

Translation and cultural adaptation of the Greek version of the confusion assessment method diagnostic algorithm and the nursing delirium screening scale and their inter-rater reliability: A prospective cohort study

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Abstract

Aim: The lack of standardized tools limits the diagnosis of postoperative delirium (POD) in the Greek population. Our aim was the translation and the cultural adaptation of the confusion assessment method (CAM) diagnostic algorithm and the nursing delirium screening scale (nu-DESC) in the Greek surgical population, and the determination of their inter-rater reliability.

Methods: After Ethical approval and registration as a clinical trial (NCT04154176), a prospective cohort study was conducted in the Department of Anesthesiology, University Hospital of Larissa, Greece. Patients at least 60 years old, undergoing elective non-cardiac surgery, under general anesthesia were included.

Results: Data from 60 patients, 180 records in total, were analyzed. There was an “almost perfect agreement” between the raters with the use of CAM (Cohen’s Kappa estimate: 0.960; 95 % CI: 0.905-1.000) and nu-DESC (Cohen’s Kappa estimate: 0.981; 95 % CI: 0.944-1.000). The agreement on each specific question of CAM and nu-DESC ranged from “substantial” to “almost perfect agreement”. Based on the CAM, the sensitivity and specificity of nu-DESC were 0.97 (95 % CI: 0.82-1.00) and 0.99 (95 % CI: 0.96-1.00), respectively. The Greek versions of CAM and nu-DESC showed a high inter-rater agreement.

Conclusion: With the translation, the cultural adaptation, and the determination of their inter-rater agreement, the CAM diagnostic algorithm and the nu-DESC may serve as reliable instruments for the detection of POD in the Greek population. HIPPOKRATIA 2020, 24(1): 8-14.

Keywords: Delirium, diagnosis, neuropsychological tests, confusion assessment method, nursing delirium screening scale, anesthesia, general, reproducibility of results, humans, Greece

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Introduction

Postoperative delirium (POD) is a prevalent complication associated with increased morbidity and mortality in patients undergoing surgery and/or anesthesia¹⁻⁴. POD is defined as a state of acute and fluctuating cerebral dysfunction, hallmarked by disturbances in attention and cognition, not attributable to pre-existing cognitive dysfunction^{1-3,5}. The prevalence of POD varies based on the type of surgery and the procedural risk. For lower-risk procedures, such as general surgery, the prevalence is <13 %, whereas, for higher-risk procedures, including major abdominal surgery, it rises up to 50 %^{3,6}.

As stated by the 2017 guidelines of the European Society of Anaesthesiology (ESA), all patients undergoing

surgery should be screened for POD with standardized tools¹. Over time, several instruments were developed to diagnose POD^{7,8}, including the Confusion Assessment Method (CAM) diagnostic algorithm⁹ and the Nursing Delirium Screening Scale (nu-DESC)¹⁰. Inouye et al⁹ initially described CAM in the early 1990s^{7,9,11}. The CAM diagnostic algorithm is based on the four cardinal elements of postoperative delirium: i) acute onset and fluctuating course, ii) inattention, iii) disorganized thinking, and iv) altered level of consciousness. It can be administered in less than five minutes by non-psychiatrist physicians^{9,11}. For the diagnosis of POD, the first two and either of the latter two features are required^{9,11}. Nu-DESC was developed in 2005 by Gaudreau et al¹⁰. It is a five items screen-

ing scale, and it can be administrated in approximately one minute¹¹. It assesses disorientation, inappropriate behavior, inappropriate communication, hallucination, and psychomotor retardation¹¹. Each item is rated on a three-point scale (0-2), and the total score varies from zero to ten. The cutoff value for POD is reported to be two¹¹. Although it was initially developed to be administrated by nurses, based on a recent study, experts suggest that there is no significant difference between the evaluation results from the nursing staff and the physician, which makes the tool equally usable by both groups¹².

