCancerRisk is VeryHigh)
(ProstateCancerRisk is		

Example model has accuracy of approximately 70%, when tested with database of 3375 diagnoses used during coevolution, but is possible to create model with better results. That can be achieved by performing longer coevolution, for what is necessary more time and computer resources. Also is possible to evolve more structure elements of the model during coevolution, such as: type of fuzzy sets used, number of fuzzy sets for one fuzzy variable, number of fuzzy rules, etc. All that should give better final fuzzy system, but it complicates coevolution process and need more time and computer resources. Very important is quality of the created fuzzy model. That quality depends primarily on the size of the database with diagnoses and the accuracy of the diagnoses. It is possible to involve more diagnostic parameters to improve the accuracy of diagnoses.

Parameter

This shows that application of cooperative coevolution on optimizing fuzzy model for this concrete problem gives good results. It also shows possibility of applying similar approach on solving any problem that we see as black box with known inputs and known outputs.





COEVOLUTION RESULTS

al fuzzy system – varian

is not About"0.6") then (ProstateCancer is Over"10") and (Age is Over"80"

l is Over"10") and (ProstateVolume ncerRisk*is* Small)

is About"4") and (<u>ProstateVolume</u> <u>cerRisk</u>is High)

is Over"10") and (<u>Aqe</u> is not Over"8 then (ProstateCancerRiskis Small)

al is About"2.5") and (Age is Over"80") a) then (<u>ProstateCancerRisk</u> is Small)

is About"2.5") and (<u>Age</u> is not Over"80)) then (<u>ProstateCancerRisk</u>is Middle)

s About"2.5") and (Age is Over"80") to

• During experiments some elements are manually changed, like size of population for both species, size of database with known diagnoses, used for fitness computation, number of rules, etc., to try to influence evolution process and gain better result. On figures 8 to 11 is presented an example of membership functions after evolution process of 150 generations, where size of database with diagnoses used was 3375 and number of rules was 14.

Set of fuzzy rules is: •*if* **PSA level** *is* <u>Very Low</u> *and* **Age** *is* <u>Yang</u> *and* **PV** *is* <u>Middle</u> *then* **PCR** *is* <u>Small</u> •*if* **PSA level** *is* <u>Very Low</u> *and* **Age** *is* <u>Early</u> Middle Age and **PV** *is* Middle *then* **PCP** *is*

No	PSA Level	Age	Prostate Volume	PCR ^[1] estimation (target syst.)	PCR ^[1] estimation (<i>fuzzy</i> syst.1)	PCR ^[1] estimation (<i>fuzzy</i> syst.2)
1.	0,10	20,00	32,90	10,00	11,70	7,81
2.	0,50	20,00	37,15	10,00	11,94	8,00
3.	0,50	66,00	32,00	10,00	12,34	7,95
4.	1,00	20,00	40,30	10,00	12,50	8,00
5.	1,50	74,00	40,00	10,00	13,32	15,10
6.	2,50	64,00	320,00	20,00	15,84	14,92
7.	3,00	35,00	320,00	22,00	17,35	19,81
8.	3,00	60,00	43,90	10,00	17,35	7,76
9.	3,00	60,00	64,00	22,00	17,35	21,25
10.	4,00	20,00	256,00	26,00	20,70	31,44
11.	5,50	58,00	288,00	32,00	27,28	30,01
12.	7,00	56,00	224,00	38,00	35,38	36,92