

World Journal of *Gastroenterology*

World J Gastroenterol 2023 February 28; 29(8): 1243-1394



REVIEW

- 1243 Predictors of early and late hepatocellular carcinoma recurrence
Nevola R, Ruocco R, Criscuolo L, Villani A, Alfano M, Beccia D, Imbriani S, Claar E, Cozzolino D, Sasso FC, Marrone A, Adinolfi LE, Rinaldi L
- 1261 Functional constipation in children: What physicians should know
Tran DL, Sintusek P
- 1289 Early-onset colorectal cancer: A review of current knowledge
Saraiva MR, Rosa I, Claro I

MINIREVIEWS

- 1304 Hamartomatous polyps: Diagnosis, surveillance, and management
Gorji L, Albrecht P

ORIGINAL ARTICLE**Basic Study**

- 1315 Stress granules inhibit endoplasmic reticulum stress-mediated apoptosis during hypoxia-induced injury in acute liver failure
Li WY, Yang F, Li X, Wang LW, Wang Y

Observational Study

- 1330 Intestinal complications in Brazilian patients with ulcerative colitis treated with conventional therapy between 2011 and 2020
Martins AL, Galhardi Gasparini R, Sasaki LY, Saad-Hossne R, Ritter AMV, Barreto TB, Marcolino T, Yang Santos C
- 1344 Establishment of a prediction model for severe acute radiation enteritis associated with cervical cancer radiotherapy
Ma CY, Zhao J, Gan GH, He XL, Xu XT, Qin SB, Wang LL, Li L, Zhou JY
- 1359 Better performance of PIVKA-II for detecting hepatocellular carcinoma in patients with chronic liver disease with normal total bilirubin
Qian XJ, Wen ZM, Huang XM, Feng HJ, Lin SS, Liu YN, Li SC, Zhang Y, Peng WG, Yang JR, Zheng ZY, Zhang L, Zhang DW, Lu FM, Liu LJ, Pan WD

SYSTEMATIC REVIEWS

- 1374 Burden of bone disease in chronic pancreatitis: A systematic review and meta-analysis
Chhoda A, Hernandez-Woodbine MJ, Addo NAA, Nasir SA, Grimshaw A, Gunderson C, Ahmed A, Freedman SD, Sheth SG

ABOUT COVER

Editorial Board Member of *World Journal of Gastroenterology*, Ragesh Babu Thandassery, MD, Assistant Professor, Doctor, Department of Medicine, Central Arkansas Veterans Healthcare System, Little Rock, AR 72205, United States. doc.ragesh@gmail.com

AIMS AND SCOPE

The primary aim of *World Journal of Gastroenterology* (WJG, *World J Gastroenterol*) is to provide scholars and readers from various fields of gastroenterology and hepatology with a platform to publish high-quality basic and clinical research articles and communicate their research findings online. WJG mainly publishes articles reporting research results and findings obtained in the field of gastroenterology and hepatology and covering a wide range of topics including gastroenterology, hepatology, gastrointestinal endoscopy, gastrointestinal surgery, gastrointestinal oncology, and pediatric gastroenterology.

INDEXING/ABSTRACTING

The WJG is now abstracted and indexed in Science Citation Index Expanded (SCIE, also known as SciSearch®), Current Contents/Clinical Medicine, Journal Citation Reports, Index Medicus, MEDLINE, PubMed, PubMed Central, Scopus, 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 WJG as 5.374; IF without journal self cites: 5.187; 5-year IF: 5.715; Journal Citation Indicator: 0.84; Ranking: 31 among 93 journals in gastroenterology and hepatology; and Quartile category: Q2. The WJG's CiteScore for 2021 is 8.1 and Scopus CiteScore rank 2021: Gastroenterology is 18/149.

RESPONSIBLE EDITORS FOR THIS ISSUE

Production Editor: *Yu-Xi Chen*; Production Department Director: *Xu Guo*; Editorial Office Director: *Jia-Ru Fan*.