As far as the Greek population is concerned, neither CAM nor nu-DESC has been translated and adapted, and thus their diagnostic accuracy is unknown. Our aim was the translation and the cultural adaptation of the CAM diagnostic algorithm and the nu-DESC in the Greek population and the determination of their inter-rater reliability under the above-mentioned setting. Our secondary aim was to estimate the diagnostic accuracy of nu-DESC and the incidence of POD in our study sample.

Materials and Methods

Setting

A prospectively designed reliability study was performed. Ethical approval was provided by the Scientific Board of University Hospital of Larissa, Greece (42627/04/10/2019). The study was conducted in accordance with the Ethical Principles for Medical Research Involving Human Subjects (Declaration of Helsinki) and was registered (NCT04154176)¹³. All patients were fully informed about the protocol and provided written informed consent before participation. The study conforms with the STROBE Statement reporting standards for cohort studies, the ISPOR guidelines for translation and cultural adaptation, and the GRRAS Statement reporting standards for reliability/agreement studies¹⁴⁻¹⁶.

Participants

We included consecutive patients who underwent elective non-cardiac surgery at the General University Hospital of Larissa during November 2019. Our inclusion criteria were: i) adult patients over 60 years of age, ii) American society of anesthesiologists (ASA) physical status I to III^{3,7}, iii) undergoing elective non-cardiac surgery under general anesthesia, iv) native speakers of the Greek language, v) eligible to leave the post-anesthesia care unit, and vi) an expected in-hospital stay of at least 24 hours following surgery. We exclude patients from our study if they had any of the following: i) refused to participate or sign the informed consent form¹³, ii) underwent surgery or anesthesia within the last 30 days^{3,17}, iii) prior or current history involving an affliction of the central nervous system^{3,5,8,18}, iv) severe hearing or visual impairment³, v) psychiatric disorders^{3,7}, vi) a score less than five according to the Geriatric Depression Scale (GDS-15)^{3,7,19}, vii) a score less than four for females and less than two for males according to the Lawton-Brody Instrumental Activities of Daily Living Scale (I. A. D.

L.)^{7,17,20}, viii) alcohol consumption more than 35 units/week^{3,7}, ix) drug dependence^{3,7}, x) previous neuropsychological testing^{3,7}, xi) hemodynamical instability^{3,7}, and xii) peri-procedural desaturation (one or more events of SpO₂ < 80 % for more than two minutes)^{3,7}.

Translation and Cultural Adaptation

The translation process from English to Greek begun after gaining permission and consent from the authors/developers of both the original instruments^{9,10}. Three authors performed three separate forward-translations, for each of the two instruments, from English to Greek. Subsequently, the translations were merged in one preliminary version¹⁴. Our next step was reconciliation. The Greek versions were adapted to the clinical setting without changing the meaning through a pilot study. Each phrase was extensively discussed between the translators. Different wording was tested throughout the process, and the final form was established after reaching a consensus between the two authors/developers¹⁴. A bilingual expert interpreter who was blinded to the original English version performed the back-translation into English¹⁴. The back translation review followed, and the back translation of the preliminary Greek version of both the instruments, was thoroughly compared with the original texts. After closed comparison and assessment, there were no substantial deviations from the original instruments¹⁴. The instrument developers reviewed the English back-translated version regarding the conformity of content and language and the agreement with the original versions. The appropriate modifications were made, and the latest step of the translation was approved (harmonization)¹⁴. The final Greek versions of CAM and nu-DESC underwent a structured evaluation by eight doctors from two different specialties (cognitive debriefing)¹⁴. The cognitive debriefing results were reviewed, and the translation was finalized¹⁴. The final translations of CAM (Figure 1) and nu-DESC (Figure 2) were proofread by three translators¹⁴. Four of the authors reviewed the process of the transaction (final report)¹⁴.

Study procedures

Two experienced anesthesiologists, who have been trained according to the original training manuals of the CAM and nu-DESC instruments, evaluated the patients for the presence of POD^{10,21}. The two raters were blinded and had no access to each other's evaluation. They independently administered the CAM and nu-DESC questionnaires during the first postoperative day after the discharge from the post-anesthesia care unit. The assessments of the two different rates were completed within 60 minutes. CAM was administrated within five minutes and nu-DESC within one in accordance with the international standards¹¹. The patients who were diagnosed with POD received the appropriate treatment according to the hospital's protocol.

