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An Evaluation of the PRE-DELIRIC (PREdiction of DELIRium in ICu patients) Delirium Prediction Model in Intensive Care Units in Turkey

PRE-DELIRIC (PREdiction of DELIRium in ICU Patients) Deliryum Öngörme Modelinin Genel Yoğun Bakım Ünitelerinde Kullanılabilirliğinin Değerlendirilmesi

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ABSTRACT Objective: The purpose of this methodological study was to perform the adaptation and validation of the "PRE-DELIRIC Score" delirium prediction model in Turkey in patients hospitalized in the intensive care unit.

Materials and Methods: The research was conducted on 172 patients treated in the intensive care units of training and research hospital between October 2019 and April 2020. The data were collected with (1) a data collection form for the participants' descriptive characteristics, (2) the PRE-DELIRIC Score, and (3) the Confusion Assessment Method for the ICU (CAM-ICU). ROC analysis and diagnostic screening tests were used to determine the cut-off point according to the groups. The sensitivity and specificity of the scores were calculated. Significance was evaluated at the $p < 0.05$ level.

Results: A statistically significant relationship was found between the cut-off point obtained for the PRE-DELIRIC Score ($\geq 7.58\%$) and the groups ($p = 0.003$). The risk of being CAM-ICU-positive was 7.404 times higher in patients with a PRE-DELIRIC score of 7.58 or more (OR: 7.404; 95% CI: 1.638-33.469). The area under the ROC curve was 64.9% (95% CI: 0.538-0.760), and the standard error was 5.6% ($p = 0.044$).

Conclusion: The PRE-DELIRIC score yielded reliable results in this study. It appears significant for patients with a likelihood of developing delirium in the intensive care unit, and its use is recommended as a functional score that is easily applied and calculated.

Keywords: Critical care, PRE-DELIRIC, delirium, model

ÖZ Amaç: Metodolojik tipteki bu çalışma, yoğun bakım ünitesinde yatan hastalarda "PRE-DELIRIC Skoru" deliryum tahmin modelinin Türkiye'ye uyarlanması ve geçerliliğinin sağlanması amacıyla yapılmıştır.

Gereç ve Yöntem: Araştırma, Ekim 2019-Nisan 2020 tarihleri arasında bir eğitim ve araştırma hastanesinin yoğun bakım ünitelerinde tedavi gören 172 hasta ile yapılmıştır. Veriler, (1) katılımcıların tanımlayıcı özelliklerine yönelik bilgi formu, (2) PRE-DELIRIC Skoru ve (3) Konfüzyon Değerlendirme Formu (CAM-ICU) ile toplanmıştır. Gruplara göre kesme noktasını saptamada ROC analizi ve tanı tarama testleri kullanıldı. Skorun duyarlılık ve özgüllük özelliği hesaplandı. Anlamlılık $p < 0.05$ düzeyinde değerlendirildi.

Bulgular: Delirium (PRE-DELIRIC) Skoru için elde edilen kesme noktası ($\geq 7,58$) ile gruplar arasında istatistiksel olarak anlamlı ilişki saptanmıştır ($p = 0,003$). Delirium (PRE-DELIRIC) Skoru 7,58 ve üzerinde olan olgularda CAM-ICU pozitif olma riski 7,404 kat fazladır (OR: 7.404; %95 GA: 1.638-33.469). ROC eğrisi altında kalan alan ise, %64.9 (%95 CI: 0.538-0.760) ve standart hata %5.6 ($p = 0.044$) olarak saptanmıştır.

Sonuç: Bu çalışmada, PRE-DELIRIC skorunun güvenilir sonuçlara sahip olduğu bulundu. Yoğun bakımlarda deliryum gelişmesi olası olan hastalar için önemli olduğu görülmekte, uygulaması ve hesaplaması kolay kullanışlı bir skor olarak kullanımı önerilmektedir.

