



Trabajo Original

Epidemiología y dietética

Utility of the EAT-10 in the detection of dysphagia in high-risk hospitalisation units at a university hospital: a cross-sectional study

Utilidad del EAT-10 en la detección de la disfagia en las unidades de hospitalización de alto riesgo de un hospital universitario: un estudio transversal

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Abstract

Introduction: the objective was to assess the utility of the Eating Assessment Tool (EAT-10) in hospitalisation units with patients at high risk of dysphagia.

Patients and methods: a cross-sectional study was conducted in the Neurology and Internal Medicine wards; patients with admission < 24 hours and in a terminal stage of disease were excluded. In the first 24-48 hours of admission the presence of dysphagia as assessed with the EAT-10, the risk of malnutrition as assessed with the Malnutrition Universal Screening Tools (MUST), and comorbidities using the Charlson index were screened.

Results: a total of 169 patients were recruited (76.0 years, 52 % women); 19.5 % were at risk of malnutrition. The EAT-10 instrument could be administered in 80.6 % of the patients, and was positive in 26.6 % (women 34.1 % vs. men 18.4 %; $p = 0.025$). When comparing patients with higher comorbidity with those with a lower Charlson index, a lower response rate to EAT-10 was observed (78.4 % vs. 93.9 %; $p = 0.038$), without differences in screening positivity (28.3 % vs. 19.4 %; $p = 0.310$). The prevalence of dysphagia risk was higher in the Internal Medicine unit than in the Neurology unit (30.4 % vs. 19.6 %; $p = 0.133$), as was the percentage of cases in which screening could not be performed (21.1 % vs. 11.1 %; $p = 0.011$). There were no significant differences in risk of malnutrition, mortality, hospital stay, or readmission according to the EAT-10.

Conclusions: the EAT-10 has limited utility in the studied hospitalisation units due to a high rate of unfeasible tests, especially among patients at higher risk of dysphagia.

Keywords:

Dysphagia.
Screening. Eating
Assessment Tool.
Malnutrition.

Resumen

Introducción: el objetivo del estudio fue evaluar la utilidad del Eating Assessment Tool (EAT-10) en unidades de hospitalización con pacientes de alto riesgo de disfagia.

Pacientes y métodos: estudio transversal de pacientes hospitalizados en Medicina Interna y Neurología; los pacientes con ingreso < 24 horas y en fase terminal de la enfermedad fueron excluidos. En las primeras 24-48 horas de ingreso se cribó la disfagia con el EAT-10, el riesgo de desnutrición con el Malnutrition Universal Screening Tool (MUST) y la comorbilidad con el índice de Charlson.

Resultados: se reclutaron 196 pacientes (76,0 años, 52 % mujeres). El 19,5 % estaban en riesgo de desnutrición. El EAT-10 se pudo realizar en el 80,6 % de la muestra y fue positivo en el 26,6 % (mujeres 34,1 % vs. hombres 18,4 %; $p = 0,025$). Al comparar a los pacientes con mayor comorbilidad con aquellos que tenían un índice de Charlson más bajo, se observó una tasa de respuesta más baja al EAT-10 (78,4 % vs. 93,9 %; $p = 0,038$), sin diferencias en la positividad del cribado (28,3 % vs. 19,4 %; $p = 0,310$). La prevalencia del riesgo de disfagia fue mayor en la unidad de Medicina Interna que en la de Neurología (30,4 % vs. 19,6 %; $p = 0,133$), así como el número de casos en que no se pudo realizar el cribado (21,1 % vs. 11,1 %; $p = 0,011$). No hubo diferencias significativas en el riesgo de desnutrición, mortalidad, estancia hospitalaria o reingreso según el EAT-10.

Conclusiones: el EAT-10 tiene una utilidad limitada en las unidades de hospitalización estudiadas debido a una alta tasa de pruebas no realizables, especialmente entre los pacientes con mayor riesgo de disfagia.

Palabras clave:

Disfagia. Cribado.
Eating Assessment
Tool. Desnutrición.

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INTRODUCTION

Swallowing is defined as the activity of transporting solid and liquid substances, as well as saliva, from the mouth to the stomach. It implies the coordinated participation of anatomical areas that, through movements and pressures, allow food to be conducted efficiently and safely through the digestive tract (1). Dysphagia represents any type of alteration or difficulty in swallowing, which can occur in any of its phases (oral, pharyngeal, or oesophageal). Depending on the swallowing phase that has been affected, we can classify it into preparatory oral dysphagia, oral phase dysphagia, pharyngeal dysphagia, and oesophageal dysphagia. However, from the practical point of view, dysphagia is classified into two large groups: oropharyngeal dysphagia and oesophageal dysphagia. From the pathophysiological perspective, functional and structural alterations can be the cause of this syndrome (2).