Statistical analysis

Descriptive statistics were used to summarize gen-

Φύλλο εργασίας Συντομομένης Μεθόδου Εκτίμησης Παραληρήματος (Συντομομένη κλίμακα-CAM)

Σημειώση: Το συγκεκριμένο φύλλο εργασίας μπορεί να χρησιμοποιηθεί ως αναλυτική ή σύντομη έκδοση του εργατολογίου CAM. Η αξιολόγηση του προσανατολισμού και της ικανότητας διατήρησης της προσοχής συνίσταται πριν την βαθμολόγηση, με την αξιολόγηση με αριθμητικά διαστήματα, μέρες της εβδομάδας ή μήνες του έτους αντίστοιχα. Το συγκεκριμένο φύλλο εργασίας μπορεί να χρησιμοποιηθεί μόνον για την αναγνώριση ασθενών με παραληρήματα. Σημειώστε ότι δεν μπορεί να χρησιμοποιηθεί για την αξιολόγηση της βαρύτητας του παραληρήματος με τη χρήση της κλίμακας CAM-5.

ΕΥΑΛΩΤΩΡ: _____ ΗΜΕΡΟΜΗΝΙΑ: _____

I. ΟΞΕΙΑ ΕΙΤΒΩΔΗ ΚΑΛΚΥΜΑΙΝΟΜΕΝΗ ΕΝΤΑΣΗ

α) Υπάρχουν αποδείξεις οξείας μεταβολής στην νοητική κατάσταση του ασθενή σε σύγκριση με τη φυσιολογική του κατάσταση: Όχι Ναι

β) Η μη φυσιολογική του συμπεριφορά μεταβάλλεται κατά τη διάρκεια της ημέρας, δηλαδή εμφανίζεται και εξαφανίζεται ή αυξάνεται και ελαττώνεται σε βαρύτητα: Όχι Ναι

II. ΔΥΝΑΜΙΑ (ΑΠΟΣΠΑΣΗ) ΠΡΟΣΟΧΗΣ

Δυσκολεύεται ο ασθενής να επικεντρωθεί, για παράδειγμα απασχούται η προσοχή του εύκολα ή εμφανίζει δυσκολία να επικεντρωθεί σε όλα τα λόγια: Όχι Ναι

III. ΜΗ ΟΡΓΑΝΩΜΕΝΗ ΣΚΕΨΗ

Είναι η σκέψη του ασθενή αναorganισμένη ή ανακόλλητη, όπως διαδοχικός ή μη συσχετισμός συζητήσεων με μη λογική και μη καθαρή ροή ιδεών, ή μη προβλεπόμενες μεταβάσεις από ένα θέμα σε άλλο: Όχι Ναι

IV. ΜΕΤΑΒΑΛΛΟΜΕΝΟ ΕΠΙΠΕΔΟ ΣΥΝΕΙΔΗΣΗΣ

Συνολικά πώς θα χαρακτηρίσετε το επίπεδο συνείδησης του ασθενή:
 -- Σύντομος (φυσιολογικό)
 -- Σε επαγρύπνηση (με τεταμένη προσοχή)
 -- Αιθαρρικός (νυσταγμένος, εύκολα αφηνιάσιμος)
 -- Βασικά ληθαργικός (δύσκολο αφηνιάσιμος)
 -- Κοιμάται (αδυναμία αφηνιάσιμος)

Υπάρχουν τετακτοίματα στο παραπάνω κουτί: Όχι Ναι

Εάν "Αδυναμία Προσοχής" και τουλάχιστον ένα ακόμη αντικείμενο στο Κουτί 1 έχει σημειωθεί και τουλάχιστον ένα αντικείμενο στο Κουτί 2 έχει σημειωθεί, προτείνεται η διάγνωση του παραληρήματος.