Anahtar Kelimeler: Yoğun bakım, predeliric, deliryum, model

Introduction

Delirium is a neuropsychiatric disorder characterized by impaired consciousness, distraction, and disorganized thinking (1). It can be seen at rates of up to 80% in intensive care patients and causes increased morbidity and mortality, prolonged duration of mechanical ventilation and intensive care unit stay, and long-term cognitive impairment (2,3).

Although various organic factors can lead to delirium, there is no laboratory test available to confirm its presence. The condition can often be overlooked if attempts are made to diagnose patients with clinical data (4). The standard methods of obtaining a general medical, psychiatric and neurological history and conducting a physical examination are required for diagnosis. Final diagnosis is based on evaluating the findings obtained through interviews. Several different scales have been developed for the screening, diagnosis and grading of symptoms. Proper diagnosis requires periodic evaluation of the diagnostic criteria and knowledge of the patient's initial mental state (5). The most common methods used for the evaluation of delirium are the Confusion Assessment Method for the Intensive Care Unit (CAM-ICU) (6) and the Intensive Care Delirium Screening Checklist (7). Delirium has been detected at a rate of 30-70% in intensive care patients using these methods (8).

Appropriate interventions are extremely important in preventing delirium. According to the guidelines on pain, agitation and delirium management (9), the application of non-pharmacological methods is highly recommended for the prevention of delirium but the evidence supporting pharmacological approaches is inadequate. In addition, the implementation of such interventions is time consuming and involves a significant increase in workload. Various prediction models that can help identify high-risk patients have therefore been developed (4). The PREdiction of DELIRium in ICu patients (PRE-DELIRIC), one such model, has been validated in various intensive care units (ICUs) and reported to be useful (10). This model was developed based on the results of a systematic review in which risk factors for delirium were investigated (11). The model estimates the development of delirium within 24 hours after admission to the ICU based on a calculation involving 10 risk factors capable of causing delirium development and that are objectively and precisely defined. This model for predicting the risk of delirium (4,12,13), was also used in a recent study from Turkey but with no evaluation of the model's predictive ability for this condition (14).

The purpose of this methodological study was to evaluate the applicability of the "PRE-DELIRIC Score" delirium prediction model in general ICU patients.

- Is the PRE-DELIRIC score confidential?
- Can it be used in the intensive care unit?

Materials and Methods

The purpose of this methodological study was to perform the adaptation and validation of the PRE-DELIRIC score delirium prediction model in Turkey in patients hospitalized in the ICU.

The study was conducted with patients who were treated in the ICUs of a training and research hospital between October 2019 and April 2020.

Participants

Patients who were hospitalized and treated for more than 24 hours in the general ICU, aged 18 years or older, with no history of chronic alcoholism, dementia, or delirium, who were not pregnant or breastfeeding, who had no communication problems, with Richmond Agitation Sedation Scale (RASS) values of +4 to -2, and for whom consent to participate was obtained from a relative were included in the study. One hundred eighty-nine 189 cases were initially included, although the study was eventually conducted with 172 patients since eight cases were excluded due to dementia, three due to history of delirium, one due to history of alcoholism, and five for being aged under 18.

Study Procedure

The data were collected using a data collection form for the participants' descriptive characteristics, the PRE-DELIRIC Score, and the Confusion Assessment Method for the ICU (CAM-ICU). Data collection took place during the study period and was performed by a physician and a nurse, who were also involved as researchers. In this study, the patient who was delirium negative at admission should have been included. The general data were collected in the first 24 hours. Data on the diagnostic and clinical characteristics were obtained from the patients' relatives and patient charts. Data on the clinical course were also collected within the first 24 hours.

The data collection form for the descriptive characteristics of the subjects was created by the researchers after reviewing the relevant literature (9,15-17). The form includes 19 questions regarding socio-demographic data and clinical characteristics.

