Comparison of NRS 2002 and PG-SGA for the assessment of nutritional status in cancer patients.

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Abstract

Purpose: To evaluate the use of the Patient-Generated Subjective Global Assessment (PG-SGA) and the Nutrition Risk Screening 2002 (NRS 2002) for the assessment of nutritional status in patients with common malignant tumors.

Methods: Patients hospitalized in Huizhou Central People's Hospital from December 2012 to May 2014 were enrolled. The diagnosis with cancer was confirmed by pathological examination and patients received chemotherapy/radiotherapy or surgery. The patients were interviewed by a trained surgeon using NRS 2002 and PG-SGA. Fasting venous blood samples were taken from all enrolled patients, and serum albumin and prealbumin levels were measured. NRS 2002 score \geq 3 indicated malnutrition risk, and PG-SGA score \geq 4 indicated malnutrition. The correlations among the scores, serum albumin and prealbumin levels, the body mass index (BMI), the length of hospital stay, and hospitalization cost were analyzed.

Results: 482 patients participated in this study, 242 (50.2%) had NRS 2002 score \geq 3 and 359 (74.5%) had PG-SGA score \geq 4. The detection rate of PG-SGA was significantly higher than that of NRS 2002. In patients with serum albumin <35 g/L and prealbumin <0.2 g/L, the detection rates of NRS 2002 and PG-SGA were 67.8% and 93.4%, and 66.4% and 88.8%, respectively, with significant difference. Both NRS 2002 and PG-SGA scores were associated with albumin, pre-albumin, and BMI (P<0.05).

Conclusions: PG-SGA has greater sensitivity than NRS 2002 in assessing nutritional status for patients with common malignancies and is more appropriate for nutritional assessment of cancer patients.

Keywords: NRS 2002, PG-SGA, Cancer, Nutritional assessment.

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Introduction

The incidence of malnutrition among hospitalized patients with cancer is high and varies dependent on the tumor location, tumor stage, therapeutic method, and nutritional assessment tool used. Several studies have reported that the occurrence rate of malnutrition related to cancer is 30-87% [1-8]. Malnutrition extends hospital stay, increases health care cost, reduces the quality of life, increases operative risks and complications, and impairs tolerance to chemotherapy and radiotherapy [9]. Some studies have suggested that approximately 20-50% of the mortality factors of cancer patients are related to malnutrition rather than to the cancer itself [10]. The quality of life and prognosis of patients with tumors will be improved and the complications reduced if their nutritional status is improved.

Nutritional therapy for patients with cancer has gradually received clinical attention and become a major component of

comprehensive treatment of patients with malignant tumors. However, nutritional therapy only benefits patients who have a nutritional risk or malnutrition. Furthermore, excessive nutritional treatment may expose patients to infection; aggravate their economic burden, and waste medical resources. Therefore, current consensus is that nutritional therapy is beneficial for patients with cancer who have a nutritional risk or malnutrition but is not needed for patients without a nutritional risk or malnutrition. Therefore, timely, accurate, and dynamic assessment of nutritional status is important for nutritional cancer treatment.

There is currently no absolute "gold standard" to determine whether a patient has malnutrition. The diagnosis of nutritional status can be divided into two stages: nutritional screening and nutritional assessment. The aim of the former is to identify malnourished patients or patients with a nutritional risk, especially those who have not yet shown symptoms of malnutrition but who have been found to have a nutritional risk. Nutritional therapy should be combined with clinical treatment for these patients and should be carried out during office visits or admission to a hospital. Nutritional assessment is more extensive for admitted patients; it should comprehensively evaluate nutritional status according to various scale scores, determine whether the patient has malnutrition and its complications, estimate the patient's nutritional requirements, develop a nutritional therapy plan, and assess the efficacy of nutritional therapy [11]. Nutritional screening has been recommended as a routine for patients admitted to a hospital [12-14]. The European Society for Clinical Nutrition and Metabolism suggested the adoption of the Nutrition Risk Screening 2002 (NRS 2002), the Mini Nutritional Assessment, and the Malnutrition Universal Screening Tool [13]. However, those nutritional screening tools were developed for patients without cancer, and their applicability for patients with cancer remains to be determined. In China, the Chinese Society of Clinical Oncology Specialist Committee of Nutritional Therapy for Cancer recommended that once patients with malignant tumors have clear diagnoses, they should immediately undergo nutritional risk screening. The widely used nutritional assessment tools for patients with malignancy are Patient-Generated Subjective Global Assessment (PG-SGA) and NRS 2002 [12]. This study was performed to compare the use of NRS 2002 and PG-SGA in the assessment of nutritional status of patients with common malignant tumors.

