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ABOUT COVER

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AIMS AND SCOPE

The primary aim of World Journal of Gastrointestinal Surgery (WJGS, World J Gastrointest Surg) is to provide scholars and readers from various fields of gastrointestinal surgery with a platform to publish high-quality basic and clinical research articles and communicate their research findings online.

WJGS mainly publishes articles reporting research results and findings obtained in the field of gastrointestinal surgery and covering a wide range of topics including biliary tract surgical procedures, biliopancreatic diversion, colectomy, esophagectomy, esophagostomy, pancreas transplantation, and pancreatectomy, etc.

INDEXING/ABSTRACTING

The WJGS is now abstracted and indexed in Science Citation Index Expanded (SCIE, also known as SciSearch®), Current Contents/Clinical Medicine, Journal Citation Reports/Science Edition, PubMed, PubMed Central, Reference Citation Analysis, China National Knowledge Infrastructure, China Science and Technology Journal Database, and Superstar Journals Database. The 2022 Edition of Journal Citation Reports® cites the 2021 impact factor (IF) for WJGS as 2.505; IF without journal self cites: 2.473; 5-year IF: 3.099; Journal Citation Indicator: 0.49; Ranking: 104 among 211 journals in surgery; Quartile category: Q2; Ranking: 81 among 93 journals in gastroenterology and hepatology; and Quartile category: Q4.

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ORIGINAL ARTICLE

Retrospective Cohort Study

Nomogram to predict permanent stoma in rectal cancer patients after sphincter-saving surgery

Chih-Yu Kuo, Po-Li Wei, Chia-Che Chen, Yen-Kuang Lin, Li-Jen Kuo

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Abstract

BACKGROUND

Approximately 20 percent of patients with a tumour localized in the low rectum still encounter the possibility of requiring permanent stoma (PS), which can cause drastic changes in lifestyle and physical perceptions.

AIM

To determine the risk factors for PS and to develop a prediction model to predict the probability of PS in rectal cancer patients after sphincter-saving surgery.

METHODS

A retrospective cohort of 421 rectal cancer patients who underwent radical surgery at Taipei Medical University Hospital between January 2012 and December 2020 was included in this study. Univariate and multivariate analyses were performed to identify the independent risk factors for PS. A nomogram was developed according to the independent risk factors obtained in the multivariate analysis. The performance of the nomogram was assessed using a receiver operating characteristic curve and a calibration curve.

RESULTS

The PS rate after sphincter-saving surgery was 15.1% (59/391) in our study after a



median follow-up of 47.3 mo (range 7-114 mo). Multivariate logistic regression analysis demonstrated that local recurrence, perirectal abscess, anastomosis site stenosis, perineural invasion, tumor size and operative time were independent risk factors for PS. These identified risk factors were incorporated into the nomogram, and the concordance index of this model was 0.903 (95% CI: 0.851-0.955). According to the calibration curves, the nomogram represents a perfect prediction model.

CONCLUSION

Several risk factors for PS after sphincter-saving surgery were identified. Our nomogram exhibited perfect predictive ability and will improve a physician's ability to communicate the benefits and risks of various treatment options in shared decision making.

Key Words: Nomogram; Permanent stoma; Risk factor; Shared decision making; Sphincter-saving operation; Rectal cancer

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Core Tip: Approximately 20 percent of patients with a tumour localized in the low rectum still encounter the possibility of requiring permanent stoma (PS), which can cause drastic changes in lifestyle and physical perceptions. The study aimed to identify the risk factors for PS in rectal cancer patients after sphincter-saving surgery. Our results showed that the predictive models constructed by clinicopathological features exhibited perfect predictive ability and will allow physicians to inform patients about the possibility of PS prior to surgery.

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INTRODUCTION

Shared decision making (SDM) is a structured process that incorporates available scientific evidence, patient values, preferences, and life situation into screening decisions[1]. The benefits of SDM include improved medical quality, improved patient satisfaction, increased patient compliance to medical treatment, and reduced patient anxiety during treatment; SDM also helps patients understand the issues with which they should be familiar before they undergo treatment[2,3]. This discussion is particularly important in cancer treatment since patients are often provided with more than one available treatment strategy[4].