Measures

PRE-DELIRIC Score

The PRE-DELIRIC Scoring System developed by van den Boogaard et al. (2014) considers the age, APACHE II score, blood urea level, amount of morphine used, sedation use, metabolic acidosis, coma state, infection, planned/emergency intensive care admission, and the reason for hospitalization to provide a score (10). A pre-delirium score ≥ 50 is reported to be associated with a high incidence of delirium. The scoring system is used within the first 24 hours after admission to the ICU. The blood results were obtained from the patient's medical chart by the researchers and recorded in the questionnaire. The PRE-DELIRIC score was determined within the first 24 hours after admission to the ICU in this study.

Once the scale had been independently translated into Turkish by three translators consisting of an English teacher, an English language linguistic specialist, and a medical doctor proficient in English language, the translators agreed on a common text in terms of the appropriateness and comprehensibility of translations. The scale thus obtained was then translated back into English by three English teachers. No change in meaning was determined in the backtranslated scale compared to the original English document and the form was finalized after preliminary administration to five intensive care nurses and 10 patients. The PRE-DELIRIC Scoring System does not include intercultural differences since it is based on objective criteria and not on patient statements or interpretation. The risk factors including these objective data have the same meaning in all languages and cultures. Determining content validity by eliciting an expert opinion was therefore not required for the PRE-DELIRIC score.

CAM-ICU

This scale is in common use and is reported to provide the best compliance with DSM IV criteria (16). It was developed by Ely et al. (2001) (6). The Turkish language reliability and validity study was performed by Akinci et al. (2005) and the scale was found to have an acceptable level of sensitivity (65-69%), together with perfect specificity (97%) and reliability (Kappa=0.96) (18). The scale consists of four subcategories consisting of changes in the patient's state of consciousness, attention disorder, impaired thought process, and level of consciousness. Sub-categories are not taken into account and a conclusion is reached in the form

of the "presence" or "absence" of delirium according to the answers to the scale questions. All the first and second category answers must be negative, and one of the third and fourth categories must be present as a condition for the "presence" of delirium. This scale can be applied to all intensive care patients aged over 18 who are not comatose and who are able to communicate. It is recommended that the scale be completed within 24 hours admission to the ICU. A repeat evaluation is performed during the day in case of any change in the patient's condition. Otherwise evaluation once a day is appropriate. CAM-ICU was measured within 24 hours after admission to the ICU in this study.

Local ethics committee approval (Decision no: 2019-SBEK-08) was obtained from the institution where the study was conducted in addition to consent from the patients who were included in the study with the permission of the relevant institution. Written permission for the use of the PRE-DELIRIC score was obtained by e-mail from van den Boogaard.

Statistical Analysis

The NCSS (Number Cruncher Statistical System) 2007 (Kaysville, Utah, USA) software program was used for the statistical analysis. Mean, standard deviation, median, frequency, ratio, minimum and maximum values were used while evaluating the study data and the compliance of the data with a normal distribution was assessed using the Shapiro-Wilk test and graphical evaluations. "Student's t-test" was used for comparisons of normally distributed variables between the two groups and the "Mann-Whitney U" test for comparisons of variables without normal distribution. ROC analysis and diagnosis screening tests were used to determine the cut-off point according to the groups. The sensitivity and specificity characteristics of the score were calculated. Significance was evaluated at the $p < 0.05$ level.

Results

The study was completed with 172 patients. CAM-ICU measurement revealed that 90.1% (n= 155) of the patients were negative and 9.9% (n= 17) positive for delirium. The age range of the subjects was 35-94 years and the mean age was 72.94 ± 13.99 years. The mean age ($p = 0.036$) and the cerebrovascular event (CVE) incidence rate ($p = 0.036$) of the patients who were positive on CAM-ICU were statistically significantly higher than those who were negative. No

statistically significant difference was found between the groups in terms of the other variables ($p>0.05$) (Table 1).