The prevalence of dysphagia varies depending on the definition used, the study population, and the sensitivity of the screening or diagnostic technique involved. It has been estimated that between 16 % and 22 % of the population over the age of 50 have dysphagia. The prevalence of swallowing disorders in those over 65 years is 40 %, and above 60 % among nursing home residents. The highest prevalence (> 80 %) has been found in hospitalised patients with dementia, especially in the more advanced stages of the disease (3,4). The ageing process causes anatomical, neurological, and muscular changes that result in a loss of functional capacity that may affect swallowing. In the elderly who do not have health problems, these changes are defined as presbyphagia, and are not necessarily pathological. However, when these changes in swallowing physiology occur in frail elderly people, with comorbidities and poly medication, the risk of dysphagia increases (5,6).

Dysphagia has a profound effect on nutritional status as a result of decreased intake due to a loss of swallowing efficacy and fear of eating. The most frequent type of malnutrition in patients with oropharyngeal dysphagia is the caloric one, with preservation of visceral protein and a significant depletion of muscle mass and the fatty compartment. In Spain, 40 % of hospitalised patients with dysphagia are at risk of malnutrition. When focusing only on patients who are 70 years of age or older, this percentage rises to 60 % (7). Because of this high risk of malnutrition, the European Society for Swallowing Disorders (ESSD) recommends continuous monitoring of the nutritional status of patients with dysphagia (8). However, despite the current ESSD recommendations, only one in four patients with dysphagia receive nutritional support (7). In addition, the decrease in fluid intake due to dysphagia causes an imbalance of body fluids that leads to increased mortality in hospitalised patients (9). One study found that daily water intake in patients with oropharyngeal dysphagia was only 22 % of the recommended amount for these patients (10).

Up to 50 % of neurological patients and the elderly have alterations in swallowing safety (penetrations and aspirations), with a high proportion of silent aspirations (not accompanied by cough), during videofluoroscopy, and a 50 % risk of developing aspirative pneumonia, which is associated with an increase in mortality for 50 % of these patients. These respiratory infections represent the

main cause of mortality in patients with oropharyngeal dysphagia, and have also been associated with an increase in hospital stay and hospital readmissions (11).

Currently, there are multiple tests for the screening and diagnosis of dysphagia. One of the most commonly used is the Eating Assessment Tool (EAT-10) questionnaire. This method, useful for the screening of dysphagia in patients with preserved cognitive levels, includes 10 questions about swallowing that the patient has to score from 0 (no problem) to 4 (a serious problem). A score greater than 3 denotes the potential presence of dysphagia (12). This tool is widely used in the ambulatory setting, but we have little data about its usefulness in hospitalised patients. We designed this study to establish the feasibility of using the EAT-10 as a screening tool for dysphagia in hospitalised patients, to determine the prevalence of dysphagia in key hospitalisation units where there are high-risk patients (such as Internal Medicine and Neurology), and to find out whether there is a relationship between EAT-10 results and nutritional status, hospital stay, in-hospital mortality, and readmissions at 30 days.

PATIENTS AND METHODS

A cross-sectional study was designed to reach the objectives, and was developed in the acute hospitalisation wards of the Departments of Internal Medicine and Neurology of the University Hospital Complex at A Coruña. This tertiary hospital assists a reference population of 505,797 people, with 1,415 hospitalisation beds installed. The study took place between January and April 2019.

This study was carried out respecting the Helsinki Declaration of the World Medical Association, the Convention on Human Rights and Biomedicine, and the Spanish legislation about human research. The protocol was reviewed and approved by the research ethics committee of the hospital.

A sample size of 118 patients was estimated, based on the prevalence of dysphagia obtained from the PREDYCES study (20.6 %), an estimate of 1500 patients admitted in the study period, a confidence level (1- α) of 95 %, and an accuracy of 7 % (7). Inclusion criteria included age over 18 years and scheduled or urgent admission to the Neurology and Internal Medicine hospitalisation units. Patients who had been hospitalised for less than 24 hours, patients admitted to the Neurology or Internal Medicine hospitalisation unit but in charge of other hospital services, and patients in terminal stages of their disease, in whom death was expected in the following hours, were excluded.