Despite innovative advancements, the management of rectal cancer remains a formidable endeavor, especially distally located rectal cancer[5]. It is extremely challenging to work in the low and narrow pelvis with laparoscopic straight instruments. Male sex, high body mass index (BMI), low rectal cancer, bulky tumor, and advanced stage are well known to increase the technical difficulty[6]. Moreover, a certain percentage of anastomosis-related complications will occur after colorectal surgery. Anastomosis complications, such as anastomotic leakage, perirectal abscess, and anastomotic stenosis, often lead to permanent stoma (PS). According to previous studies, 3%-24% of rectal cancer patients experience anastomosis complications after sphincter-saving surgery^[7-9].

A nomogram is a statistical tool that can transform a complex regression equation result into a simple and visual graph[10]. Thus, the results of prediction models become more readable and valuable. The aim of this study was to develop and validate a nomogram that incorporated both the clinical and pathologic risk factors for individual preoperative prediction of PS in patients with rectal cancer who underwent sphincter-saving surgery.

MATERIALS AND METHODS

Patient selection

We retrospectively reviewed records of patients with rectal cancer who underwent surgery at Taipei Medical University Hospital from January 2012 to December 2020. The inclusion criteria were as follows: (1) Patients older than 18 years; (2) Underwent radical surgery [low anterior resection,



intersphincteric resection, or abdominoperineal resection (APR)]; (3) Pathological diagnosis of malignancy; and (4) lesion located within 12 cm from the anal verge. The exclusion criteria were as follows: (1) Patients with stage IV disease; (2) Those who underwent emergency surgery; and (3) Those who underwent other organ resection during primary surgery. Defunctioning stoma was performed if any of the following conditions applied: (1) Positive air leak test; (2) Patient received preoperative chemoradiotherapy (CRT); (3) Anastomosis had tension or poor blood supply; (4) Presence of incomplete anastomotic ring; (5) Very low anastomosis; (6) Patients' clinical condition indicated defunctioning stoma; and (7) The surgeon elected to perform this procedure based on his/her experience. The condition of PS included non-reversal temporary stoma and stoma re-creation after reversal surgery (Figure 1). This study was approved by the Joint Institutional Review Board of Taipei Medical University (TMU-JIRB No: N202103023).

Data collection and definition of postoperative complications

Patient demographics and potential risk factors for PS were retrospectively collected and included sex, age, BMI, comorbidities (diabetes mellitus, hypertension, heart disease, chronic obstructive pulmonary disease, chronic kidney disease, liver disease), smoking status, clinical tumor-node-metastasis stage, whether the patient received neoadjuvant CRT, American Society of Anesthesiologists (ASA) score, tumor location (distance from the anal verge), tumor markers, such as carcinoembryonic antigen (CEA), preoperative lab data (hemoglobin and albumin), surgical approach, blood loss, operative time, stoma status, postoperative hospital stay, histologic grade, lymph vascular invasion, perineural invasion, circumferential resection margin (CRM) status, whether the patient received adjuvant chemotherapy, local recurrence, postoperative leakage, anastomosis site stenosis, perirectal abscess, and recto-visceral fistula.

Anastomotic leakage was defined as peritonitis that was clinically apparent (discharge containing pus or fecal material) or radiologically evident (contrast leakage or abscess around the anastomosis). Perirectal abscess (late anastomotic leak) was defined as a leak that was diagnosed more than 30 d after surgery. Anastomotic stricture was defined as the inability of a 12-mm proctoscope to pass through the anastomosis. A PS was defined when a closure procedure had not been performed or scheduled within the follow-up period (median, 47 mo; range, 7-114 mo).

Postoperative follow-up

Patients were followed-up every 3 mo during the first 2 years and then every 6 mo until the fifth year. Clinical examination and serum CEA testing were performed during each follow-up visit. Surveillance colonoscopy was performed within 12 mo after the initial surgery and every other year thereafter. Contrast-enhanced computed tomography scan of the thorax, abdomen, and pelvis was performed annually for 3 years and subsequently only when clinically indicated.