Patients who were positive on the CAM-ICU required statistically significantly more physical restriction ($p= 0.07$) and developed more pressure ulcers ($p= 0.001$) than those who were negative. Statistically significantly larger amounts of sedatives were required for patients who were negative on the CAM-ICU than those who were positive ($p= 0.017$). A statistically significant difference was found between the groups in terms of the form of discharge from intensive care ($p= 0.047$) and those who died were usually negative while those who were transferred were usually positive. No statistically significant difference was found between the groups in terms of the other variables ($p>0.05$) (Table 2).

The 1st PRE-DELIRIC score of the patients who were negative on CAM-ICU was 3.9-11.9 with a mean value of 7.85 ± 1.91 . The 1st PRE-DELIRIC score of the patients who were positive for CAM-ICU was 6.7-14.9 with a mean value of 9.04 ± 2.09 (Figure 1). A statistically significant difference was found between the 1st PRE-DELIRIC scores of the CAM-ICU negative and positive groups ($p= 0.017$), and the scores of the positive patients were higher than the negative

patients. An increase of one unit in the PRE-DELIRIC scores increased the risk of CAM-ICU positivity 1.358 times (OR: 1.358; 95% CI: 1.047-1.761) (Table 3).

Determining the Cut-off Point for PRE-DELIRIC Scores Based on CAM-ICU Status

A statistically significant difference was found between the 1st PRE-DELIRIC scores of the CAM-ICU negative and positive patients ($p= 0.017$), with the CAM-ICU positive patients having higher scores (Table 3). Based on this significance, the cut-off point for the PRE-DELIRIC score was calculated. ROC analysis and diagnostic screening tests were used to determine this cut-off point by group. The cut-off point for the PRE-DELIRIC score was 7.58. This PRE-DELIRIC score cut-off value exhibited sensitivity of 88.24%, specificity of 49.68%, a positive predictive value of 16.13%, a negative predictive value of 97.47%, and accuracy of 53.49% (Table 4). The area under the ROC curve was 64.9% (95% CI:0.538-0.760) and the standard error was 5.6% ($p=0.044$) (Figure 2).

A statistically significant relationship was found between the cut-off point obtained for the PRE-DELIRIC score (≥ 7.58) and the groups ($p= 0.003$). The risk of CAM-ICU

Table 1. Evaluation of descriptive features according to CAM-ICU status

		Total	CAM-ICU		p
			Negative(n=155)	Positive (n=17)	
Age (years)	Min-Max (Median)	35-94 (77)	35-94 (74)	66-94 (84)	^a 0.006**
	Mean \pm SD	72.94 \pm 13.99	71.98 \pm 14.18	81.65 \pm 8.12	
Gender	Female	90 (52.3)	80 (51.6)	10 (58.8)	^c 0.572
	Male	82 (47.7)	75 (48.4)	7 (41.2)	
BMI (kg/m²)	Min-Max (Median)	17.2-40 (25)	17.2-40 (25)	19.5-28.3 (24.6)	^a 0.595
	Mean \pm SD	24.88 \pm 3.64	24.93 \pm 3.74	24,44 \pm 2,59	
Chronic disease status		149 (86.6)	133 (85.8)	16 (94.1)	^d 0.475
Chronic disease type[†]					
Hypertension		95 (55.2)	88 (56.8)	7 (41.2)	^c 0.220
Diabetes		42 (24.4)	39 (25.2)	3 (17.6)	^d 0.766
Chronic heart failure		50 (29.1)	47 (30.3)	3 (17.6)	^d 0.401
Chronic arterial failure		8 (4.7)	8 (5.2)	0 (0)	^d 1.000
COPD		24 (14.0)	22 (14.2)	2 (11.8)	^d 1.000
Asthma		2 (1.2)	1 (0.6)	1 (5.9)	^d 0.188
Alzheimer Disease		2 (1.2)	1 (0.6)	1 (5.9)	^d 0.188
Cerebrovascular Attack		14 (8.1)	10 (6.5)	4 (23.5)	^d 0.036*
Use Cigarette – Alcohol		18 (10.5)	18 (11.6)	0 (0)	^d 0.222