The recruitment was performed consecutively based on the patients admitted to the above-mentioned wards within the first 24 hours of hospitalisation, upon request of an informed consent. Patients admitted during the weekend or non-working days were recruited the next working day (maximum, 48 hours after admission).

The presence of dysphagia was screened using the EAT-10, and the risk of malnutrition was assessed using the Malnutrition Universal Screening Tool (MUST). Both questionnaires were

administered by the researchers. Height and weight were measured with the stadiometer and scales at each ward, which are periodically calibrated. When it was not possible to obtain these measures directly, researchers used the alternative measurements described by the British Association of Parenteral and Enteral Nutrition (BAPEN) (13). The comorbidities of patients were measured using the Charlson comorbidity index (CCI), considering a high comorbidity level when a result higher than 3 points was obtained (14). Data such as admission date, discharge date, main diagnosis, mortality during admission, and 30-day readmissions were obtained from the medical records.

Qualitative data were summarised as percentages. The normal distribution of quantitative data was checked with the Shapiro-Wilk test and summarised using mean and standard deviation. Qualitative variables (e.g., gender, mortality, readmissions, prevalence of malnutrition) were compared using the chi-squared test. Continuous quantitative variables (e.g., hospital length of stay or age) were compared using Student's t-test for independent measurements (or the Mann-Whitney U-test). A p-value less than 0.05 was considered significant.

RESULTS

The study recruited 196 patients, whose general characteristics are presented in table I. Most patients were recruited at the Internal Medicine wards (67.9 %). The differences between patients in each ward are summarised in table II.

The EAT-10 was administered to 158 of the 196 patients included in the study (80.6 %), and was positive in 42 (26.6 %) of the 158 patients who could be screened. Patients where EAT-10 could not be administered were older (82.5 [18.0] yrs vs. 74.4 [17.5] yrs; $p = 0.012$) and had a higher CCI (6.3 [2.2] points vs. 5.1 [2.9] points; $p = 0.018$). Patients admitted to Internal Medicine were more frequently unable to answer the questionnaire (23.3 % vs. 11.1 %; $p = 0.044$). There were no significant differences in the prevalence of positive results in the EAT-10 according to hospitalization ward: Internal Medicine, 30.4 % vs. Neurology, 19.6 % ($p = 0.143$). The differences found between patients according to EAT-10 screening are analysed in table III.

According to MUST, 19.6 % of patients were at risk of malnutrition, without significant differences between hospitalisation units: Internal Medicine, 23.7 % vs. Neurology, 12.3 % ($p = 0.084$). There were no differences in the prevalence of patients with risk of malnutrition between patients with a positive or negative EAT-10 (15.8 % vs. 16.3 %, $p = 0.936$). MUST was more frequently positive among patients who could not answer the EAT-10 (58.3 % vs. 6.2 %, $p < 0.001$).

The results of the EAT-10 did not predict clinical outcomes: there were no differences in hospital length of stay [positive, 11.9 (13.3) [days vs. negative, 9.8 (11.2) days; $p = 0.256$], in-hospital mortality (positive, 4.9 % vs. negative, 4.3 %; $p = 0.888$), or readmission rate (positive, 9.4 % vs. negative, 12.9 %; $p = 0.596$). Mortality was higher in patients whose EAT-10 could not be performed (18.9 % vs. 4.5 %, $p = 0.002$).

Table I. General characteristics of the patients

Female	52.0 %
Age (years)	76.0 (17.9)
CCI (points)	5.3 (2.7)
High CCI (> 3 points)	26.6 %
EAT-10 (points)	2.3 (4.5)
MUST (points)	0.7 (1.0)
Length of stay (days)	6.5 (32.0)
Mortality	7.3 %
30-day readmission	12.0 %

CCI: Charlson comorbidity index; EAT-10: Eating Assessment Tool; MUST: Malnutrition Universal Screening Tool. Quantitative data are presented as mean (standard deviation).

Table II. Comparison of patients admitted to the Neurology and Internal Medicine units

	Internal Medicine	Neurology	p
n	133	63	-
Female	54.9 %	46.0 %	0.246
Age (years)	82.8 (11.8)	61.5 (19.7)	< 0.001
CCI (points)	6.3 (2.1)	3.3 (2.9)	< 0.001
High CCI (> 3 points)	30.4 %	16.6 %	0.143
EAT-10 (points)	2.7 (4.6)	1.7 (4.1)	0.164
MUST (points)	0.8 (1.1)	0.5 (0.7)	0.075
Length of stay (days)	10.9 (8.6)	10.3 (17.0)	0.793
Mortality	9.2 %	3.2 %	0.128
30-day readmission	12.1 %	11.9 %	0.967

CCI: Charlson comorbidity index; EAT-10: Eating Assessment Tool; MUST: Malnutrition Universal Screening Tool. Quantitative data are presented as mean (standard deviation).

