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Application of soft sets to diagnose the prostate cancer risk

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Abstract

In recent years the artificial intelligence has been developed rapidly since it can be applied easily to several areas like medical diagnosis, engineering and economics, among others. In this study we have devised a soft expert system (SES) as a prediction system for prostate cancer by using the prostate specific antigen (PSA), prostate volume (PV) and age factors of patients based on fuzzy sets and soft sets and have calculated the patients' prostate cancer risk. Our data set has been provided by the Department of Urology, Meram Medical Faculty in Necmettin Erbakan University, Konya, Turkey.

Keywords: fuzzy set; soft set; prostate cancer; soft expert system

1 Introduction

In recent years vague concepts have been used in different areas such as medical applications, pharmacology, economics and engineering since the classical mathematics methods are inadequate to solve many complex problems in these areas. Traditionally mathematics uses a crisp (well-defined) property P(x), *i.e.*, properties that are either true or false. Each property defines a set: $\{x: x \text{ has a property } P\}$ [1].

The most successful theoretical approach to vagueness is undoubtedly fuzzy set theory introduced by Zadeh [2]. The theory is used commonly in different areas as engineering, medicine and economics, among others. The fuzzy set theory is based on the fuzzy membership function $\mu: X \to [0,1]$. By the fuzzy membership function, we can determine the membership grade of an element with respect to a set. A fuzzy set F is described by its membership function μ_F . The fuzzy set theory has become very popular and has been used to solve problems in different areas. But there exists a difficulty: how to set the membership function in each particular case. The reason for these difficulties is, possibly, the inadequacy of a parametrization tool of the theory [3]. Soft set theory was initiated by Molodtsov [3] as a new method for vagueness. Molodtsov showed in his paper that the theory can be applied to several areas successfully; for example, the smoothness of functions, game theory, Riemann-integration, Perron-integration, etc. He also showed that soft set theory is free from the parametrization inadequacy syndrome of other theories developed for vagueness. A soft set can be represented by Boolean-valued information system, and so it can be used to represent a dataset. Also, the hybrid models of the vague sets take attention of researchers. Maji et al. [4] defined a hybrid model called fuzzy soft sets. This new model is a combination of fuzzy and soft sets and is a generalization of soft sets. Irfan Ali and Shabir [5] developed the theory. To address decision making problems based on



fuzzy soft sets, Feng *et al.* introduced the concept of level soft sets of fuzzy soft sets and initiated an adjustable decision-making scheme using fuzzy soft sets [6]. Feng *et al.* [7] first considered the combination of soft sets, fuzzy sets and rough sets. Using soft sets as the granulation structures, Feng *et al.* [8] defined soft approximation spaces, soft rough approximations and soft rough sets, which are generalizations of Pawlak's rough set model based on soft sets. It has been proven that in some cases Feng's soft rough set model could provide better approximations than classical rough sets. Simsekler (Dizman) and Yuksel [9] contributed to fuzzy soft topological structures.

Prostate cancer is the second most common cause of cancer death among men in most industrialized countries, and it depends on various factors such as family cancer history, age, ethnic background and the level of prostate specific antigen (PSA) in blood. The level of PSA in blood is very important method to an initial diagnosis for patients [10-12]. However the level of PSA in blood can be increased by inflammation of prostate and benign prostate hyperplasia (BPH). For this reason, it is difficult to differentiate it from benign prostate hyperplasia (BPH). The definitive diagnose of the prostate cancer is possible with prostate biopsy. The results of PSA test, rectal examination and transrectal findings help the doctor to decide whether biopsy is necessary or not [1, 13, 14]. However the patients with low cancer risk have to avoid this process due to possible complications and its high cost. Because of this reason, before agreeing to biopsy, the patients with low cancer risk can be determined. There are several research works in the area of the prostate cancer prognosis or diagnosis. One of them is FES which is a rule-based fuzzy expert system using the laboratory data PSA, PV and age of the patient and it aims to help to an expert-doctor to determine the necessity of biopsy and the risk factor [15]. Benecchi [16] developed a neuro-fuzzy system by using both serum data (total prostate specific antigen and free prostate specific antigen) and clinical data (age of patients) to enhance the performance of tPSA (total prostate specific antigen) to distinguish prostate cancer. Keles et al. [17] built a neuro-fuzzy classifier to be used in the diagnosis of prostate cancer and BPH diseases. Since the symptoms of these two illnesses are very close to each other, the differentiation between them is an important problem. Saritas et al. [18] have devised an artificial neural network that provides a prognostic result indicating whether patients have cancer or not by using their free prostate specific antigen, total prostate specific antigen and age data.