^aStudent t-test, ^bMann-Whitney U test, ^cPearson chi-square test, ^dFisher's Exact test, * $p<0.05$, ** $p<0.01$ [†]More than one chronic disease

		Total n (%)	CAM-ICU		p
			Negative (n=155)	Positive (n=17)	
Coming to intensive care	Emergency room	147 (85.5)	131 (84.5)	16 (94.1)	°0.473
	Clinic	25 (14.5)	24 (15.5)	1 (5.9)	
Reason for hospitalization	Surgery	19 (11.0)	18 (11.6)	1 (5.9)	°0.977
	Medical	134 (77.9)	118 (76.1)	16 (94.1)	
	Trauma	11 (6.4)	11 (7.1)	0 (0)	
	Neurosurgery	8 (4.7)	8 (5.2)	0 (0)	
Feeding type	Enteral	97 (56.4)	86 (55.5)	11 (64.7)	°0.668
	Parenteral	48 (27.9)	43 (27.7)	5 (29.4)	
	Oral	17 (9.9)	16 (10.3)	1 (5.9)	
	Enteral+Parenteral	4 (2.3)	4 (2.6)	0 (0)	
	Parenteral+Oral	6 (3.5)	6 (3.9)	0 (0)	
Physical restriction status		70 (40.7)	58 (37.4)	12 (70.6)	°0.017*
Drain-Tube-Ostomy status		117 (68.0)	104 (67.1)	13 (76.5)	°0.431
Drain-Tube-Ostomy type (n=117)	Drain	13 (11.1)	13 (12.5)	0 (0)	°0.671
	Tube	84 (71.8)	72 (69.2)	12 (92.3)	
	Ostomy	6 (5.1)	6 (5.8)	0 (0)	
	Drain+Tube	8 (6.8)	7 (6.7)	1 (7.7)	
	Tube+Ostomy	5 (4.3)	5 (4.8)	0 (0)	
	Drain+Tube+Ostomy	1 (0.9)	1 (1.0)	0 (0)	
Catheter status		171 (99.4)	154 (99.4)	17 (100)	°1.000
Catheter type (n=171)	SVC	3 (1.8)	3 (1.9)	0 (0)	°0.868
	SVC+Foley	24 (14.0)	21 (13.6)	3 (17.6)	
	PVC+Foley	100 (58.5)	91 (59.1)	9 (52.9)	
	SVC+PVC+Foley	44 (25.7)	39 (25.3)	5 (29.4)	
Pressure ulcer status		45 (26.2)	34 (21.9)	11 (64.7)	°0.001**
Pressure ulcer phase (n=45)	Phase 1	8 (17.8)	5 (14.7)	3 (27.3)	°0.540
	Phase 2	33 (73.3)	26 (76.5)	7 (63.6)	
	Phase 3	3 (6.7)	2 (5.9)	1 (9.1)	
	Phase 4	1 (2.2)	1 (2.9)	0 (0)	
MV status		68 (39.5)	64 (41.3)	4 (23.5)	°0.155
MV type (n=68)	Invasive	59 (86.8)	57 (89.1)	2 (50.0)	°0.082
	Non-Invasive	9 (13.2)	7 (10.9)	2 (50.0)	
Invasive MV time (day) (n=59)	Min-Max (Median)	1-24 (6)	1-24 (6)	3-4 (3.5)	°0.164
	Mean±SD	8.54±6.30	8.72±6.34	3.50±0.71	
Non-Invasive MV time (day) (n=9)	Min-Max (Median)	2-12 (3)	2-12 (2)	3-3 (3)	°0.533
	Mean±SD	3.78±3.23	4.00±3.70	3.00±0	
Sedation status		54 (31,4)	53 (34.2)	1 (5.9)	°0.017*
Sedation time (day) (n=54)	Min-Max (Median)	1-30 (6)	1-30 (6)	3-3 (3)	°0.332
	Mean±SD	8.78±7.62	8.89±7.65	3.00±0	