Table III. Differences among patients according to the EAT-10 screening

	EAT-10 positive	EAT-10 negative	p
Female sex	66.7 %	46.6 %	0.025
Age (years)	76.9 (15.4)	73.5 (18.1)	0.274
CCI (points)	0.5 (0.9)	0.6 (0.8)	0.598
High CCI (> 3 points)	85.7 %	78.4 %	0.310
MUST (points)	0.5 (0.9)	0.6 (0.8)	0.598

CCI: Charlson comorbidity index; EAT-10: Eating Assessment Tool; MUST: Malnutrition Universal Screening Tool. Quantitative data are presented as mean (standard deviation).

DISCUSSION

In this cross-sectional study carried out in the Neurology and Internal Medicine hospitalisation units of a university hospital, the prevalence of patients at risk of dysphagia detected with the EAT-10 was 26.6 %, a higher rate than that obtained in the PREDyCES® study (20.5 %) (7). There were no differences in the prevalence of dysphagia between the two studied departments, in which presumably we can find patients at higher risk of swallowing disorders due to ageing, frailty, and neurological disorders (e.g., dementia, cerebrovascular disease). According to studies, any type of cognitive impairment is associated with dysphagia. The prevalence of swallowing disorders in patients with dementia could reach 93 % (3,15). In cerebrovascular disease, it is the most common cause of admission to the Neurology hospitalisation unit; prevalence increases to 37-45 % when questionnaires are used, and to 78 % when assessed with videofluoroscopy (3). The questionnaire was more frequently positive in women, unlike what was found in the study that validated the Spanish EAT-10, which could be possibly related to older age and higher morbidity (12).

The EAT-10 was realisable in 80.6 % of the studied patients. In the Neurology unit, 88.9 % of subjects could answer the questionnaire, compared to 76.7 % in the Internal Medicine unit. No differences were found between men and women when they were able to answer the questionnaire; however, differences were found in age. Thus, patients who cannot fill in the screening test were older and had a higher score in the CCI as well. Patients who could not perform the test exhibited a higher mortality rate than those who were able to fill it out. Many studies that used the EAT-10 as a screening tool excluded patients with cognitive impairment or serious neurological disease who were unable to answer the items in the questionnaire, so they did not test the actual feasibility of this tool in a hospital environment (12,16-18).

In this study, malnutrition, in-hospital mortality, mean hospital stay, and 30-day readmissions were not increased in patients with a positive EAT-10. These results are in contrast with what previous studies showed (3,16-18). In patients who were unable to perform the EAT-10 there was a higher mortality rate than in those who could perform it. Together, these data indicate that the EAT-10 cannot be used in patients at higher risk of worse clinical outcomes. In contrast, MUST could be performed in all the recruited patients. Both the weight and height of the patient can be indicated by the patient himself or by their family or caregiver. If they fail to provide the data, height can be estimated by the length of the ulna.

The study was carried out at a time of the year with a large number of patients admitted for influenza, so perhaps, if the study were repeated at another time of the year, the prevalence of dysphagia in Internal Medicine could change, and even the feasibility of the test, since there would be other types of patients admitted to this unit. In addition, a larger sample size would possibly reveal significant differences among some parameters related to dysphagia. As a strength of the study, we can highlight that patients with potential risk of dysphagia were not excluded, so the results show the real utility of the EAT-10 in the analysed hospital units.

CONCLUSIONS

The prevalence of dysphagia risk as estimated with the EAT-10 in our Neurology and Internal Medicine hospitalisation units is 26.6 %. The questionnaire was answered only by 80.6 % of the patients in these departments, and it was less frequently practicable in older patients, patients with more comorbidity, and patients at risk of malnutrition. The results of the EAT-10 were not related to worse clinical outcomes, such as hospital stay, mortality, and readmission. We can conclude that the EAT-10 has limited utility for inpatients at Internal Medicine and Neurology wards, and other strategies focused on the detection of dysphagia should be studied.