NAME OF JOURNAL

World Journal of Gastroenterology

ISSN

ISSN 1007-9327 (print) ISSN 2219-2840 (online)

LAUNCH DATE

October 1, 1995

FREQUENCY

Weekly

EDITORS-IN-CHIEF

Andrzej S Tarnawski

EDITORIAL BOARD MEMBERS

<http://www.wjgnet.com/1007-9327/editorialboard.htm>

PUBLICATION DATE

February 28, 2023

COPYRIGHT

© 2023 Baishideng Publishing Group Inc

INSTRUCTIONS TO AUTHORS

<https://www.wjgnet.com/bpg/gerinfo/204>

GUIDELINES FOR ETHICS DOCUMENTS

<https://www.wjgnet.com/bpg/GerInfo/287>

GUIDELINES FOR NON-NATIVE SPEAKERS OF ENGLISH

<https://www.wjgnet.com/bpg/gerinfo/240>

PUBLICATION ETHICS

<https://www.wjgnet.com/bpg/GerInfo/288>

PUBLICATION MISCONDUCT

<https://www.wjgnet.com/bpg/gerinfo/208>

ARTICLE PROCESSING CHARGE

<https://www.wjgnet.com/bpg/gerinfo/242>

STEPS FOR SUBMITTING MANUSCRIPTS

<https://www.wjgnet.com/bpg/GerInfo/239>

ONLINE SUBMISSION

<https://www.f6publishing.com>

Observational Study

Establishment of a prediction model for severe acute radiation enteritis associated with cervical cancer radiotherapy

Chen-Ying Ma, Jing Zhao, Guang-Hui Gan, Xiao-Lan He, Xiao-Ting Xu, Song-Bing Qin, Li-Li Wang, Li Li, Ju-Ying Zhou

Specialty type: Health care sciences and services

Provenance and peer review:

Unsolicited article; Externally peer reviewed.

Peer-review model: Single blind

Peer-review report's scientific quality classification

Grade A (Excellent): A

Grade B (Very good): B

Grade C (Good): 0

Grade D (Fair): 0

Grade E (Poor): 0

P-Reviewer: Aydin S, Turkey; Šarenac TM, Serbia

Received: October 17, 2022

Peer-review started: October 17, 2022

First decision: January 3, 2023

Revised: January 13, 2023

Accepted: February 14, 2023

Article in press: February 14, 2023

Published online: February 28, 2023



Chen-Ying Ma, Jing Zhao, Guang-Hui Gan, Xiao-Lan He, Xiao-Ting Xu, Song-Bing Qin, Li-Li Wang, Li Li, Ju-Ying Zhou, Radiation Oncology, The First Affiliated Hospital of Soochow University, Suzhou 215123, Jiangsu Province, China

Ju-Ying Zhou, State Key Laboratory of Radiation Medicine and Protection, Soochow University, Suzhou 215123, Jiangsu Province, China

Corresponding author: Ju-Ying Zhou, MD, Radiation Oncology, The First Affiliated Hospital of Soochow University, No. 188 Shizi Street, Suzhou 215123, Jiangsu Province, China. zhoujuyingsy@163.com

Abstract**BACKGROUND**

Cervical cancer is one of the most common gynecological malignant tumors. Radiation enteritis (RE) leads to radiotherapy intolerance or termination of radiotherapy, which negatively impacts the therapeutic effect and seriously affects the quality of life of patients. If the incidence of RE in patients can be predicted in advance, and targeted clinical preventive treatment can be carried out, the side effects of radiotherapy in cervical cancer patients can be significantly reduced. Furthermore, accurate prediction of RE is essential for the selection of individualized radiation dose and the optimization of the radiotherapy plan.

AIM

To analyze the relationships between severe acute RE (SARE) of cervical cancer radiotherapy and clinical factors and dose-volume parameters retrospectively.

METHODS

We included 50 cervical cancer patients who received volumetric modulated arc therapy (VMAT) from September 2017 to June 2018 in the Department of Radiotherapy at The First Affiliated Hospital Soochow University. Clinical and dose-volume histogram factors of patients were collected. Logistic regression analysis was used to evaluate the predictive value of each factor for SARE. A nomogram to predict SARE was developed (SARE scoring system ≥ 3 points) based on the multiple regression coefficients; validity was verified by an internal verification method.