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Figure 1: The translated and adapted Greek version of the confusion assessment method (CAM) diagnostic algorithm.

eral study parameters, including the demographic data and the questionnaire scores. Count data were summarized in counts and percentages, whereas continuous parameters by their central tendency and range. Moreover, we used repeated-measures analysis of variance to test for statistically important differences among repeated measurements. The index of dissimilarity, percentage of agreement, and Cohen’s kappa estimate were used to estimate the inter-rater agreement analysis. The former is a statistic used to measure the overall difference between two percent distributions. It ranges from 0 to 100 and indicates the proportion of items required to be reallocated to make the two distributions the same²². The percentage of agreement is a basic measure for inter-rater reliability calculated by the ratio of the number of ratings in agreement over the total number of ratings²³. Additionally, the kappa statistic represents a quantitative measure of the magnitude of agreement between observers beyond what would be expected by chance. Kappa estimates range from “0.01 to 0.20”, “0.21 to 0.40”, “0.41 to 0.60”, “0.61 to 0.80”, and “0.81 to 1.00”, and correspond to “slight agreement”, “fair agreement”, “moderate agreement”, “substantial agreement”, and “almost perfect or perfect agreement”, respectively²⁴. We used Cronbach’s alpha statistic to evaluate the internal consistency for both scales. Values of $\alpha \geq 0.9$, $0.9 > \alpha \geq 0.8$, $0.8 > \alpha \geq 0.7$, $0.7 > \alpha \geq 0.6$, $0.6 > \alpha \geq 0.5$, $\alpha < 0.5$ represented excellent, good, acceptable, questionable, and

Nursing delirium screening scale (Nu-DESC)

Ημερομηνία: _____

Χαρακτηριστικά και περιγραφή	Βαθμολογία συμπτωμάτων 0-2		
	24-8.00	8.00-16.00	16.00-24
Αποπροσανατολισμός Λεκτικές ή συμπεριφορικές εκδηλώσεις αποπροσανατολισμού στον χρόνο ή τον τόπο ή λανθασμένη αντίληψη των ατόμων στο περιβάλλον			
Ανάρμοστη συμπεριφορά Ανάρμοστη συμπεριφορά στον τόπο ή τα άτομα			
Ψευδαισθήσεις, παραισθήσεις Όραση ή ακοή πραγμάτων που δεν υπάρχουν			
Ψυχοκινητική υστέρηση Καθυστερημένη αντίδραση, λίγες ή καθόλου αυτόματες πράξεις/λέξεις, πχ όταν ο ασθενής ενοχλείται η αντίδραση καθυστερεί ή/και ο ασθενής δεν αφηνιάζεται			
Συνολική βαθμολογία (από το σύνολο των 10)			
Τα συμπτώματα βαθμολογούνται από το 0 έως το 10 ανάλογα με την παρουσία και την ένταση των συμπτωμάτων, και η κάθε βαθμολογία σθροίζεται για να οριστεί η τελική συνολική βαθμολογία.			
Βαθμολογία >2 στο nu-DESC δηλώνει την ύπαρξη παραληρήματος στο 86% των ασθενών.			

Figure 2: The translated and adapted Greek version of the nursing delirium screening scale (nu-DESC).

poor scale internal consistency, respectively. The diagnostics accuracy of nu-DESC was assessed in terms of sensitivity, specificity, positive predictive value, negative predictive value, and positive and negative likelihood ratios, using CAM recordings as the reference tool. All data were analyzed using the statistical environment R²⁵. Statistical significance for all analyses was set at $p < 0.05$.

Study size

The required sample size (questionnaires/records) for the inter-observer agreement analysis using Cohen’s Kappa statistic (delirium prevalence: 0.25; the true Cohen’s Kappa statistic: 0.96; kappa under the null hypothesis: 0.7; power analysis: 0.90; type I error: 0.05; two-sided) was estimated to be 65 questionnaires/records. With an anticipated dropout rate of 20 %, our study required a minimum of 78 records.

Results

A total of 79 patients were eligible and included initially. Of them, 19 patients refused re-evaluation (Figure 3). Three pairs of records were available for each patient; at 8, 16, and 22 hours post-operatively. Our final sample size consisted of 180 records/questionnaires in total. There were 33 (55 %) females and 27 (45 %) males with a mean age of 71 (standard deviation: 6.23) years. Table 1 and Table 2 summarize the main characteristics of our study sample. There were not any missing data for each variable of interest.