Data and risk factor analysis

Categorical variables are presented as counts and percentages, while continuous variables are depicted as the mean ± SD. Differences between both groups were assessed with the chi-square test or Fisher exact test depending on the sample size. Univariate analyses for risk factors related to a PS were performed. Multivariate logistic regression was conducted to identify the independent risk factors. A two-tailed P value < 0.05 was considered statistically significant. Statistical analyses were performed using SAS v9.4 (Cary, NY)

Nomogram development

Statistical analyses of the nomogram were conducted using SAS v 9.4 and R (ver. 3.0.1, Vienna, Austria). The rms package in R was used to plot the nomogram as a graphical calculating device that visualizes an approximation of mathematical function. Features of the nomogram are based on logistic regression models. The nomogram function in the rms package was adopted to generate nomograms from the fitted logistic statistical model. As a result, the performance of the nomogram is dependent on the regression models. We assessed the predictive power of the nomogram using receiver operating characteristic curve analysis. Calibration curves were used to explore the performance of the nomogram.

RESULTS

Patient characteristics

In all, 421 patients who underwent radical surgery are included in our study, including 391 (92.9%) who underwent sphincter-saving surgery and 30 (7.1%) who underwent APR. Moreover, 136/391 (34.8%) patients who underwent a sphincter-saving procedure had a temporary stoma after primary surgery. After a median follow-up of 47.3 mo (range 7–114 mo), 59/391 (15.1%) patients were confirmed to have PS, and the details of the stoma condition are shown in Figure 1. According to our data, 332 patients are in the stoma free group, while 89 patients are in the PS group. In summary, the PS rate after sphincter-





Figure 1 Study flow chart.

saving surgery at our hospital from January 2012 to December 2020 is 15.1% (59/391), and the total sphincter-saving rate is 78.9% (89/421). All data compared between the stoma free and PS groups are presented in Table 1.

Feature selection

Data from the univariate and multivariate analyses for PS are provided in Table 2. According to the multivariate logistic regression analysis, seven features were significantly related to PS. The independent risk factors for PS by multivariate logistic regression were local recurrence [odd ratio (OR), 111.578; 95% CI: 7.964-> 999; P < 0.001], perirectal abscess (OR, 369.397; 95% CI: 17.137-> 999; P < 0.001), anastomosis site stenosis (OR, 211.256; 95% CI: 13.705-> 999; P < 0.001), perineural invasion (OR, 7.674; 95% CI: 1.138-51.745; P = 0.036), tumor size (OR, 1.076; 95% CI: 1.015-1.14; P = 0.014), liver disease (OR, 0.054; 95% CI: 0.004-0.698; P = 0.025), and operative time (min) (OR, 1.008; 95% CI: 1.002-1.014; P = 0.01). We excluded liver disease because of OR < 1. Thus, these six variables were selected to construct the nomogram.

Construction of the nomogram

The prognostic nomogram that integrated all potential risk factors for PS in the cohort is shown in Figure 2. The nomogram model was validated by computing the concordance index (C-index) of the nomogram sample. The nomogram provides a visualization of accumulated risk by mapping the predicted probabilities into points on a scale from 0 to 1 in a graphical interface. The total points accumulated by each covariate correspond to the predicted probability in a given patient. To further illustrate this, the point system functions by ranking the effect estimates, regardless of statistical significance, and this ranking is influenced by the presence of other covariates. Despite statistical significance, the risk factor whose absolute value has the largest regression coefficient will be assigned 100 points on the scale, while the remaining variables are assigned a smaller number of points proportional to their effect size. As shown in Figure 2, perirectal abscess has the highest effect, and thus, this variable is assigned 100 points. Whereas a patient with perirectal abscess would be assigned 100 points, a patient with perirectal abscess would be assigned 100 points. For example, a patient with perirectal abscess, perineural invasion, and a tumor size of 20 mm would be assigned 150 points overall, which is mapped to an approximate predicted probability of 70%.