RESULTS

Gastrointestinal and hematological toxicity of cervical cancer VMAT gradually increased with radiotherapy and reached the peak at the end of radiotherapy. The main adverse reactions were diarrhea, abdominal pain, colitis, anal swelling, and blood in the stool. There was no significant difference in the incidence of gastrointestinal toxicity between the radical and postoperative adjuvant radiotherapy groups ($P > 0.05$). There were significant differences in the small intestine V_{20} , V_{30} , V_{40} , and rectal V_{40} between adjuvant radiotherapy and radical radiotherapy after surgery ($P < 0.05$). Univariate and multivariate analyses revealed anal bulge rating (OR: 14.779, 95%CI: 1.281-170.547, $P = 0.031$) and disease activity index (DAI) score (OR: 53.928, 95%CI: 3.822-760.948, $P = 0.003$) as independent predictors of SARE.

CONCLUSION

Anal bulge rating (> 0.500 grade) and DAI score (> 2.165 points) can predict SARE. The nomogram shows potential value in clinical practice.

Key Words: Cervical cancer; Intensity-modulated radiotherapy; Radiation enteritis; Nomogram; Predictor

©The Author(s) 2023. Published by Baishideng Publishing Group Inc. All rights reserved.

Core Tip: Radiation enteritis (RE) not only seriously affects the quality of life of patients, but it also leads to radiotherapy intolerance or termination of radiotherapy. The aim of our study was to determine the cumulative incidence of acute RE associated with cervical cancer radiotherapy in patients with RE in organs at risk and changes in dose-volume histogram indices. The nomogram of severe acute RE (SARE) was further developed according to the clinical factors, cumulative incidence of SARE and dosimetric parameters of volumetric modulated arc therapy patients, which may be useful for individualized risk assessment and accurate prediction of SARE to guide clinical treatment strategies.

Citation: Ma CY, Zhao J, Gan GH, He XL, Xu XT, Qin SB, Wang LL, Li L, Zhou JY. Establishment of a prediction model for severe acute radiation enteritis associated with cervical cancer radiotherapy. *World J Gastroenterol* 2023; 29(8): 1344-1358

URL: <https://www.wjgnet.com/1007-9327/full/v29/i8/1344.htm>

DOI: <https://dx.doi.org/10.3748/wjg.v29.i8.1344>

INTRODUCTION

Cervical cancer is one of the most common gynecological malignant tumors, ranking fourth among malignant tumors in females worldwide[1]. Radiotherapy plays an important role in the postoperative adjuvant treatment of cervical cancer patients and the radical treatment of patients who are ineligible for surgery. Radiation enteritis (RE) is a common and potentially dose-limiting toxic reaction of pelvic radiotherapy. The clinical manifestations include nausea, vomiting, abdominal pain, bloody stool, mucous stool, diarrhea, tenesmus, and urinary incontinence. RE not only seriously affects the quality of life of patients, but it also leads to radiotherapy intolerance or termination of radiotherapy, which negatively impacts the therapeutic effect. If the incidence of RE in patients can be predicted in advance, and targeted clinical preventive treatment can be carried out, the side effects of radiotherapy in cervical cancer patients can be significantly reduced. Furthermore, accurate prediction of RE is essential for the selection of individualized radiation dose and the optimization of the radiotherapy plan.

Many studies have shown that the occurrence and development of RE are related to intestinal radiotherapy technology, total dose, total volume, fraction dose, ratio of dose to volume, and uniformity of radiation dose distribution[2-6]. RE is also influenced by factors such as inflammatory bowel disease [7], collagen vascular disease[8], history of abdominal and pelvic surgery[9], history of pelvic inflammatory disease, diabetes mellitus[10], and acquired immunodeficiency syndrome[11]. However, there is no consensus on the relative importance of these predictors. Only a few studies have explored the dosimetric risk factors for RE in cervical cancer patients[12]. How to predict RE and reduce adverse reactions is the core issue of clinical radiotherapy for cervical cancer. Notably, a single predictor has limited predictive power and cases can show substantial differences in heterogeneity of RE; these issues should be addressed in further research[13]. To specifically and accurately identify RE, there is an urgent need for an accurate predictive model that combines multiple factors.