Inter-observer reliability and incidence of POD

According to the first observer, eight (13.3 %), eleven (18.3 %), and twelve (20 %) patients were diagnosed with POD using CAM at the initial examination, 16-hour, and 22-hour, respectively. For the second observer, there were eight (13.3 %), eleven (18.3 %), and ten (16.7 %) patients with POD at the 8-hour, 16-hour, and 22-hour examination, respectively. Based on the nu-DESC, the first observer identified nine (15 %) cases with POD at the 8-hour evaluation, and eleven (18 %) and thirteen (22 %) cases at the 16-hour and 22-hour re-examination, respectively. At the same time points, the second observer recorded eight (13 %), eleven (18 %), and thirteen (22 %) patients with POD. According to the repeated measurement analysis, there was no statistically important difference in POD occurrence (Figure 4). The two independent raters completed 180 paired evaluations (Table 3 and Table 4). There was an “almost perfect agreement” between the two raters with the use of CAM [Cohen’s

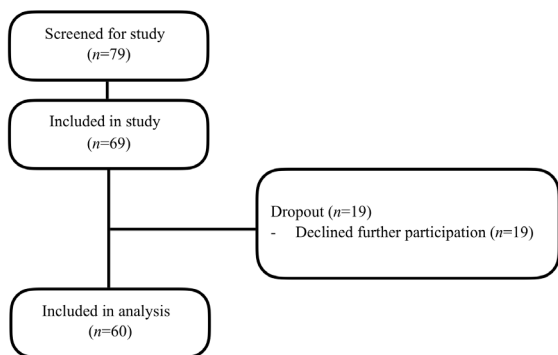


Figure 3: Flowchart depicting the patient sample selection.

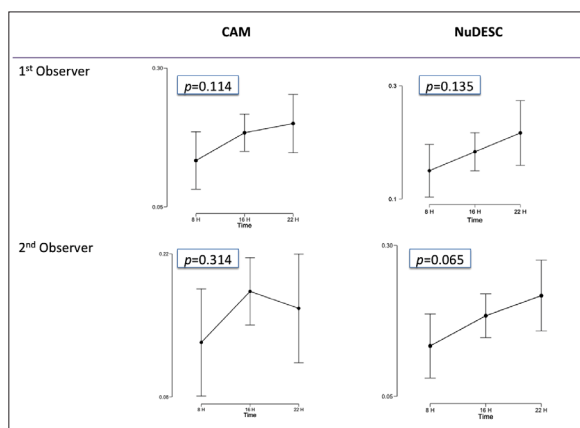


Figure 4: A two-by-two graph visualizing the interaction plots between the two observers scores (mean values and 95 % confidence interval of the occurrence of postoperative delirium) and time for the confusion assessment method (CAM) diagnostic algorithm and the nursing delirium screening scale (nu-DESC). The scores were measured at 8, 16, and 22 hours after surgery; p values indicate the statistical significance among the repeated measurements. Accordingly, there was no statistically important change in the scores with time in our study sample.

Kappa estimate: 0.960; 95 % Confidence Interval (CI): 0.905-1.000] and nu-DESC (Cohen’s Kappa estimate: 0.981; 95 % CI: 0.944-1.000). The agreement on each specific question of CAM and nu-DESC ranged from “substantial agreement” to “almost perfect agreement”, as depicted in Table 5 and Table 6. The internal consistency of both questionnaires ranged between “excellent” and “good” (Table 7). Considering the Cronbach’s alpha values, the measurements at 16 hours provide significant information if a measurement is dropped.

Table 1: Summary of the count parameters of the 60 consecutive patients who underwent elective non-cardiac surgery and were included in the study.