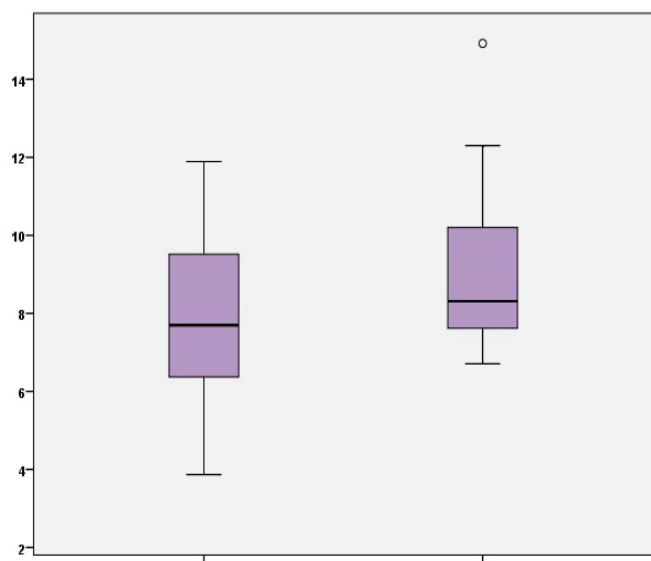
CAM-ICU 2. measurement	Negative	91 (52.9)	85 (54.8)	6 (35.3)	°0.125
	Positive	81 (47.1)	70 (45.2)	11 (64.7)	
Discharge type in intensive care	Exitus	76 (44.2)	71 (45.8)	5 (29.4)	°0.047*
	Referral	95 (55.2)	84 (54.2)	11 (64.7)	
	Discharge	1 (0,6)	0 (0)	1 (5.9)	
Length of stay in intensive care (day)	Min-Max (Median)	1-49 (5)	1-49 (5)	2-13 (4)	°0.393
	Mean±SD	6.60±6.39	6.79±6.64	4.82±2.86	
Length of stay in the hospital (day)	Min-Max (Median)	2-49 (5)	2-49 (5)	2-13 (4)	°0.218
	Mean±SD	8.56±8.71	8.89±9.04	5.53±3.45	

^bMann-Whitney U test, ^cPearson chi-square test, ^dFisher's Exact Test, ^eFisher Freeman Halton test, *p<0.05, **p<0.01

Table 3. Evaluation of Pre-Deliric Scores According to CAM-ICU Status

		Total	CAM-ICU		P
			Negative (n=155)	Positive (n=17)	
1. measurement Pre-Deliric score	Min-Max (Median)	3.9-14.9 (7.9)	3.9-11.9 (7.7)	6.7-14.9 (8.3)	°0.017*
	Mean±SD	7.97±1.95	7.85±1.91	9.04±2.09	
2.* measurement Pre-Deliric score	Min-Max (Median)	2.9-15.8 (8.3)	2.9-15.8 (8.2)	6.4-10.0 (9.2)	°0.053
	Mean±SD	8.28±2.04	8.21±2.11	8.87±1.15	

*Student t-test, °p<0.05, *24 h later

**Figure 1.** Pre-Deliric scores of the cases with negative and positive CAM-ICU in the 1st measurement

positivity was 7.404 times higher in patients with a PRE-DELIRIC score of 7.58 or more (OR:7.404; 95% CI:1.638-33.469). An increase of 5% or more in the PRE-DELIRIC score exhibited sensitivity of 100%, a specificity of 6.45%, a positive predictive value of 10.49%, a negative predictive value of 100%, and accuracy of 15.70%. An increase of 10% or more in the PRE-DELIRIC score exhibited sensitivity of 29.41%, a specificity of 29.41%, a positive predictive value

of 17.85%, a negative predictive value of 91.67%, and accuracy of 79.65%.