The initial objective of this study was to determine the cumulative incidence of acute RE associated with intensity modulated radiation therapy (IMRT) for cervical cancer in patients with RE in organs at risk (OAR) and changes in dose-volume histogram indices. The severe acute RE scoring system (SARE-

SS) was defined by integrating clinical factors and dosimetric physical parameters. Finally, the nomogram of severe acute RE (SARE) was further developed according to the clinical factors, cumulative incidence of SARE and dosimetric parameters of volumetric modulated arc therapy (VMAT) patients. This nomogram may be useful for individualized risk assessment and accurate prediction of SARE to guide clinical treatment strategies.

MATERIALS AND METHODS

Patient selection

Fifty patients with cervical cancer who received pelvic local radiotherapy in the Department of Radiation Oncology of The First Affiliated Hospital of Soochow University from August 2017 to August 2018 were selected. The inclusion criteria were as follows: (1) 18-80 years old; (2) cervical cancer confirmed by pathology; (3) no history of intestinal diseases or metabolic diseases; and (4) good understanding and communication skills. The exclusion criteria were as follows: (1) Severe heart, liver and kidney dysfunction; (2) termination of treatment because of serious complications during and after radiotherapy (such as cardiopulmonary, hepatic and renal insufficiency, severe infection, severe bone marrow suppression, massive hemorrhage, vaginorectal fistula and vaginovesical fistula); and (3) refusal to participate in the study.

This study was approved by the ethics committee of The First Affiliated Hospital of Soochow University Medical Ethics Committee [approval No. 2016(100)]. All enrolled patients provided signed informed consent.

External irradiation method for cervical cancer

Radiotherapy simulation positioning: On the night before radiotherapy, all patients took laxatives to clean the intestinal track. One hour before positioning, the bladder was emptied and patients drank 800 mL of water to fill the bladder. The patients then took the supine position. A Philips [Brilliance computed tomography (CT) Big Bore] large aperture CT machine was used for simulated CT localization, and enhanced CT scanning was performed with a slice thickness of 5 mm. The scanning range was from the upper edge of T11 vertebral body to 5 cm below the ischial tuberosity. The CT localization images were transferred to the treatment planning system (V5.1.1, Elekta, Monaco, Sweden).

Radiotherapy target delineation: Magnetic resonance imaging (MRI), CT or positron emission tomography-CT (PET-CT) examinations were routinely performed during the pre-radiotherapy evaluation. The target volume and OAR were delineated according to the standards of Radiation Therapy Oncology Group and Japan Clinical Oncology Group.

Radiotherapy plan design: The Monaco treatment planning system was used to design the 7-field reverse dynamic VMAT plan using unified parameters. The X-ray beam energy was 6 MV. At least 95% of the pelvic lymphatic drainage (PTV) is required to reach the prescribed dose, and no hot spot of $\geq 110\%$ dose can occur outside the PTV. The OAR dose limit was uniform (OAR dose limitation: Bladder $V_{40} < 0\%$, rectum $V_{40} < 40\%$, colon $V_{40} < 30\%$, small intestine $V_{40} < 20\%$, femoral head $V_{45} < 5\%$).

Prescription dose of the VMAT target volume after operation: We administered 45 Gy/25 fractions for moderate risk included lymph node metastasis, paracytal invasion, and positive margin) PTV1 and 50 Gy/25 fractions for high risk PTV1 50. 4 Gy/28 fractions, the prescription dose of the remaining target volume can be executed according to the radical external irradiation scheme.

Prescription dose of the radical VMAT target volume: We administered the following doses: PTV1 45 Gy/25 fractions, parametrial area (PTV2) 50 Gy/25 fractions, and positive lymph nodes (PGTVnd) 56 Gy/28 fractions. In cases with para-aortic area, the dose for the para-aortic lymphatic drainage area (PTV3) was 36-40 Gy/20 fractions.

Implementation of the VMAT plan: VMAT was performed using the Elekta Synergy linear accelerator. All patients were treated five times a week, with one treatment a day. Cone-beam CT (Elekta) image-guided VMAT was performed at least once a week, and the error of each treatment was kept within 3 mm.