In this study we aim to discuss how soft set theory can be used for developing knowledge-based system in medicine and devise a prediction system named soft expert system (SES) by using the PSA, PV and age data of patients based on fuzzy sets and soft sets and calculate the patients prostate cancer risk. It is a rule-based system, and according to the rules, we determine the risk of prostate cancer. Our aim is to help the doctor to determine whether the patient needs biopsy or not.

2 Preliminaries

Definition 2.1 [2] A fuzzy set A in U is a set of ordered pairs:

 $A = \{(x, \mu_A(x)) : x \in U\}$, where $\mu_A : U \longrightarrow [0,1] = I$ is a mapping and $\mu_A(x)$ (or A(x)) states the grade of belonging of x in A. The family of all fuzzy sets in U is denoted by I^U .

A fuzzy set can be related to a family of crisp sets through the notion of an α -level set. The α -level set of a fuzzy set F is defined by

$$F(\alpha) = \{x \in U : \mu_F(x) \ge \alpha\},\$$

where $\alpha \in [0,1]$.

Definition 2.2 [3] Let $A \subseteq E$. A pair (F,A) is called a soft set over U, where F is a mapping given by $F: A \to P(U)$, where E is the set of parameters. In other words, the soft set is a parametrized family of the subsets of U. Every set F(e), $e \in E$ from this family may be considered as the set of e-elements of the soft set (F,E), or the set of e-approximate elements of the soft set.

Example 2.1 Mr. X and Miss Y are going to marry and they want to rent a wedding room. The soft set (F,E) describes the 'capacity of the wedding room'. Let $U = \{u_1, u_2, u_3, u_4, u_5, u_6\}$ be the wedding rooms under consideration, and $E = \{e_1, e_2, e_3, e_4, e_5\}$ be the parameter set

$$F(e_1) = \{u_2, u_4\},$$

$$F(e_2) = \{u_1, u_3, u_4\},$$

$$F(e_3) = \emptyset,$$

$$F(e_4) = \{u_1, u_3, u_5\},$$

$$F(e_5) = \{u_1, u_6\}.$$

The soft set (F, E) is as follows:

$$(F,E) = \{e_1 = \{u_2, u_4\}, e_2 = \{u_1, u_3, u_4\}, e_3 = \emptyset, e_4 = \{u_1, u_3, u_5\}, e_5 = \{u_1, u_6\}\}.$$

The tabular presentation of (F, E) is shown in Table 1.

Definition 2.3 [7] Let (F,A) and (G,B) be two soft sets over U. (F,A) is called a soft subset of (G,B) denoted by $(F,A)\subseteq (G,B)$ if $A\subseteq B$ and for every $a\in A$, $F(a)\subseteq G(a)$. Two soft sets (F,A) and (G,B) over U are said to be equal, denoted by (F,A)=(G,B) if $(F,A)\subseteq (G,B)$ and $(G,B)\subseteq (F,A)$.

Definition 2.4 [19] A soft set (F,A) over U is said to be a NULL soft set denoted by Φ if $\forall e \in A, F(e) = \phi$.

Table 1 Tabular presentation of the soft set

U	e_1	e_2	e_3	e ₄	e ₅
<i>u</i> ₁	0	1	0	1	1
u_2	1	0	0	0	0
U ₃	0	1	0	1	0
<i>U</i> 4	1	1	0	0	0
U_5	0	0	0	1	0
И6	0	0	0	0	1

Definition 2.5 [19] A soft set (F,A) over U is said to be an absolute soft set denoted by \tilde{A} if $\forall e \in A, F(e) = U$.