Table 1 Comparison of patient-related characteristics between the stoma free and permanent stoma groups					
Characteristic	Stoma free (<i>n</i> = 332)	Permanent stoma (<i>n</i> = 89)	<i>P</i> value		
Age, yr	60.78 ± 12.80	60.56 ± 12.60	0.888		
Sex (<i>n</i>)			0.716		
Male	196 (59.04%)	50 (56.18%)			
Female	136 (40.96%)	39 (43.82%)			
Body mass index, kg/m ²	24.00 ± 3.97	24.47 ± 4.32	0.331		
Comorbidity (n)					
DM	68 (20.48%)	14 (15.73%)	0.393		
Hypertension	103 (31.02%)	33 (7.08%)	0.339		
Heart disease	25 (7.53%)	8 (8.99%)	0.816		
COPD	2 (0.60%)	2 (2.25%)	0.421		
Chronic kidney disease	36 (10.84%)	9 (10.11%)	0.996		
Liver disease	39 (11.75%)	10 (11.24%)	1		
Smoker (n)	49 (14.76%)	9 (10.11%)	0.339		
Distance to anus verge, cm	7.06 ± 3.52	4.68 ± 3.96	< 0.001		
Clinical T stage (<i>n</i>)			0.002		
ТО	8 (2.41%)	1 (1.13%)			
T1	12 (3.61%)	1 (1.13%)			
T2	50 (15.06%)	8 (8.98%)			
Т3	218 (65.66%)	56 (62.92%)			
T4	20 (6.03%)	17 (19.10%)			
Data loss	24 (7.23%)	6 (6.74%)			
Clinical N stage (<i>n</i>)			0.44		
N0	108 (32.53%)	23 (25.84%)			
N1	100 (30.12%)	31 (34.83%)			
N2	100 (30.12%)	29 (32.59%)			
Data loss	24 (7.23%)	6 (6.74%)			
AJCC c TNM stage (<i>n</i>)			0.002		
Stage 0	8 (2.41%)	1 (1.13%)			
Stage I	49 (14.76%)	7 (7.86%)			
Stage II	52 (15.66%)	15 (16.85%)			
Stage III	199 (59.94%)	60 (67.42%)			
Data loss	24 (7.23%)	6 (6.74%)			
NACR (n)	222 (66.87%)	69 (77.53%)	0.026		
Hb, g/dL	12.78 ± 1.57	12.52 ± 1.72	0.169		
Albumin, g/dL	4.14 ± 0.36	4.08 ± 0.37	0.19		
CEA, ng/mL	4.81 ± 8.58	6.15 ± 8.69	0.198		
ASA score (<i>n</i>)			0.182		
Ι	26 (7.83%)	3 (3.37%)			
П	271 (81.63%)	73 (82.02%)			
III	30 (9.03%)	12 (13.48%)			
Data loss	5 (1.51%)	1 (1.13%)			

ASA: American Society of Anesthesiologists; COPD: Chronic obstructive pulmonary disease; CEA: Carcinoembryonic antigen; DM: Diabetes mellitus; AJCC: American Joint Committee on Cancer; Hb: Hemoglobin; NCRT: Neoadjuvant chemoradiotherapy; TNM: Tumor-node-metastasis; NACR: Neoadjuvant chemoradiation.



Figure 2 The established nomogram for predicting permanent stoma was developed by incorporating the following six parameters: Local recurrence, perineural invasion, tumor size (mm), rectal stenosis, perirectal abscess and operative time. First, the nomogram is used by giving each variable a score on the "Points" scale. The scores for all variables are then added to obtain the total score after which a vertical line is drawn from the "Total points" row to estimate the predicted probability of permanent stoma.

Validation and performance of the nomogram

After these six factors were incorporated, the nomogram achieved an outstanding C-index of 0.903 (95%CI: 0.851–0.955). The area under the receiver operating characteristic curve of our model (0.903) was higher than that of any single factor (local recurrence: 0.641; perineural invasion: 0.636; tumor size: 0.638; rectal stenosis: 0.645; perirectal abscess: 0.565; operative time: 0.669), which indicates that this model was more accurate than other models (Figure 3A). According to the calibration curve, the nomogram calibration plot demonstrated high reliability (Figure 3B). Predicted PS rates based on the model and the observed outcomes on calibration fit best at PS probability rates above 40%. However, the nomogram showed less consistent but high performance in the lower PS rate ranges, as the calibration curve fluctuates below 40% probability.

DISCUSSION

For the past three decades, dramatic improvements have been made in rectal cancer treatment, including advances in surgical pathology, refinements in surgical techniques and instrumentation, new imaging modalities, and the widespread use of neoadjuvant therapy^[11]. No matter how advanced the surgical technique, restoration of bowel continuity in patients with rectal cancer is still currently a challenge. Whenever possible, sphincter preservation should be sought. The sphincter can generally be preserved if the tumor can be resected with a 1-cm distal margin[12]. However, not all patients meet the surgical indications for sphincter-saving surgery. Even if patients undergo resection for rectal cancer, a common dilemma faced by surgeons is whether or not to create a defunctioning stoma. According to a recent meta-analysis published in 2017, which included ten studies consisting of 8568 patients, the rate of non-reversal of temporary stoma was 19% [13]. Patients still encounter multiple possible complications and the risk of perioperative mortality after surgery. Anastomotic complications are the primary reason for the necessity of a PS, and thus, these complications are more frequent than local recurrence [14-16]. Therefore, surgical decision making in the setting of rectal cancer is often complex, and detailed meetings for SDM are necessary. Patients and physicians arrive at treatment decisions together based on