Chemotherapy for cervical cancer

Of the 50 cervical cancer patients included in this study, 18 were treated with concurrent chemoradiotherapy (CCRT) and 32 received sequential chemoradiotherapy (SCRT). All doses and adjustments of the chemotherapy regimen followed the NCCN guidelines[14].

End point definitions

General data collection: Data on age, weight, height, body mass index (BMI)[15], age-adjusted Charlson Comorbidity Index (aCCI), pathological data, HPV, squamous cell carcinoma antigen, risk rating, FIGO

2018 staging information, surgery, radiotherapy and chemotherapy were collected.

Data collection of acute RE: The adverse reactions of all patients were monitored at baseline and 2 wk, 4 wk, and 6 wk after the start of radiotherapy. Adverse reactions included diarrhea, abdominal pain, colitis, anal distension, hematochezia and bone marrow suppression. The diagnosis and classification of RE were determined following the National Institutes of Health Common Adverse Events Evaluation Criteria (CTCAE 5.0).

We established the SARE-SS to reflect the adverse reaction score as follows: SARE-SS = the sum of CTCAE scores of diarrhea + abdominal pain + colitis + anal bulging + hematochezia. A score of ≥ 3 indicated the presence of SARE. The diagnosis of SARE was determined by at least two experienced radiation oncologists on the basis of clinical symptoms and changes in the results of ancillary tests.

Disease activity index determination: The disease activity index (DAI) is based on scores that reflect the weight loss of patients (weight unchanged is 0, 1%-5% is 1 point, 5%-10% is 2 points, 10%-15% is 3 points, more than 15% is 4 points), stool viscosity (normal is 0 points, loose stool is 2 points, diarrhea is 4 points) and stool bleeding (normal is 0 points, occult blood positive is 2 points, dominant bleeding 4 points). The sum of the three scores is divided by 3 to obtain the final DAI score.

Radiotherapy planning parameter collection: Statistics of different OAR dose-volume relationship (percentage of PTV volume receiving prescription dose), recorded as V_x .

Statistical analysis

Statistical analysis was performed using SPSS 20.0. The quantitative data of normal distribution were expressed as (mean \pm SD), and *t* test was used for comparisons between groups. The data of skew distribution were tested by Mann-Whitney *U* test. Qualitative data were expressed as number of cases and percentage; Fisher's exact probability method or χ^2 test was used to compare unordered categorical data, and Mantel-Haenszel χ^2 and Mann-Whitney *U* test were used to compare ordered categorical data. $P < 0.05$ was considered to indicate statistical significance. Receiver operating characteristic curve (ROC curve) was used to analyze the specificity and sensitivity of OAR dose-volume parameters in predicting RE. Two-factor repeated-measures ANOVA was used to analyze the relationship between DAI scores and time in different groups, and one-way ANOVA was used to compare the differences of DAI scores among four time points in each group.

To establish the prediction model, univariate logistic regression model was used to evaluate the predictive ability of each factor for SARE. Multivariate analysis was performed for significant factors from the univariate analysis. Kendall correlation analysis was used to avoid multicollinearity between factors. Factors with significant predictive value in the multivariate analysis were used to construct a nomogram. The area under the ROC curve (AUC), calibration curve and decision curve analysis (DCA) were used for nomogram validation. Data analysis was performed using R software (software version 4.0.2, R package version rmda 1.6).

RESULTS

Clinical characteristics of patients

This study included 50 patients with cervical cancer. Among the 50 patients, 14 patients received radical radiotherapy, 35 patients received postoperative adjuvant pelvic radiotherapy and 1 patient received pelvic radiotherapy combined with after loading vaginal radiotherapy because the vaginal margin was 2 cm away from the tumor boundary. The specific clinical factors are detailed in Table 1. The mean age of the postoperative adjuvant radiotherapy group was significantly lower than that of radical radiotherapy group ($P < 0.05$) (Table 2). The BMI of the postoperative adjuvant radiotherapy group was also significantly lower ($P < 0.05$), but there was no significant difference in weight, pathological type, cumulative incidence of RE and SARE and irradiation of para-aortic extension field between the two groups ($P > 0.05$).