Definition 2.6 [19] If (F,A) and (G,B) are two soft sets, then (F,A) and (G,B) denoted by $(F,A) \wedge (G,B)$ is defined by $(F,A) \wedge (G,B) = (H,A \times B)$, where $H(\alpha,\beta) = F(\alpha) \cap G(\beta)$, $\forall (\alpha,\beta) \in A \times B$.

Definition 2.7 [19] Let (F,A) and (G,B) be two soft sets over U. The union of (F,A) and (G,B) denoted by $(F,A) \tilde{\cup} (G,B)$ is defined as the soft set (H,C), where $C = A \cup B$, and $\forall e \in C$,

$$H(e) = \begin{cases} F(e) & \text{if } e \in A - B, \\ G(e) & \text{if } e \in B - A, \\ F(e) \cup G(e) & \text{if } e \in A \cap B. \end{cases}$$

Definition 2.8 [20] Let (F,A) and (G,B) be two soft sets over U.

1. The extended intersection of (F,A) and (G,B) denoted by $(F,A) \sqcap_{\wp} (G,B)$ is defined as the soft set (H,C), where $C = A \cup B$, and for all $e \in C$,

$$H(e) = \begin{cases} F(e) & \text{if } e \in A - B, \\ G(e) & \text{if } e \in B - A, \\ F(e) \cap G(e) & \text{if } e \in A \cap B. \end{cases}$$

2. The restricted intersection of (F,A) and (G,B) denoted by $(F,A) \cap (G,B)$ is defined as the soft set (H,C), where $C = A \cap B$, and for every $c \in C$, $H(c) = F(c) \cap G(c)$.

Theorem 2.1 [21] Every fuzzy set can be considered as a soft set.

Definition 2.9 [22] An information system is a 4-tuple S = (U, A, V, f), where $U = \{u_1, u_2, \dots, u_{|U|}\}$ is a non-empty finite set of objects, $A = \{a_1, a_2, \dots, a_{|A|}\}$ is a non-empty finite set of attributes, $V = \bigcup_{a \in A} V_a$, V_a is the domain of attribute $a, f : U \times A \to V$ is an information function, such that $f(u, a) \in V_a$ for every $(u, a) \in U \times A$, called information (knowledge) function. An information system can be expressed in terms of an information table (see Table 2). In an information system S = (U, A, V, f), if $V_a = \{0, 1\}$, for every $a \in A$, then S is called a Boolean-valued information system.

Proposition 2.2 [22] If (F,E) is a soft set over the universe U, then (F,E) is a Boolean-valued information system.

Table 2 An information system

U	<i>a</i> ₁	a ₂	•••	a _k	•••	$a_{ A }$
<i>u</i> ₁	f(u ₁ , a ₁)	f(u ₁ , a ₂)		$f(u_1, a_k)$		$f(u_1,a_{ A })$
u_2	$f(u_2, a_1)$	$f(u_2, a_2)$		$f(u_2, a_k)$		$f(u_2, a_{ A })$
<i>U</i> ₃	$f(u_3, a_1)$	$f(u_3, a_2)$		$f(u_3,a_k)$		$f(u_3,a_{ A })$
:	:	:	٠.	:	٠.	:
$u_{ U }$	$f(u_{ U },a_1)$	$f(u_{ U },a_2)$		$f(u_{ U },a_k)$		$f(u_{ U },a_{ A })$

The reduction of parameters of soft sets has taken attention of several researchers. Kong [23] gave an algorithm for the normal parameter reduction of soft sets in 2008. In 2011 Ma [24] gave a new algorithm for the normal parameter reduction of soft sets and compared this new method with Kong's method. These two algorithms calculate the same reduction, but Kong's method is more difficult and complex. Ma gave a new algorithm that is more understandable and easier to avoid the difficulty of Kong's algorithm.

3 Soft expert system

The prostate data set was provided by the Department of Urology, Meram Medical Faculty in Necmettin Erbakan University, Konya, Turkey. The true data set contains the PSA, PV and age data of 78 patients (see Table 3). For the design process PSA, age and PV were used as input values and prostate cancer risk was used as an output.