Table 2 Risk factors for permanent stoma after sphincter-preserving surgery according to univariate and multivariable analyses						
Variable	Univariable analysis OR (95%Cl)	<i>P</i> value	Multivariable analysis OR (95%Cl)	P value		
Age, yr	0.99 (0.969-1.012)	0.369	0.959 (0.895-1.027)	0.232		
Sex (Ref. = female)						
Male	0.822 (0.472-1.443)	0.491	0.273 (1.044-1.7)	0.164		
Body mass index, kg/m ²	1.022 (0.953-1.092)	0.532	0.949 (0.807-1.116)	0.525		
DM (Ref. = No)						
Yes	0.792 (0.363-1.586)	0.532	0.307 (0.032-2.9)	0.303		
Hypertension (Ref. = No)						
Yes	1.229 (0.678-2.179)	0.488	0.819 (0.121-5.542)	0.838		
Heart disease (Ref. = No)						
Yes	0.893 (0.256-2.413)	0.84	0.229 (0.008-6.382)	0.385		
COPD (Ref. = No)						
Yes	5.795 (0.684-49.02)	0.082	451.125 (0.376->999)	0.091		
CKD (Ref. = No)						
Yes	0.931 (0.34-2.172)	0.878	0.421 (0.019-9.234)	0.583		
Liver disease (Ref. = No)						
Yes	1.179 (0.488-2.55)	0.694	0.054 (0.004-0.698)	0.025		
Smoker (Ref. = No)						
Yes	0.906 (0.379-1.932)	0.81	0.125 (0.007-2.148)	0.152		
Distance to anus verge, cm	0.838 (0.758-0.921)	< 0.001	0.834 (0.618-1.127)	0.238		
Clinical T stage (Ref. = T0)						
T1	< 0.001 (NA-4.239)	0.98	1.081 (< 0.001-> 999)	0.999		
T2	1.28 (0.193-25.357)	0.827	> 999 (< 0.001-> 999)	0.968		
T3	1.394 (0.246-26.24)	0.757	> 999 (< 0.001-> 999)	0.976		
T4	3.2 (0.468-64.31)	0.308	> 999 (< 0.001-> 999)	0.971		
Clinical N stage (Ref. = N0)						
N1	1.697 (0.831-3.568)	0.152	0.017 (< 0.001-> 999)	0.986		
N2	1.466 (0.701-3.129)	0.313	0.003 (< 0.001-> 999)	0.981		
AJCC c TNM stage (Ref. = Stage 0)						
Stage I	0.98 (0.139-19.76)	0.986	0.015 (< 0.001-> 999)	0.986		
Stage II	1.077 (0.159-21.492)	0.948	0.007 (< 0.001-> 999)	0.983		
Stage III	1.648 (0.291-30.993)	0.642	NA	NA		
Pre-operative CCRT (Ref. = No)						
Yes	1.332 (0.731-2.533)	0.364	1.873 (0.137-25.575)	0.638		
Hb, g/dL	0.987 (0.832-1.18)	0.887	1.404 (0.768-2.568)	0.27		
Albumin, g/dL	0.821 (0.361-1.928)	0.643	0.66 (0.041-10.497)	0.769		
CEA, ng/mL	1.011 (0.978-1.038)	0.443	0.936 (0.804-1.09)	0.396		
ASA score (Ref. = I)						
П	2.02 (1.046-3.891)	0.036	7.967 (0.64-99.127)	0.107		
III	NA	NA	NA	NA		
Surgical Approach way (Ref. $= 0$)						