Patients and Methods

Subjects

Patients with common malignant tumors hospitalized in Huizhou Central People's Hospital from December 2012 to May 2014 were enrolled in this study. The inclusion criteria were as follows: age between 18 and 90 years; a diagnosis of malignancy on the basis of pathological examination; the ability to answer questions without any communication obstacles; the willingness to volunteer for the study; a common malignancy (lung, gastric, liver, colon/rectum, breast, esophageal, cervical, endometrial, nasopharyngeal, pancreatic, ovarian, prostatic, bladder, and brain cancer; malignant lymphoma; and leukemia); and a hospital stay of more than 1 day with no operation before the next morning.

A total of 482 cancer patients were selected, including 255 cases of colon/rectum cancer, 61 of lung cancer, 58 of gastric cancer, 31 of breast cancer, 22 of esophageal cancer, 15 of malignant lymphoma, 10 of cervical cancer, 11 of ovarian cancer, 6 of nasopharyngeal carcinoma, 4 of liver cancer, 3 of bladder cancer, 3 of endometrial cancer, 2 of pancreatic cancer, and 1 of prostatic cancer. 206 were male (42.7%) and 276 were female (57.3%). Their average age was 57.23 ± 12.19 years.

Methods

A single surgeon from a Class III Grade I hospital who had received standard nutritional screening training examined all of

the eligible patients. The examination included NRS 2002 and PG-SGA and was completed for each patient within 24 h of hospitalization. The height and body weight were measured to the nearest 0.5 kg and 0.5 cm, respectively, and the patients were in patient dress, did not wear shoes, and fasted that morning. In addition, the changes in body weight during the previous 3 months and the patient's diet for the previous 2 weeks were recorded. The details of nutritional status assessment standards were as follows:

(1) A body mass index (BMI) $<18.5 \text{ kg/m}^2$ was considered to be low, a BMI of 18.5-23.9 kg/m² was considered normal, and a BMI of 24.0-28 kg/m² was considered overweight [15].

(2) The patients' nutritional status and nutritional risk were assessed according to NRS 2002, including the severity of illnesses, diet, recent changes in body weight, and somatometry; the total scores ranged from 0 to 7. The patients at least 70 years old received one additional point. The patients were divided into two groups, a group with no nutritional risk (score<3) and a group with a nutritional risk (score \geq 3). Moreover, patients with a BMI <18.5 kg/m² were automatically given a score of 3 and assessed as having malnutrition [15].

(3) PG-SGA was divided into two parts and completed independently by the patients and checked by the physician. The patients provided information about historical symptoms, body weight, current mobility, and diet. Total scores were calculated and the results fell into three grades: grade A (total: 0-3) with normal nutrition; grade B (total: 4-8) with moderate malnutrition; and grade C (total: \geq 9) with severe malnutrition [16].

In addition to completing the questionnaire, in the early morning of the second day after admission, fasting venous blood samples were obtained from each enrolled patient to measure serum albumin and pre-albumin levels.

Statistical analysis

SPSS Statistics 20.0 software package was used for statistical analysis. Rank sum tests, t-tests, and chi-square tests were used to compare the averages and detection rates of the two groups. A matching McNemar chi-square test was used to compare the relevance coincidence of two scales with the same group of subjects (e.g., BMI<18.5). Pearson/Spearman correlation coefficient analysis was conducted for the analysis of the BMI and biochemical indexes (albumin, pre-albumin). The correlation coefficient average was compared by estimating the 95% confidence interval by bootstrap. The inspection level of the hypothesis test was assumed to be α =0.05. In addition, Kolmogorov-Smirnov (n>50) and Shapiro-Wilk (n ≤ 50) tests were used to check normality of quantitative data; the test of normality inspection level was set at α =0.10. P<0.05 was considered as statistically significant.

Results

General data of the patients

Of the 482 patients who participated in the study, 242 (50.2%) scored \geq 3 on NRS 2002 (147 male and 95 female), and 359 patients (74.5%) scored \geq 4 on PG-SGA (225 male and 134 female). The patients in the group with malnutrition risk or malnutrition were older than those in the group without malnutrition risk or malnutrition. The gender distribution of the groups when sorted by a PG-SGA score \geq 4 was statistically significant (P<0.05), whereas that of the groups when classified according to NRS 2002 score \geq 3 was not (P>0.05).

Table 1. Detection rate of NRS 2002 and PG-SGA in cases with serum albumin <35 g/L.