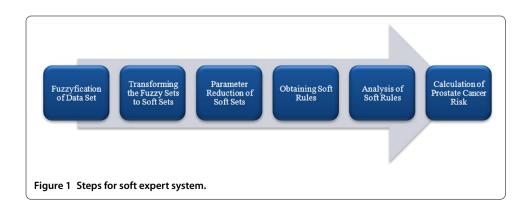
The steps for our designed system are as shown in Figure 1.

3.1 First step: fuzzyfication of data set

The data set used in this work is 78 patients who appealed to Meram Medical Faculty urology department for the prostate complaint. The data set is not convenient for applying to soft sets directly (see Table 3). For this reason, we first fuzzyficate the data set. For fuzzyfication of the factors, the linguistic variables are (for PSA) very low (VL), low (L), middle (M), high (H), very high (VH), (for PV) very small (VS), small (S), middle (M), big (B), very big (VB), (for age) young (Y), middle (M), old (O). Fuzzyfication of the used factors is made by the membership functions (1), (2) and (3). These formulas are determined by

Table 3 The input values of several patients

U	PSA	PV	Age
<i>U</i> ₃	100	44	58
U19	20	37	69
U ₂₅	38	36	72
U42	25	48	60
U46	4,03	60	63
U55	10	62	71
U ₆₀	31	72	79
U ₆₈	20,6	78	67
U72	8,5	82	60
u_{75}	41	79	80



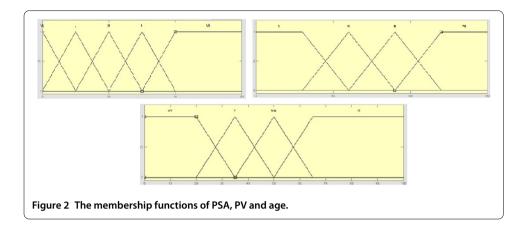


Table 4 The fuzzy membership values of factors

U	PSA	PV	Age
<i>U</i> ₃	1 VH	0.53 S, 0.47 M	0.47 M, 0.53 O
U19	0.2 VL, 0.8 L	0.77 S, 0.23 M	10
U ₂₅	0.48 L, 0.52 M	0.8 S, 0.2 M	10
U42	0.28 VL, 0.72 L	0,4 S 0,6 M	0,33 M 0,67 O
U ₄₆	0,84 VL 0,16 L	1 M	0,13 M 0,87 O
U55	0,6 VL 0,4 L	0,93 M 0,07 B	10
U ₆₀	0,41 L 0,59 M	0,6 M 0,4 B	10
U ₆₈	0,18 VL 0,82 L	0,4 M 0,6 B	10
U72	0,66 VL 0,34 L	0,27 M 0,73 B	0,33 M 0,67 O
U75	0,36 L 0,64 M	0,37 M 0,63 B	1 O

the expert doctor and literature.

$$PSA(a) = \begin{cases} \mu_a & \text{if } 0 < a < 100, \\ 1 & \text{if } 100 \le a, \end{cases}$$
 (1)

$$Age(b) = \begin{cases} 0 & \text{if } b \le 20, \\ \mu_b & \text{if } 20 < b < 65, \\ 1 & \text{if } b \ge 65, \end{cases}$$
 (2)

$$PV(c) = \begin{cases} \mu_c & \text{if } 30 < c < 120, \\ 1 & \text{if } c \ge 120. \end{cases}$$
 (3)

We get the memberships of the input variables from the formulas (1), (2) and (3) and show them in Figure 2.

We fuzzificated all data of the patients by using these membership functions. We can see the membership functions of some patients in Table 4.