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LPS (1)	NA	NA	> 999 (< 0.001-> 999)	0.859
Robotic (2)	NA	NA	> 999 (< 0.001-> 999)	0.872
Type of operation (Ref. = LAR)				
CAA	3.46 (1.958-6.266)	< 0.001	0.221 (0.027-1.796)	0.158
Estimated blood loss	1.002 (1-1.005)	0.072	1.001 (0.987-1.016)	0.889
Operative time	1.004 (1.002-1.007)	< 0.001	1.011 (1.001-1.02)	0.026
Histologic tumor grade (Ref. = Grade I)				
Grade II	1.622 (0.883-3.05)	0.124	1.203 (0.22-6.586)	0.831
Grade III	2.507 (0.645-8.203)	0.147	1.53 (0.038-61.785)	0.822
Tumor size, mm	1.026 (1.011-1.041)	< 0.001	1.076 (1.015-1.14)	0.014
Circumferential resection margin (Ref. = No)				
Yes	6.575 (2.955-14.604)	< 0.001	0.936 (0.064-13.699)	0.961
Lymph vascular invasion (Ref. = No)				
Yes	1.99 (1.071-3.617)	0.026	0.94 (0.132-6.715)	0.951
Perineural invasion (Ref. = No)				
Yes	3.085 (1.726-5.518)	< 0.001	7.674 (1.138-51.745)	0.036
Postoperative hospital stays	1.05 (1.02-1.083)	0.001	1.003 (0.911-1.104)	0.953
Postoperative chemotherapy (Ref. = No)				
Yes	1.907 (0.963-4.134)	0.079	4.281 (0.247-74.107)	0.318
Anastomosis site stenosis (Ref. = No)				
Yes	11.648 (5.499-25.374)	< 0.001	211.256 (13.705-> 999)	< 0.001
Local recurrence (Ref. = No)				
Yes	12.584 (5.874-27.885)	< 0.001	111.578 (7.964-> 999)	< 0.001
Postoperative leakage (Ref. = No)				
Yes	2.659 (0.982-6.557)	0.041	0.743 (0.047-11.833)	0.833
Perirectal abscess (Ref. = No)				
Yes	11.037 (3.22-43.367)	< 0.001	369.397 (17.137-> 999)	< 0.001
Recto visceral fistula (Ref. = No)				
Yes	44.557 (7.71-841.643)	< 0.001	> 999 (< 0.001-> 999)	0.963

ASA: American Society of Anesthesiologists; COPD: Chronic obstructive pulmonary disease; CKD: Chronic kidney disease; AJCC: American Joint Committee on Cancer; CCRT: Concurrent chemoradiotherapy; CEA: Carcinoembryonic antigen; CAA: Coloanal anastomosis; DM: Ciabetes mellitus; Hb: Hemoglobin; LPS: Laparoscopic surgery; LAR: Low anterior resection; OR: Odd ratio; TNM: Tumor-node-metastasis.

> clinical evidence within the context of a patient's personal preferences^[4]. Prior to surgery, patients should be informed that a certain percentage of postoperative anastomosis complications may occur, which in turn may lead to PS. In addition, the physician should carefully judge whether sphinctersaving surgery or APR should be performed. Many factors should be carefully considered, including the effects of neoadjuvant CRT, sufficient tumor resection margins, the patient's functional status/comorbid disease, and his or her personal wishes [17]. If patients who are at a higher risk of a PS after surgery can be identified, a physician's ability to communicate the benefits and risks of various treatment options in an SDM setting will be improved.

> Postoperative leakage and stricture are the most well-known anastomotic healing complications that have continued to plague surgeons. Both are primary reasons for PS. Although numerous studies have attempted to determine the healing process of colorectal anastomoses, the pathophysiologic mechanisms that govern the process of anastomotic regeneration remain poorly understood[18]. One major obstacle has been the lack of access to observe, sample, and analyze an anastomosis as it heals. Traditional dogma suggests that the most common factors implicated in anastomotic healing include





Figure 3 The nomogram calibration plot demonstrated high reliability. A: The area under the receiver operating characteristic curve for the nomogram was 0.903 (95%CI: 0.851-0.955); B: In the calibration curve, the predicted probability of stoma is plotted on the x-axis, while the actual probability of stoma is plotted on the y-axis. The dotted line represents an ideal nomogram, and the solid blue line represents the current nomogram.