	Subgroup	Counts	%
Gender	M	33	55
	F	27	45
ASA physical status	2	35	58.3
	3	25	41.7
Type of Surgery	General	21	35
	Vascular	11	18.3
	Gynaecological	12	20
	Urogical	8	13.3
	ENT	3	5
	Orthopaedic	2	3.3
Neurosurgical	2	3.3	
Orthognathic	1	1.7	

M: male, F: female, ASA: American society of anesthesiologists, ENT: Ear, nose, throat.

Table 2: Summary of the continuous parameters of the 60 consecutive patients who underwent elective non-cardiac surgery and were included in the study.

	Age (years)	Anesthesia Duration	Surgery Duration
Mean	71.1	138	123
Median	71.07	130	110
Standard deviation	6.23	66.35	62.1
Minimum	60	30	25
Maximum	85	420	370

Duration of anesthesia and surgery is reported in minutes.

Table 3: Two by two table between the two observers for the confusion assessment method (CAM) diagnostic algorithm.

	2 nd Observer	
	0	1
1 st Observer	0	146
	1	4

Table 4: Two by two table between the two observers for the nursing delirium screening scale (nu-DESC).

	2 nd Observer	
	0	1
1 st Observer	0	147
	1	1

Table 5: Inter-rater agreement analysis of the confusion assessment method (CAM) diagnostic algorithm.

	Index of dissimilarity	Percentage of agreement	Cohen's kappa estimate *	
Acute onset / fluctuating course	0.097	98.3	0.930 (0.870-1.00)	Almost perfect agreement
Inattention	0.033	99.4	0.979 (0.939-1.00)	Almost perfect agreement
Disorganized thinking	0.413	92.7	0.683 (0.518-0.849)	Substantial agreement
Altered level of consciousness	0.448	92.1	0.654 (0.479-0.828)	Substantial agreement
Total	0.0645	98.9	0.960 (0.905-1.000)	Almost perfect agreement

Values in brackets for Cohen's kappa represent 95 % confidence interval.

Table 6: Inter-rater agreement analysis of the nursing delirium screening scale (nu-DESC).

	Index of dissimilarity	Percentage of agreement	Cohen's kappa estimate *	
Disorientation	0.037	99.4	0.9731 (0.920-1.000)	Almost perfect agreement
Inappropriate behavior	0.210	97.1	0.815 (0.656-0.975)	Almost perfect agreement
Inappropriate communication	0.157	98.9	0.928 (0.821-1.000)	Almost perfect agreement
Illusions / Hallucination	0.307	97.7	0.806 (0.619-0.993)	Almost perfect agreement
Psychomotor retardation	0.272	97.1	0.709 (0.458-0.960)	Substantial agreement
Total	0.033	99.4	0.981 (0.944-1.000)	Almost perfect agreement

Values in brackets for Cohen's kappa represent 95 % confidence interval.

Table 7: Scale reliability statistics. Considering the Cronbach's alpha values and Cronbach's alpha values if a measurement is dropped.

Scale	1 st Observer		2 nd Observer			
	Cronbach's alpha	Cronbach's alpha (if item dropped)	Cronbach's alpha	Cronbach's alpha (if item dropped)		
CAM	0.912	8 hours	0.904	8 hours	0.906	
		16 hours		0.813	16 hours	0.759
		22 hours		0.940	22 hours	0.902
nu-DESC	0.918	8 hours	0.897	8 hours	0.886	
		16 hours		0.813	16 hours	0.762
		22 hours		0.940	22 hours	0.902

CAM: Confusion assessment method, nu-DESC: Nursing delirium screening scale. Sixty of the observations were used, 0 were excluded list wise, and 60 were provided. Values of $\alpha \geq 0.9$, $0.9 > \alpha \geq 0.8$, $0.8 > \alpha \geq 0.7$, $0.7 > \alpha \geq 0.6$, $0.6 > \alpha \geq 0.5$, $\alpha < 0.5$ represented excellent, good, acceptable, questionable, and poor scale internal consistency, respectively.