3.2 Second step: transforming the fuzzy sets to soft sets

We know that every fuzzy set can be considered as a soft set. First we choose the parameter set by using the membership functions. Hence we have numerical values for a parameter

set. Some of the soft sets obtained by the relation with fuzzy sets are as follows:

$$U = \{0 = u_1, u_2, \dots, u_{78}\}, \qquad E = \{0, 0.25, 0.5, 0.75, 1\},$$

$$(F_{M PSA}, E) = \{0 = \{u_4, u_5, u_6, u_{11}, u_{13}, u_{15}, u_{20}, u_{22}, u_{23}, u_{25}, u_{30}, u_{32}, u_{34}, u_{38}, u_{41}, u_{42}, u_{43}, u_{44}, u_{53}, u_{60}, u_{64}, u_{73}, u_{75}\},$$

$$0.25 = \{u_4, u_6, u_{11}, u_{13}, u_{15}, u_{20}, u_{22}, u_{23}, u_{25}, u_{34}, u_{38}, u_{41}, u_{44}, u_{60}, u_{64}, u_{75}\},$$

$$0.5 = \{u_4, u_{11}, u_{13}, u_{15}, u_{20}, u_{22}, u_{23}, u_{25}, u_{38}, u_{41}, u_{44}, u_{60}, u_{64}, u_{75}\},$$

$$0.75 = \{u_{13}, u_{20}, u_{23}, u_{38}, u_{41}\},$$

$$1 = \{u_{20}, u_{38}\},$$

$$U = \{u_{11}, u_{21}, u_{21}, u_{22}, u_{23}, u_{25}, u_{38}, u_{41}, u_{44}, u_{60}, u_{64}, u_{75}\},$$

$$0.185 = \{u_{11}, u_{17}, u_{35}, u_{36}, u_{45}, u_{46}, u_{49}, u_{53}, u_{55}, u_{60}, u_{65}, u_{68}, u_{72}, u_{73}, u_{75}\},$$

$$0.37 = \{u_{36}, u_{45}, u_{60}, u_{68}, u_{72}, u_{73}, u_{75}\},$$

$$0.555 = \{u_{45}, u_{68}, u_{72}, u_{75}\},$$

$$0.74 = \phi\},$$

$$U = \{u_{11}, u_{22}, \dots, u_{78}\}, \qquad E = \{0.06, 0.31, 0.56, 0.81, 0.94\},$$

$$(F_{M Age}, E) = \{0.06 = \{u_{31}, u_{81}, u_{91}, u_{222}, u_{322}, u_{331}, u_{35}, u_{422}, u_{431}, u_{441}, u_{46}, u_{48}, u_{491}, u_{522}, u_{561}, u_{581}, u_{631}, u_{661}, u_{661}, u_{661}, u_{661}, u_{662}, u_{662}, u_{663}, u_{662}, u_{663}, u_{662}, u_{663}, u_{663}, u_{664}, u_$$

3.3 Third step: parameter reduction of soft sets

In Step 2 we obtain the soft sets corresponding to each fuzzy set. Then we use the parameter reduction of soft sets given by Ma [24]. Hence we have new soft sets. Some of them are shown in the following:

$$U = \{u_1, u_2, \dots, u_{78}\}, \qquad E = \{0.25, 0.5, 0.75, 1\},$$

$$(F_{M PSA}, E) = \{0.25 = \{u_4, u_6, u_{11}, u_{13}, u_{15}, u_{20}, u_{22}, u_{23}, u_{25}, u_{34}, u_{38}, u_{41}, u_{43}, u_{44}, u_{60}, u_{64}, u_{75}\},$$

$$0.5 = \{u_4, u_{11}, u_{13}, u_{15}, u_{20}, u_{22}, u_{23}, u_{25}, u_{38}, u_{41}, u_{44}, u_{60}, u_{64}, u_{75}\},$$

$$0.75 = \{u_{13}, u_{20}, u_{23}, u_{38}, u_{41}\},$$

$$1 = \{u_{20}, u_{38}\}\},$$

$$U = \{u_1, u_2, \dots, u_{78}\}, \qquad E = \{0.185, 0.37, 0.555\},$$

$$(F_{BPV}, E) = \{0.185 = \{u_{11}, u_{17}, u_{36}, u_{45}, u_{60}, u_{68}, u_{72}, u_{73}, u_{75}\},$$