tissue perfusion/ischemia, tissue tension, and patient nutritional status[19]. However, surgeons still cannot predict which anastomoses will leak or undergo stenosis. Even a well-constructed anastomosis by the most skilled surgeon with good perfusion and no tension can still develop leakage or stricture. Therefore, many retrospective studies attempt to determine the incidence and potential risk factors of anastomotic complications, which can help us predict the probability of PS. According to recent studies, the incidence of anastomotic leakage in the literature varies from 1% to 29%[20], and over half of patients with symptomatic anastomotic leakage will have PS[21,22]. A systematic search by Qu et al[23] indicated that common risk factors for anastomotic leakage include male gender, high BMI, high ASA score, large tumor size, preoperative chemotherapy, intraoperative adverse events, and low rectal anastomosis. While many studies have thoroughly analyzed the risk factors of anastomotic leakage, relatively few studies have focused on risk factors of anastomotic stricture. Rates have been shown to vary from 2%-30% in the literature, but these rates are usually under-reported due to the requirement for long-term follow-up[24]. In addition, while high-grade strictures are immediately recognized due to patient symptoms, low-grade strictures are not always identified[18]. According to recent studies, neoadjuvant CRT, clinical anastomotic leakage, and hand-sewn coloanal anastomosis have all been shown to be associated with independent risk factors of anastomotic stricture[25,26]. Endoscopic balloon dilation is the most common and effective way to treat symptomatic anastomotic stricture, but the recurrence rates after this procedure range from 6%-25% [27]. Some patients with recurrent anastomotic stricture have to accept PS to avoid the symptoms of anastomotic stricture and maintain a good quality of life.

Histology and pathology have played an important role in cancer diagnosis and prognostic prediction for decades. Some markers may potentially reflect the biological aggressiveness of the tumor, such as tumor type, tumor differentiation, growth pattern, tumor budding, and involvement of the serosa, nerves, lymphatic vessels, intramural, and extramural veins[28]. Patients with these high-risk

tumor patterns may easily develop local recurrence (LR), which can lead to PS. Perineural invasion and lymphovascular invasion have been demonstrated to be independent prognostic factors of recurrence in many cancers. Perineural invasion is characterized by tumor invasion of nervous structures and spread along nerve sheaths, while lymphovascular invasion is characterized by tumor invasion of small lymphatic or blood vessels^[29]. According to a study in rectal cancer by Peng *et al*^[30], the 5-year LR rate of the perineural invasion-positive group was more than 2.5-fold higher than that of the perineural invasion-negative group (22.7% vs 7.9%; P = 0.017). In addition, in terms of lymphovascular invasion, Dresen *et al*[28] indicated that the presence of lymphovascular invasion (OR 4.66, P < 0.001) was associated with an increased risk for the development of local recurrence in patients with rectal cancer. Another key factor for the development of local recurrence is positive CRM. Agger *et al*^[31] reported that the local recurrence rate was 17.0% in patients without any microscopic margin (CRM 0 mm) and 6.7% in patients with a CRM of 0-1 mm. With advancements in surgical techniques, the ratio of CRM has continued to decrease. In the study by Quentin et al[32], the rate of positive CRM decreased significantly after perineal dissection compared with after abdominal rectal dissection (4% vs 18%; P = 0.025). Moreover, it was beyond our expectations that tumor size was an independent risk factor for PS according to the results of the multivariate analysis. In previous studies, the results of the correlation between tumor size and the prognosis of rectal cancer are often contradictory, and multivariate analyses are seldom performed. However, in more recent studies, Kornprat et al[33] indicated that tumors larger than 4.5 cm are associated with high T and N classification, UICC stage, and tumor grade. Moreover, Chen *et al*[34] reported that pathological tumor size ≥ 5 cm is an independent prognostic factor for local recurrence in rectal adenocarcinoma. In our current study, the univariate analysis revealed that the independent risk factors for PS were lymphovascular invasion (OR, 1.99; 95% CI: 1.071–3.617; P = 0.026) and positive CRM (OR, 6.575; 95% CI: 2.955–14.604; P < 0.001), while the multivariate analysis revealed that the independent risk factors for PS were perineural invasion (OR, 3.085; 95% CI: 1.726–5.518; P < 0.001) and tumor size (OR, 1.076; 95%CI: 1.015-1.14; P = 0.014). The above four factors have been confirmed to be related to tumor recurrence, which can cause intestinal obstruction and affect intestinal continuity. The patient has no choice but to accept PS when the disease recurs because it is impossible for the physician to close the stoma in these patients.