REFERENCES

1. Hennessy M, Goldenberg D. Surgical anatomy and physiology of swallowing. *Oper Tech Otolaryngol* 2016;27(2):60-6. DOI: 10.1016/j.otot.2016.04.002
2. Clavé P, Terré R, Kraa M, Serra M. Approaching oropharyngeal dysphagia. *Rev Esp Enf Dig* 2004;96(2):119-31. DOI: 10.4321/S1130-01082004000200005
3. Bajjens L, Clavé P, Cras P, Eckberg O, Forster A, Kolb GF, et al. European Society for Swallowing Disorders, European Union Geriatric Medicine Society white paper: oropharyngeal dysphagia as a geriatric syndrome. *Clin Interv Aging* 2016;11:1403-28. DOI: 10.2147/CIA.S107750
4. Rofes L, Arreola V, Almirall J, Cabré M, Campins L, García-Peris P, et al. Diagnosis and management of oropharyngeal dysphagia and its nutritional and respiratory complications in the elderly. *Gastroenterol Res Pract* 2010;2011:1-13. DOI: 10.1155/2011/818979
5. Clavé P, Shaker R. Dysphagia: Current reality and scope of the problem. *Nat Rev Gastroenterol Hepatol* 2015;12(5):259-70. DOI: 10.1038/nrgastro.2015.49
6. Roy N, Stemple J, Merrill RM, Thomas L. Dysphagia in the elderly: preliminary evidence of prevalence, risk factors, and socioemotional effects. *Ann Otol Rhinol Laryngol* 2007;116(11):858-65. DOI: 10.1177/000348940711601112
7. Álvarez Hernández J, León Sanz M, Planas Vilá M, Araujo K, García de Lorenzo A, Celaya Pérez S. Prevalence and costs of malnutrition in hospitalized dysphagic patients: a subanalysis of the PREDyCES study. *Nutr Hosp* 2015;32(4):1830-6.
8. European Society for Swallowing Disorders. Position statements and meeting abstracts. *Dysphagia* 2013;28:280-335.
9. Warren JL, Bacon WE, Harris T, McBean AM, Foley DJ, Phillips C. The burden and outcomes associated with dehydration among US elderly, 1991. *Am J Public Health* 1994;84(8):1265-9. DOI: 10.2105/AJPH.84.8.1265
10. Whelan K. Inadequate fluid intakes in dysphagic acute stroke. *Clin Nutr* 2001;20(5):423-8. DOI: 10.1054/clnu.2001.0467
11. Almirall J, Cabré M, Clavé P. Aspiration pneumonia. *Med Clin (Barc)* 2007;129(11):424-32. DOI: 10.1157/13110467
12. Burgos R, Sarto B, Seguro H, Romagosa A, Puiggrócs C, Vázquez C, et al. Traducción y validación de la versión española del EAT-10 (Eating Assessment Tool-10) para el cribado de disfagia. *Nutr Hosp* 2012;27(6):2048-54.
13. BAPEN: The MUST toolkit [Internet]. Redditch, Worcestershire, United Kingdom: British Association for Parenteral and Enteral Nutrition; 2018 Jul 4 [Citado 22 diciembre 2019]. Disponible de: <https://www.bapen.org.uk/screening-and-must/must/must-toolkit>
14. Charlson ME, Pompei P, Ales KL, MacKenzie CR. A new method of classifying prognostic comorbidity in longitudinal studies: development and validation. *J Chronic Dis* 1987;40(5):373-83. DOI: 10.1016/0021-9681(87)90171-8
15. Paranjani S, Paranjani N, Wright S, Chandra S. A nationwide study of the impact of dysphagia on hospital outcomes among patients with dementia. *Am J Alzheimers Dis Other Dement* 2017;32(1):5-11. DOI: 10.1177/1533317516673464
16. Mañas-Martínez A, Búcar-Barjud M, Campos-Fernández J, Gimeno-Orna JA, Pérez-Calvo J, Ocón-Bretón J. Asociación de un cribado positivo para disfagia con el estado nutricional y la mortalidad a largo plazo en pacientes ancianos hospitalizados. *Endocrinol Diabetes y Nutr* 2018;65(7):402-8. DOI: 10.1016/j.endinu.2018.02.004
17. Izaola O, Gómez Hoyos E, López JJ, Ortola A, Torres B, Primo D, et al. The 10-item eating assessment tool is associated with nutritional status, mortality and hospital stay in the elderly individuals requiring hospitalization with acute diseases. *Nutr Hosp*. 2018;35(4):827-32.
18. Matsuo H, Yoshimura Y, Ishizaki N, Ueno T. Dysphagia is associated with functional decline during acute-care hospitalization of older patients. *Geriatr Gerontol Int* 2017;17(10):1610-6.