$$0.37 = \{u_{36}, u_{45}, u_{60}, u_{68}, u_{72}, u_{73}, u_{75}\},$$

$$0.555 = \{u_{45}, u_{68}, u_{72}, u_{75}\}\},$$

$$U = \{u_1, u_2, \dots, u_{78}\}, \qquad E = \{0.31, 0.56, 0.81\},$$

$$(F_{MAge}, E) = \{0.31 = \{u_3, u_{22}, u_{33}, u_{35}, u_{42}, u_{43}, u_{44}, u_{48}, u_{52}, u_{58}, u_{63}, u_{66}, u_{69}, u_{70}, u_{72}, u_{74}, u_{76}, u_{78}\},$$

$$0.56 = \{u_{43}, u_{48}, u_{52}, u_{58}, u_{63}, u_{70}, u_{74}, u_{78}\},$$

$$0.81 = \{u_{48}, u_{52}, u_{70}\}\}.$$

3.4 Fourth step: obtaining soft rules

We get the soft rules by the 'AND' operation of the soft sets we obtained in the second step, and we observe which patient provides which rule. Some of the rules we obtained are as follows:

$$F_{VL\ PSA}(0.35) \wedge F_{M\ PV}(0.25) \wedge F_{OAge}(0.59)$$

$$= \{u_7, u_{10}, u_{14}, u_{16}, u_{27}, u_{31}, u_{33}, u_{39}, u_{46}, u_{49}, u_{50}, u_{51}, u_{54}, u_{55}, u_{56}, u_{57}, u_{61}, u_{62}, u_{63}, u_{65}, u_{67}, u_{71}, u_{72}\},$$

$$F_{L\ PSA}(0.2875) \wedge F_{S\ PV}(0.275) \wedge F_{M\ Age}(0.31)$$

$$= \{u_{22}, u_{33}, u_{42}, u_{43}, u_{44}, u_{48}, u_{52}, u_{58}, u_{63}, u_{70}, u_{74}, u_{78}\},$$

$$F_{M\ PSA}(0.25) \wedge F_{M\ PV}(0.25) \wedge F_{O\ Age}(0.325) = \{u_{6}, u_{11}, u_{15}, u_{20}, u_{34}, u_{41}, u_{44}, u_{60}, u_{75}\},$$

$$F_{M\ PSA}(0.25) \wedge F_{M\ PV}(0.5) \wedge F_{O\ Age}(0.325) = \{u_{6}, u_{11}, u_{15}, u_{34}, u_{41}, u_{44}, u_{60}\},$$

$$F_{H\ PSA}(0.2225) \wedge F_{S\ PV}(0.785) \wedge F_{O\ Age}(0.59) = \{u_{8}\},$$

$$F_{VH\ PSA}(0.6875) \wedge F_{S\ PV}(0.785) \wedge F_{O\ Age}(0.59) = \{u_{8}\},$$

$$F_{VH\ PSA}(0.6875) \wedge F_{S\ PV}(0.785) \wedge F_{O\ Age}(0.59) = \{u_{8}\},$$

$$F_{VH\ PSA}(0.6875) \wedge F_{S\ PV}(0.785) \wedge F_{O\ Age}(0.59) = \{u_{8}\},$$

$$F_{VH\ PSA}(0.6875) \wedge F_{S\ PV}(0.785) \wedge F_{O\ Age}(0.59) = \{u_{8}\},$$

In this way, we obtain 400 rules. Then we eliminate some rules that have the same output (the same patient set), and hence we get 285 rules.

3.5 Fifth step: analysis of soft rules

In this step we analyze the soft rules and calculate the prostate cancer risk percentage. The patients set for each rule was obtained in the fourth step. We consider these sets and observe how many of the patients in the set have prostate cancer, then we rate the patients with prostate cancer to each patient in the set. Therefore we have the prostate cancer risk percentage for each rule. If a patient's data is convenient to more than one rule and so has more than one rate, then we accept the highest one.