Here, we developed a nomogram to predict the incidence of PS in patients with rectal cancer who undergo sphincter-saving surgery. To our knowledge, nomograms are widely used in many cancers to predict patient prognosis and cancer behavior (e.g., lymph node metastasis, recurrence, and distant metastasis)[35-37]. In addition, some studies have used nomograms to predict the rate of postoperative complications, such as infection, anastomotic leakage, and stenosis[38,39]. Currently, only a few predictive models of PS for patients with rectal cancer have been published [40-42]. We collected 391 cases for analysis, which is the largest case number to date among all relevant studies. The C-index for the nomogram is 0.903 (95%CI: 0.851-0.955), which indicates a perfect prediction model. According to the calibration curve, the nomogram calibration plot demonstrated high reliability. Patients with these risk factors would be classified as high-risk patients with PS, and they should be informed of their status prior to surgery. We propose that this nomogram provides more individualized outcome predictions and could aid clinicians and patients in the treatment decision making process.

The present study has some limitations. First, this was a retrospective study and was not randomized in nature. In some incomplete patient records, the details of stoma complications after hospital discharge may be difficult to evaluate. Second, the study period was relatively long, and differences may exist in surgeon discretion and surgical techniques. Finally, this analysis was based on data from a single center. External validation using data from other centers is needed to certify the discriminatory ability of this model. More representative prediction models can be developed using data from multiple centers.

CONCLUSION

This study reports that risk factors leading to PS were highly correlated with local recurrence, perirectal abscess, anastomosis site stenosis, perineural invasion, tumor size and operative time (min). Our established nomogram enables a relatively accurate assessment of the risk of PS after sphincter-saving surgery. The ease of use of this nomogram can improve a physician's ability to communicate the benefits and risks of various treatment options in SDM.

ARTICLE HIGHLIGHTS

Research background

Despite innovative advancements, the management of rectal cancer remains a formidable endeavor, especially distally located rectal cancer. According to previous studies, 3%-24% of rectal cancer patients experience anastomosis complications after sphincter-saving surgery, which may lead to permanent



stoma (PS).

Research motivation

Patients fail to achieve stoma closure can cause drastic changes in lifestyle and physical perceptions.

Research objectives

The purpose of this study was to determine the risk factors for PS and to develop a prediction model to predict the probability of PS in rectal cancer patients after sphincter-saving surgery.

Research methods

A retrospective cohort of 421 rectal cancer patients who underwent radical surgery at Taipei Medical University Hospital between January 2012 and December 2020 was included in this study. Univariate and multivariate analyses were performed to identify the independent risk factors for PS. A nomogram was developed according to the independent risk factors obtained in the multivariate analysis. The performance of the nomogram was assessed using a receiver operating characteristic curve and a calibration curve.

Research results

The PS stoma rate after sphincter-saving surgery was 15.1% (59/391) in our study after a median followup of 47.3 mo (range 7-114 mo). Multivariate logistic regression analysis demonstrated that local recurrence, perirectal abscess, anastomosis site stenosis, perineural invasion, tumor size, liver disease, and operative time were independent risk factors for PS. After exclude liver disease, these identified risk factors were incorporated into the nomogram, and the concordance index of this model was 0.903 (95%CI: 0.851-0.955). According to the calibration curves, the nomogram represents a perfect prediction model

Research conclusions

This study reports that risk factors leading to PS were highly correlated with local recurrence, perirectal abscess, anastomosis site stenosis, perineural invasion, tumor size and operative time (min). Our established nomogram enables a relatively accurate assessment of the risk of PS after sphincter-saving surgery. The ease of use of this nomogram can improve a physician's ability to communicate the benefits and risks of various treatment options in shared decision making.

Research perspectives

The present study has some limitations. First, this was a retrospective study and was not randomized in nature. In some incomplete patient records, the details of stoma complications after hospital discharge may be difficult to evaluate. Second, the study period was relatively long, and differences may exist in surgeon discretion and surgical techniques. Finally, this analysis was based on data from a single center. External validation using data from other centers is needed to certify the discriminatory ability of this model. More representative prediction models can be developed using data from multiple centers.

FOOTNOTES

Author contributions: Kuo CY manuscript writing; Wei PL and Chen CC participate in patients' treatment and collection; Lin YK data interpretation and statistical analysis; Kuo LJ original idea creator, project coordinator, manuscript reviewing and proofing; Lin YK and Kuo LJ contributed equally to this work; All authors have read and approve the finial manuscript.

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