Diagnostic accuracy of nu-DESC

For the first observer, CAM and nu-DESC were at an "almost perfect agreement" (Cohen's Kappa estimate: 0.960; 95 % CI: 0.91-1.000). Based on CAM, nu-DESC was characterized by a sensitivity and specificity as high as 0.94 (95 % CI: 0.80-0.99) and 1.00 (95 % CI: 0.98-1.00), respectively (Figure 5). Likewise, for the second reviewer, there was an "almost perfect agreement" between the two scales (Cohen's Kappa estimate: 0.90; 95 % CI: 0.82-0.98). With the later values, based on CAM, the sensitivity and specificity of nu-DESC were 0.97 (95 % CI: 0.82-1.00) and 0.99 (95 % CI: 0.96-1.00), respectively (Table 8).

Discussion

POD can prove to be a devastating complication for

the surgical population^{3,5}. Experts strongly recommend screening all patients for delirium, in each shift, up to the 5th postoperative day¹. CAM and nu-DESC are among the recommended tools based on the guidelines released by ESA in 2017¹. Of note, both instruments are not time-consuming; they do not require extensive training and can be administrated by non-psychiatrists^{1,3,5,11}.

This study provides the first official Greek translation and cultural adaptation of CAM and nu-DESC, according to the ISPOR guidelines, and the first evaluation of their inter-rater agreement. The Greek version of CAM and nu-DESC have a high inter-rater agreement, and ultimately, nu-DESC was characterized by significant diagnostic accuracy based on CAM. The incidence of POD in our cohort was less than 20 %, using CAM and nu-DESC.

Our findings are comparable with the originally de-

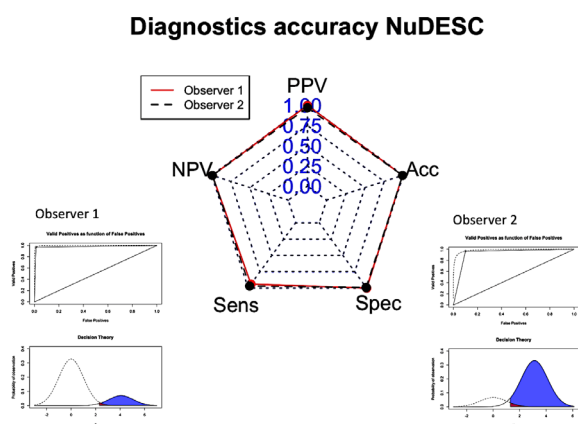


Figure 5: Nursing delirium screening scale (nu-DESC) is an excellent tool in screening for postoperative delirium. The spider plot visualizes the accuracy of nu-DESC as a screening tool for delirium in postoperative patients based on the confusion assessment method (CAM) diagnostic algorithm as a gold standard. The inserts highlight the diagnostic accuracy analysis's inherent details, including the area under the curve (AUC).

Spec: specificity, Sens: sensitivity, Acc: accuracy, PPV: positive predictive value, NPV: negative predictive value.

veloped versions of CAM and nu-DESC. In particular, Inouye et al reported inter-rater reliability for CAM as high as 93 %⁹. Regarding the features of CAM, the agreement ranged from “substantial agreement” for the features of disorganized thinking and altered level of consciousness to “almost perfect agreement” for the acute onset/fluctuating course and inattention. Inouye et al described agreement at 56 % and 100 % for the altered level of consciousness and the disorganized thinking, respectively⁹. In another study, Martins et al validated the European Portuguese version of CAM and stated that the inter-rater reliability in their study was 1.0 for the acute onset, 0.78 for inattention, 0.65 for disorganized thinking, and 1.0 for altered level of consciousness²⁶. These minor disparities in the agreement among the different translations could be attributed to the distinctive characteristics in the two raters' experience, the non-homogeneity of the study populations, and, more importantly, the patients' dissimilarities regarding their educational background.

Likewise, Gaudreau et al focused on the agreement between CAM and nu-DESC, and the reported sensitivity

and specificity rates were as high as 85.7 % and 86.8 %, respectively^{9,10}. In our study, the features disorientation, inappropriate behavior, inappropriate communication, and illusions/hallucinations were associated with “almost perfect agreement”. However, the feature psychomotor retardation was characterized by “substantial agreement”. One possible explanation could be the high proportion of the hyperactive type of POD (23/60) in our study sample. Moreover, our sample's homogeneity (patients over the age of sixty and without dementia, who underwent elective non-cardiac surgery under general anesthesia) could be another reasonable justification. The predictive value of nu-DESC can vary considerably between populations with low versus high prevalence of delirium, and the positive and negative predictive values of our results (Table 8) highlight the significance of the homogeneity of our sample^{10,12}.