NRS 2002 ≥ 3	PG-SGA≥4		Р
	Malnutrion 1	No malnutrion 0	
Malnutrition risk 1	59 (65.6)	2 (2.2)	<0.001
No malnutrition risk 0	25 (27.8)	4 (4.4)	

Table 2. Detection rate of NRS 2002 and PG-SGA in cases with serum prealbumin <0.2 g/L.

NRS 2002 ≥ 3	PG-SGA≥4		Р	
	Malnutrition 1	No malnutrition 0		
Malnutrition risk 1	140 (62.8)	8 (3.6)	<0.001	
No malnutrition risk 0	58 (26.0)	17 (7.6)	_	

Table 3. Correlation of indexes with NRS 2002 and PG-SGA.

	NRS 2002 Score	PG-SGA Score	Р
	(bootstrap 95% CI)	(bootstrap 95% CI)	
Albumin	-0.297	-0.355	>0.05
	(-0.335, -0.157)**	(-0.412, -0.252)**	
Prealbumin	-0.360	-0.396	>0.05
	(-0.404, -0.242)**	(-0.424, -0.275)**	
BMI	-0.378	-0.257	>0.05
	(-0.462, -0.304)**	(-0.344, -0.170)**	

Spearman correlation coefficient was adopted, and comparison of correlation coefficient average was performed by estimating the 95% confidence interval of the correlation coefficient by bootstrap (midpoint crossing indicated P>0.05; no crossing P<0.05). **Hypothesis testing of each correlation coefficient P<0.01.

Comparison of positive rate between NRS 2002 and PG-SGA

For all 482 patients, the incidence of malnutrition risk was 50.2% according to NRS 2002, while the rate of malnutrition was 74.5% according to PG-SGA. In addition, we found that the positive rate of PG-SGA was significantly higher than that of NRS 2002 in these patients (P<0.05). When a PG-SGA score \geq 4 was set as the standard for a diagnosis of malnutrition, 359 patients were determined to have

Table 4. Comparison of Length of Stay (LOS) and costs of patients

 evaluated by NRS 2002.

	Operation Group		Chemotherap Radiotherapy	•
	1. Positive ⊼±s/M	Neative x ± s/ M	Positive $\overline{x} \pm s/M$	Negative x ± s/ M
	(P25-P75)	(P25-P75)	(P25-P75)	(P25-P75)
LOS (days)	17 (13-21)	15 (13-18.3)	6(3.8-12)	4(3-6)
Z	1.81		2.68	
Р	0.07		0.007	
Costs (×10,000 yuan)	4.0 (2.8-5.1)	3.7 (2.3-4.6)	0.8 (0.5-1.4)	0.7 (0.5-0.9)
Z	1.60		2.09	
Р	0.11		0.036	

Table 5. Comparison of Length of Stay (LOS) and costs of patients evaluated by PG-SGA.

	Operation Grou			
	Operation Group		Chemotherapy Radiotherapy Group	
	Positive x ± s/ M (P25-P75)	Negative $\overline{x} \pm s/$ M (P25-P75)	Positive	Negative x ± s/ M (P25-P75)
LOS (days)	17 (13-21)	15 (13.5-19.5)	5 (4-8.5)	4 (3-6)
Z	0.66		4.08	
Р	0.51		<0.001	
Costs (×10,000 yuan)	4.0 (2.8-5.0)	3.6 (2.5-4.5)	0.8 (0.5-1.2)	0.7 (0.5-0.9)
Z	0.98		2.36	
Р	0.33		0.019	

Correlation of NRS 2002 and PG-SGA scores with serum albumin and prealbumin levels and body mass index

In the patients with serum albumin <35 g/L, the detection rates of NRS 2002 and PG-SGA were 67.8% and 93.4%, respectively, but in the patients with serum prealbumin <0.2 g/L, the detection rates were 66.4% and 88.8%, respectively, the differences were statistically significant. The probability of detecting malnutrition in patients with serum albumin <35 g/L and serum prealbumin <0.2 g/L was higher when PG-SGA scale was used (Tables 1 and 2). The correlation of NRS 2002 and PG-SGA scores with serum albumin and prealbumin levels and BMI was negative (P<0.05). The average difference between the scores and the biochemical index had no statistical significance, indicating that the correlation intensity was the same (Table 3).