Dose-volume comparison of OAR in the VMAT plan between the postoperative adjuvant radiotherapy group and radical radiotherapy group

Patients in the radical radiotherapy group were treated with total pelvic and parauterine local radiotherapy, and patients in the postoperative adjuvant radiotherapy group were treated with total pelvic prophylactic radiotherapy. The dose and volume of small intestine in the radical radiotherapy group were significantly reduced compared with that in the postoperative adjuvant radiotherapy group, and the decrease of V_{20} , V_{30} and V_{40} were statistically significant ($P < 0.05$) (Table 3). Rectal V_{30} , V_{35} and V_{40} in the radical radiotherapy group were significantly higher than those in the postoperative adjuvant radiotherapy group. The increase of V_{40} was statistically significant ($P < 0.05$). There was no significant difference in the dose-volume relationship of colon, bladder and femoral head between the two groups ($P > 0.05$).

Table 1 Baseline characteristics of all patients (*n* = 50)

Characteristic	<i>n</i> (%)
Age, median (IQR)	51.0 (45.5-62.0)
Weight, IQR	55.0 (50.5-61.8)
BMI/kg/m ² , IQR	21.5 (20.7-22.9)
aCCI	
2-3/4-5	38 (76)/12 (24)
Pathological diagnosis	
Squamous cell carcinoma	47 (94)
Adenocarcinoma	1 (2)
Adenosquamous carcinoma	2 (4)
T	
> 4 cm/≤ 4 cm	24 (48)/26 (52)
N	
Negative/positive	15 (30)/35 (70)
Metastatic pelvic lymph nodes	
Positive/negative	14 (28)/36 (72)
Metastatic common iliac lymph nodes	
Positive/negative	5 (10)/45 (90)
Para-aortic lymph nodes	
Positive/negative	2 (4)/48 (96)
FIGO staging	
I/II/III/IVA	19 (38)/21 (42)/5 (10)/5 (10)
LVSI	
Positive/negative	16 (32)/34 (68)
Degree of differentiation	
Low and medium differentiation/high differentiation	40 (80)/10 (20)
Degree of infiltration	
Shallow 1/3/Medium 1/3/Deep 1/3	10 (20)/16 (32)/24 (48)
Incised margin	
R1 + R2/R0	5 (10)/45 (90)
Danger degree	
Low/medium/high risk	15 (30)/12 (24)/23 (46)
HPV	
Positive/negative	46 (92)/4 (8)
SCC	
Abnormally elevated/no abnormality observed	17 (34)/33 (66)
RE	
Occurred/not occurred	42 (84)/8 (16)
SARE/Non-SARE	15 (30)/35 (70)
Operation	
Radical surgery/no surgery	36 (72)/14 (28)
Chemotherapy	

CCRT/SCRT	18 (36)/32 (64)
Pelvic External Radiation Dose, IQR	45.0 (45.0-48.0)
Para-aortic extension field	
Radiotherapy/no radiotherapy	4 (8)/46 (92)

aCCI: Adjusted Charlson Comorbidity Index; SARE: Severe acute radiation enteritis; CCRT: Concurrent chemoradiotherapy; SCRT: Sequential chemoradiotherapy; SCC: Squamous cell carcinoma; HPV: Human papillomavirus.

Table 2 Comparison of general data between postoperative adjuvant radiotherapy and radical radiotherapy for patients with cervical cancer, *n* (%)

Factor	Postoperative adjuvant radiotherapy group	Radical radiotherapy group	T/ χ^2 values	P value
Age/yr	48.25 ± 8.05	61.86 ± 13.17	-4.450	< 0.001
Young and middle-aged	33 (92)	5 (36)	17.301	< 0.001
Old age	3 (8)	9 (64)		
Weight	57.31 ± 8.00	53.07 ± 8.74	1.638	0.108
BMI/kg/m ²	21.86 ± 2.07	20.21 ± 2.79	2.284	0.027
Tumor type				0.186 ¹
Squamous cell carcinoma	35 (97)	12 (86)		
Adenocarcinoma	0 (0)	1 (7)		
Adenosquamous carcinoma	1 (3)	1 (7)		
CCRT			0.397	0.529
Implement	12 (33)	6 (43)		
Not implemented	24 (67)	8 (57)		
Para-aortic extended field radiotherapy				0.186 ¹
Implement	1 (3)	2 (14)		
Not implemented	35 (97)	12 (86)		
RE			-	1.000 ¹
Occurred	30 (83)	12 (86)		
Did not occur	6 (17)	2 (14)		
SARE			2.286	0.131
Occurred	13 (36)	2 (14)		
Did not occur	23 (64)	12 (86)		

¹Fisher exact probability method.