Now we calculate the risk percentage of the first rule:

Rule 1:

```
F_{VL\ PSA}(0.35) \wedge F_{M\ PV}(0.25) \wedge F_{O\ Age}(0.59)
= \{u_7, u_{10}, u_{14}, u_{16}, u_{27}, u_{31}, u_{33}, u_{39}, u_{46}, u_{49}, u_{50}, u_{51}, u_{54}, u_{55}, u_{56}, u_{57}, u_{61}, u_{62}, u_{63}, u_{65}, u_{67}, u_{71}, u_{72}\}.
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There are 23 patients who have the properties stated in Rule 1. Prostate cancer is found in eight of these patients. Hence, the risk percentage for first rule is $(8 \div 23) \times 100 = 34.78$. We can easily say that the patients whose values of PSA, PV and age are convenient to the first rule have cancer risk of 34%. The values of patient u_{34} are convenient to Rule 3, Rule 4 and Rule 8. When we look at the risk percentage of these rules, we see that Rule 8 has the highest rate. Hence the risk percentage of u_{34} is 100% (the percentage of Rule 8).

The risk percentage for some rules is as follows:

- Rule 1: If a patient has $F_{VL\ PSA}(0.35)$ and $F_{M\ PV}(0.25)$ and $F_{O\ Age}(0.59)$, then the cancer risk is 28%.
- Rule 2: If a patient has $F_{L\ PSA}(0.2875)$ and $F_{S\ PV}(0.275)$ and $F_{M\ Age}(0.31)$, then the cancer risk is 34%.
- Rule 3: If a patient has $F_{M\ PSA}(0.25)$ and $F_{M\ PV}(0.25)$ and $F_{O\ Age}(0.325)$, then the cancer risk is 74%.
- Rule 4: If a patient has $F_{M\ PSA}(0.25)$ and $F_{M\ PV}(0.5)$ and $F_{O\ Age}(0.325)$, then the cancer risk is 83%.
- Rule 5: If a patient has $F_{H\ PSA}(0.2225)$ and $F_{S\ PV}(0.785)$ and $F_{O\ Age}(0.59)$, then the cancer risk is 100%.
- Rule 6: If a patient has $F_{H\ PSA}(0.2225)$ and $F_{S\ PV}(0.53)$ and $F_{O\ Age}(0.325)$, then the cancer risk is 100%.
- Rule 7: If a patient has $F_{VH\ PSA}(0.6875)$ and $F_{S\ PV}(0.785)$ and $F_{O\ Age}(0.59)$, then the cancer risk is 100%.
- Rule 8: If a patient has $F_{VH\ PSA}(1)$ and $F_{S\ PV}(0.275)$ and $F_{O\ Age}(0.59)$, then the cancer risk is 100%.

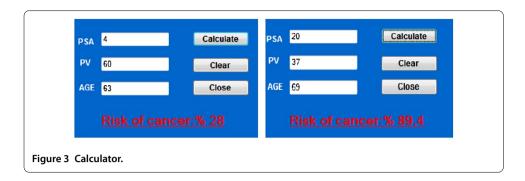
Finally, we write the soft expert system which calculates the prostate cancer risk by input variables PSA, PV and age.

3.6 Calculation of prostate cancer risk

We used MicrosoftVisual Studio 2008 and C Sharp programming language when we devised all the steps of the soft expert system. Figure 3 shows two results from the calculation system.

3.7 Conclusion

In this work we designed an expert system SES by using a soft set and it is a pioneering work for applying the soft sets to a medical diagnosis. We also used fuzzy membership functions and an algorithm to reduce the parameter set of soft sets. The expert doctor can reduce unnecessary biopsies in patients undergoing evaluation for prostate cancer by calculating the percentage of prostate cancer risk in the soft expert system. According to our devised system, if the risk percentage is bigger than 50%, then biopsy is necessary. Our data set contains 78 patients. These patients have high values of PSA, PV and age and



they are potential prostate cancer patients. For this reason, the biopsy was applied to these patients; however, after biopsy it was seen that 44 of them had cancer. When we calculated the risk percentage of these 78 patients in the soft expert system, we saw that 51 patients needed biopsy, and 27 patients who really had low cancer risk had to avoid biopsy. Our aim is to help the doctor to decide whether the patient needs biopsy or not.

Competing interests

The authors declare that they have no competing interests.

Authors' contributions

All authors contributed equally and significantly in writing this paper. All authors read and approved the final manuscript.

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