Discussion

We investigated the applicability of the PRE-DELIRIC model in intensive care, and the sensitivity of the model in predicting delirium with various physical and medical parameters compared with the CAM-ICU, the current gold standard.

The rate of development of delirium in this study was (9.9%). The reported delirium rate in intensive care units ranges widely, from 10% to 80%, and it is important to obtain a measurement that can easily detect delirium, since this is vital for follow-up, treatment, and care (19-21). Delirium development is affected by many factors, and its prediction and prevention will make it possible to reduce the rate of disorders capable of leading to mortality and morbidity (14). There is evidence that delirium increases the length of hospital stay, exacerbates the risk of transmission of in-hospital infections, and puts the patient at risk of pressure ulcers and injuries (22-24).

CAM-ICU exhibited sensitivity of 88.24% (95% confidence interval ([CI] 0.538-0.760) in this study and specificity of 85.16%. In their prospective, observational study, Guenther et al. (2021) determined a risk of delirium of

Table 4. Diagnostic screening tests and ROC curve results for Pre-Deliric scores

Pre-Deliric (%)	Diagnostic Scan				
	Cut off	Sensitivite	Spesifisite	Positive Predictive Value	Negative Predictive Value
Low risk	≥7,58	88,24	49,68	16,13	97,47
Medium risk	≥8,06	64,71	52,90	13,10	93,18
High risk	≥8,32	47,06	60,00	11,43	91,18
Very high risk	≥9,32	29,41	72,90	10,64	90,40
Predeliric>5		100	6,45	10,49	100
Predeliric>10		29,41	85,16	17,85	91,67

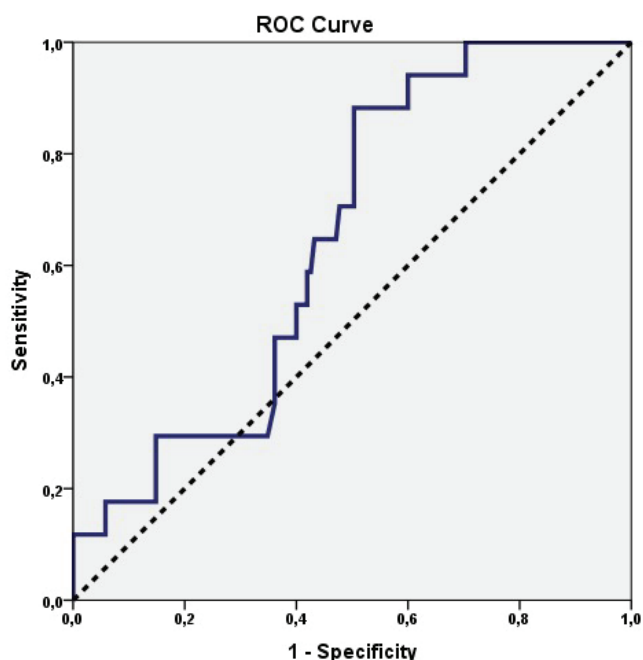


Figure 2. ROC curve for Pre-Deliric score by groups

19.8%, with CAM-ICU exhibiting sensitivity of 71% ([CI] 44-90%) and specificity of 100%, while another study reported a risk of delirium development of 25.2%, and CAM-ICU sensitivity of 100% ([CI] 92-99%) and specificity of 98% (6,25). A meta-analysis of nine separate studies concluded that CAM-ICU exhibited 80.0% sensitivity (95% CI: 77.1% and 82.6%) and 95.9% specificity (95% CI: 94.8% and 96.8%) (26). These inconsistencies may be due to variations in sampling and patient diagnoses.