Data are shown as *n* (%) or mean ± SD. SARE: Severe acute radiation enteritis; CCRT: Concurrent chemoradiotherapy.

Comparison of the cumulative incidence of VMAT-related acute RE in cervical cancer between the postoperative adjuvant radiotherapy group and the radical radiotherapy group

The main radiotherapy-related adverse reactions included diarrhea, abdominal pain, colitis, anal distension, hematochezia and bone marrow suppression. All adverse reactions were grade 1 to 3; no adverse reactions above grade 3 occurred. RE was monitored as follows (Table 4). There was no significant difference in the incidences of adverse reaction between the radical radiotherapy group and postoperative adjuvant radiotherapy group ($P > 0.05$).

Evaluation of the specificity and sensitivity of OAR dose-volume parameters in the IMRT plan for predicting acute/SARE

ROC curve was used to analyze the specificity and sensitivity of small intestine V_{20r} , V_{30r} , V_{40} and rectum

Table 3 Organs at risk dose-volume comparison between adjuvant radiotherapy (*n* = 36) and radical radiotherapy (*n* = 14) after cervical cancer surgery (%)

OAR	Group	V ₂₀	V ₂₅	V ₃₀	V ₃₅	V ₄₀
Small intestine	Postoperative adjuvant radiotherapy group	77.41 ± 16.49	51.90 ± 10.99	39.51 ± 8.24	26.40 ± 5.50	19.25 ± 4.12
	Radical radiotherapy group	67.75 ± 14.10	50.11 ± 9.86	32.92 ± 7.91	25.18 ± 7.12	15.07 ± 4.65
	Z value	-2.377	-0.864	-2.701	-0.367	-3.003
	P value	0.017	0.387	0.007	0.713	0.003
Colon	Postoperative adjuvant radiotherapy group	100.00 ± 0.00	99.88 ± 0.73	98.26 ± 6.09	91.36 ± 11.67	75.64 ± 12.59
	Radical radiotherapy group	95.57 ± 16.59	95.34 ± 17.44	94.21 ± 21.66	91.09 ± 21.12	84.17 ± 7.94
	Z value	-1.604	-0.732	-0.332	-0.750	-2.247
	P value	0.109	0.464	0.740	0.453	0.025
Rectum	Postoperative adjuvant radiotherapy group	100.00 ± 0.00	100.00 ± 0.00	99.17 ± 2.90	95.63 ± 9.41	75.64 ± 12.59
	Radical radiotherapy group	100.00 ± 0.00	100.00 ± 0.00	100.00 ± 0.00	99.60 ± 1.48	84.17 ± 7.94
	Z value	0.000	0.000	-1.102	-1.745	-2.247
	P value	1.000	1.000	0.270	0.081	0.025
Bladder	Postoperative adjuvant radiotherapy group	70.87 ± 28.16	68.75 ± 24.41	47.18 ± 23.63	35.58 ± 16.04	27.33 ± 12.22
	Radical radiotherapy group	74.08 ± 27.93	71.94 ± 25.87	52.54 ± 28.93	40.40 ± 21.04	31.18 ± 16.28
	T-value	-0.354	-0.523	-0.519	-0.605	-0.627
	P value	0.723	0.601	0.604	0.545	0.531
Femoral head	Postoperative adjuvant radiotherapy group	48.55 ± 28.80	34.45 ± 22.27	24.73 ± 16.00	18.12 ± 11.68	12.23 ± 8.10
	Radical radiotherapy group	39.60 ± 19.76	27.13 ± 14.03	19.54 ± 10.07	14.42 ± 7.69	9.47 ± 5.37
	T-value	-0.951	-0.929	-0.994	-0.929	-0.983
	P value	0.342	0.353	0.320	0.353	0.326

OAR: Organs at risk.