Additionally, the fact that the postoperative delirium's assessment was performed by anesthesiologists, who were able to evaluate the fluctuating daily course of POD, may strengthen the high inter-rater reliability of our study. The fluctuating course of postoperative delirium makes its clinical routine detection and diagnosis quite challenging. Thus, the dedicated and experienced physicians play a central role in the diagnosis of postoperative delirium³. Although nu-DESC was initially developed to be administrated by nurses, Hägi-Pedersen et al performed the validation of the Danish version of nu-DESC successfully by estimating the inter-rater agreement between medical doctors and nurses¹². Their study showed that the nu-DESC is comprehensible and that there is no significant difference between the nursing staff and medical doctors' evaluation results, which suggests that the tool may be equally usable by both groups¹².

A limitation in the present study could be the selection of the population. Besides the numerous advantages of our study sample's homogeneity, further studies are necessary for assessing the validity of CAM and nu-DESC to other surgical populations (non-elective procedures, cardiac surgery). Additionally, we did not choose to compare either CAM or nu-DESC to Diagnostic and Statistical Manual of Mental Disorders (DSM) V or IV because an isolated comparison did not seem meaningful^{9-12,21,27,28}. Other CAM and nu-DESC translations were only compared to DSM IV, and the whole postoperative delirium focus has shifted significantly with the progression from DSM IV to V^{1,10-12,26,29}. Moreover, the utility

Table 8: Accuracy parameters of the nursing delirium screening scale (nu-DESC) as a screening tool for postoperative delirium, based on the confusion assessment method (CAM) diagnostic algorithm.

	1 st Observer	2 nd Observer
Apparent prevalence	0.17 (0.12-0.24)	0.16 (0.11-0.23)
True prevalence	0.18 (0.13-0.25)	0.16 (0.11-0.23)
Sensitivity	0.94 (0.80-0.99)	0.97 (0.82-1.00)
Specificity	1.00 (0.98-1.00)	0.99 (0.96-1.00)
Positive predictive value	1.00 (0.89-1.00)	0.97 (0.82-1.00)
Negative predictive value	0.99 (0.95-1.00)	0.99 (0.96-1.00)
Positive likelihood ratio	Inf (NaN, Inf)	142.90 (20.24-1008.95)
Negative likelihood ratio	0.06 (0.02-0.23)	0.03 (0.01-0.24)

Values in brackets represent 95 % confidence interval, Inf: infinity, NaN: not a number.

of DSM has been questioned during the past years and, based on current literature, there is only a 30 % overlap between the DSM-IV and DSM-V in delirium diagnosis under strict criteria^{3,4,10-12,28,29}. Of note, both screening tools, CAM and nu-DESC, have been abundantly validated in comparison to DSM in numerous different languages and settings^{1,10-12,21}.

With the translation, the cultural adaptation, and the determination of the inter-rater agreement of the Greek versions of the CAM and nu-DESC, our study provides two reliable screening tools for the diagnosis of POD in our country. Moreover, as nu-DESC can detect all psychomotor types of delirium, its implementation in daily practice may increase awareness about POD in Greece. Both CAM and nu-DESC are internationally recognized scales recommended by the ESA. Thus, they may be used in multi-center national and international clinical research, improving the safety and the peri-operative care of patients undergoing surgery and/or anaesthesia.

Conflict of Interest

Authors declare no conflict of interest and not receiving any specific grant from any funding agency in the public, commercial, or not-for-profit sectors.

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Adapted from: Gaudreau JD, et al. *J Pain Symptom Manage.* 2005; 29: 368-375. Fast, systematic, and continuous delirium assessment in hospitalized patients: the nursing delirium screening scale.

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