CAM-ICU-positive cases exhibited higher scores than negative cases in this study. The risk of CAM-ICU-positivity was 7.404 times higher in cases with PRE-DELIRIC scores of 7.58 or higher (OR:7,404; 95% CI:1.638-33.469). Previous studies have emphasized that PRE-DELIRIC and CAM-ICU scores exhibit a better performance in identifying delirium,

and that PRE-DELIRIC scores are important due to their simplicity, reliability, and rapid use (10,27). High PRE-DELIRIC scores have been reported in patients with positive CAM-ICU values in the literature (28,29). Similarly in the present study, a positive correlation was determined between CAM-ICU and PRE-DELIRIC.

The relevant factors in patients developing delirium according to the CAM-ICU in the present study were age, previous CVE, being physically restrained, presence of a pressure ulcer, and the form of intensive care discharge. Delirium was found to develop more commonly in elderly patients, those under physical restraint, in patients with a history of CVE, and in those with pressure ulcers. The predisposing factors reported to be related to delirium in the literature are similar to those found in the present study (12,30,31). Alcohol abuse, a history of dementia, hypertension, sedation, a high APACHE II score, mechanical ventilation, and metabolic acidosis have also been described as risk factors for the development of delirium in other studies (4,12,30,31). Since we only included intensive care patients with no history of chronic alcohol abuse, delirium or dementia, and no communication problems, and with a Richmond Agitation Sedation Scale (RASS) score of +4 to -2 in this study, we may have been unable to detect all predisposing factors. However, the question of whether the PRE-DELIRIC model should not be taken into account in patients with a history of dementia or alcohol abuse and those who may have significant risk factors for delirium is a controversial one. Van den Boogaard et al. (2014) excluded groups of patients with a history of alcohol abuse and dementia from their PRE-DELIRIC regression model due to the low prevalence rate and reported that preventive measures can be taken directly instead of predicting the delirium risk, since the present evidence shows that these patients are already at a high risk of delirium (10). This has

been criticized as a deficiency of the PRE-DELIRIC model by many researchers (4,31,32).

The PRE-DELIRIC score is easily applied and uses objective data, without the need for the patient to be conscious. This score can also be used as an important screening tool in detecting delirium in patients who are unable to communicate. We recommend the use of the PRE-DELIRIC score in determining the risk of developing delirium in intensive care patients since it is easy to use and calculate and can make a useful contribution to clinical practice.

This study cannot be generalized to the general population since it was conducted at a single center within a limited time frame. The number of patients included in the study was also quite low. In addition, the inclusion of only conscious and communicating patients at the intensive care unit, in which delirium evaluation was not routinely performed, may have resulted in a lower incidence than usual in this study in which we observed an incidence rate at the lower limit of the range reported in the literature.

Conclusion

Routine use of the PRE-DELIRIC score will make it possible to safely and easily predicting the risk of delirium within the first 24 hours after admission to the ICU. Evidence-based literature support of whether the model provides a fully valid and reliable risk estimate will require its common use in intensive care patients in addition to further interventional and observational studies to decrease the risk of delirium. In addition, CAM-ICU can predict the presence of delirium, and the PRE-DELIRIC model is beneficial when making a preliminary prediction and evaluation. The PRE-DELIRIC score is convenient for determining the risk of delirium development in patients hospitalized in the ICU and connected to a mechanical ventilator. There is currently no suitable screening test for delirium diagnosis, especially in disorientated patients. The present delirium screening tests require evaluation using a subjective method, in other words by means of answers to questions put to conscious and communicating patients.

Ethics

Ethics Committee Approval: Ethics committee approval was obtained from the Ethics Committee of the Institute of Health Sciences of Kırklareli University (decision no: 3, date: 11.10.2019).

Informed Consent: Consent was obtained from the patients participating in the study.

Peer-review: Externally peer-reviewed.

Conflict of Interest: No conflict of interest was declared by the authors.

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