Correlation of nutrition scores with hospital stay and hospitalization costs

In chemotherapy or radiotherapy group, the differences in the length of hospital stay and the hospitalization costs between patients with and without malnutrition risk and nutrition scores were significant (P<0.05). Therefore, NRS 2002 and PG-SGA scores could predict patients' length of hospital stay and hospitalization costs. However, in surgery group, the differences between the malnourished and well-nourished patients evaluated by NRS 2002 and PG-SGA scores were not significant (Tables 4 and 5).

Discussion

The incidence of malnutrition in patients with malignant tumors has been reported to be 30-87% [1-8]. In this study, we assessed the nutritional status of patients with malignancies, and the incidence of malnutrition was 74.5% evaluated by PG-SGA but was only 50.2% evaluated by NRS 2002. The incidence of malnutrition or nutritional risk varies because of different subjects, nutritional screening or assessment tools, tumor locations, stages, and therapeutic aims or methods. There is currently no gold standard for a diagnosis of malnutrition. The ability to estimate a patient's nutritional status quickly, accurately, conveniently, and non-invasively is the first step in nutritional cancer therapy.

Traditional indices for nutritional status include anthropometric indicators, such as BMI, triceps skin fold thickness, mid-arm muscle circumference, and grip strength, and biochemical indicators, such as serum levels of albumin, pre-albumin, transferrin, and C-reactive protein and total lymphocyte counts. However, each of those nutritional assessment indicators has limitation and could not accurately and comprehensively reflect patients' nutritional status. Commonly used nutritional screening tools include NRS 2002, Subjective Global PG-SGA, Assessment, Mini-Nutritional Assessment, Malnutrition Universal Screening Tool, Nutrition Risk Index, and Malnutrition Screening Tool. These familiar tools have advantages and disadvantages, but not all of them can be used for patients with malignancies. American Society for Parenteral and Enteral Nutrition and European Society for Parenteral and Enteral Nutrition recommend SGA as a nutritional assessment tool [13,17]. American Nutrition Association recommends PG-SGA as a nutritional screening tool for patients with malignancy. Chinese Society of Clinical Oncology Specialist Committee of Nutritional Therapy for Cancer recommends PG-SGA and NRS 2002 as the nutritional assessment tools for patients with malignancy [11]. No consensus has been achieved about which tool is the most suitable for patients with malignant tumors.

Bauer et al. used SGA and PG-SGA to assess nutritional status. When SGA was set as the standard, the sensitivity of PG-SGA was 98% and the specificity 82%. They concluded that PG-

SGA was a fast, effective, and reliable tool for the assessment of nutritional status of patients with malignancies [18]. Additional studies have shown that PG-SGA score is closely related to weight loss, the length of hospital stay, quality of life, and energy intake of the patients [19,20]. In this study, PG-SGA and NRS 2002 scores both had weakly positive correlations with the length of hospital stay and hospitalization costs of cancer patients who were hospitalized for chemotherapy or radiotherapy, but were not correlated with those variables in patients hospitalized for surgery. These results may be associated with the fact that comprehensive standardized surgery has not been performed at our hospital, and we have different standards for hospital discharge and surgical methods.

An international multi-center trial validated that NRS 2002 score was closely related to the length of hospital stay, the incidence of complications, and the death rate in patients undergoing gastrointestinal surgery [3]. Schiesser et al. suggested that NRS 2002 score could be applied to predict the incidence of postoperative complications and even postoperative death rate [21]. In this study, the incidence of malnutrition in patients with common malignant tumors was 74.5% and 50.2% when PG-SGA and NRS 2002 were used for nutritional screening, respectively, and the difference was statistically significant. PG-SGA has greater sensitivity in nutritional assessment of patients with common tumors. When a PG-SGA score \geq 4 was set as a standard for the diagnosis of malnutrition, the sensitivity of an NRS 2002 score \geq 3 was 61.8% and the specificity was 83.7%.

PG-SGA is mainly based on diet, recent changes in body weight, clinical symptoms, physical examination, and some laboratory measurements. However, in this study PG-SGA score had a weak negative correlation with BMI and serum albumin and prealbumin levels. In contrast, NRS 2002 is based on BMI and albumin level. NRS 2002 has lower sensitivity than PG-SGA because the latter is based on objective laboratory data. In addition, the cost of using the PG-SGA as a nutritional assessment tool was lower, and it is more easily accepted by patients during outpatient reexamination. Nevertheless, PG-SGA failed to screen for malnutrition according to a BMI <18.5 (10.9%), serum prealbumin <35 g/L (6.6%), and serum prealbumin <0.2 g/L (11.2%) in potentially malnourished patients, which indicates that PG-SGA still has some flaws and remains to be improved.

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