V₄₀ in predicting acute/SARE. The AUC was less than 0.60 and *P* > 0.05. These results suggest that a single OAR dose-volume parameter is not enough to predict the occurrence of acute/SARE.

DAI score changes

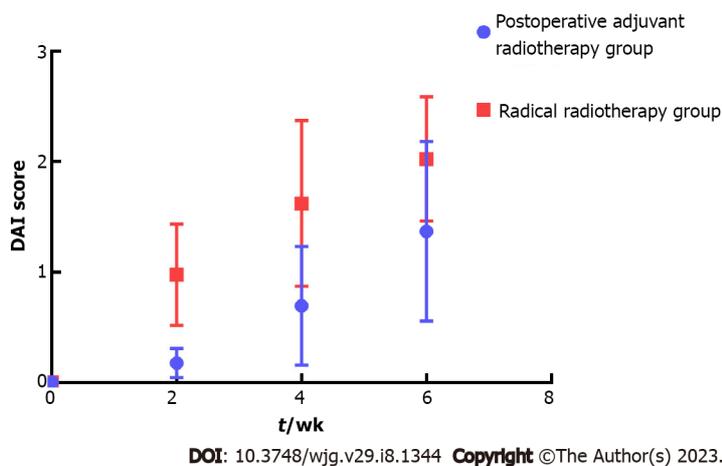
The DAI scores of the radical radiotherapy group were higher than those of the postoperative adjuvant radiotherapy group at the second week, fourth week and the end of radiotherapy (Figure 1). There was an interaction effect between different groups and time ($\chi^2 = 77.238, P < 0.01$). The DAI scores of the two groups showed an increasing trend over time (*P* < 0.05), and the DAI scores of the radical radiotherapy group increased the most.

Establishment of the SARE prediction model

Univariate analysis showed that abdominal pain, colitis, anal bulging, hematochezia, DAI score, age, and CCRT were significantly correlated with the occurrence of severe acute RE (all *P* < 0.05) (Table 5). Kendall analysis showed a relatively strong correlation between abdominal pain, hematochezia and DAI score (*R* = 0.715, 0.622, *P* < 0.001); the correlation between other factors was very weak (Figure 2A). To avoid multicollinearity, abdominal pain, hematochezia, and DAI scores were included in the multivariate analysis. Multivariate logistic regression analysis was conducted to analyze the factors of colitis, anal bulge, DAI score and age. In multivariate analysis, anal bulge rating (OR: 14.779, 95%CI: 1.281-170.547, *P* = 0.031) and DAI score (OR: 53.928, 95%CI: 3.822-760.948, *P* = 0.003) were independent predictors of SARE (Table 6). These factors were used to construct the nomogram. The constant in the logistic regression equation was -10.039, and the logistic regression equation was Logit (*P*) = 3.988 × DAI + 2.693 × anal bulge rating -10.039.

Table 4 Comparison of the incidence of acute radiation enteritis in patients with cervical cancer treated with adjuvant radiotherapy ($n = 36$) and radical radiotherapy ($n = 14$), n (%)

Adverse reaction	Group	Level 0	Level 1	Level 2	Level 3	χ^2 value	P value
Diarrhea	Postoperative adjuvant radiotherapy group	10 (28)	19 (53)	5 (14)	2 (5)	2.748	0.432
	Radical radiotherapy group	3 (21)	10 (72)	0 (0)	1 (7)		
Abdominal pain	Postoperative adjuvant radiotherapy group	21 (58)	12 (33)	3 (9)	0 (0)	1.423	0.491
	Radical radiotherapy group	8 (57)	6 (43)	0 (0)	0 (0)		
Colitis	Postoperative adjuvant radiotherapy group	25 (69)	11 (31)	0 (0)	0 (0)	-	0.140 ¹
	Radical radiotherapy group	13 (93)	1 (7)	0 (0)	0 (0)		
Anal bulging	Postoperative adjuvant radiotherapy group	22 (61)	9 (25)	5 (14)	0 (0)	1.374	0.503
	Radical radiotherapy group	11 (79)	2 (14)	1 (7)	0 (0)		
Blood in the stool	Postoperative adjuvant radiotherapy group	23 (64)	13 (36)	0 (0)	0 (0)		0.179 ¹
	Radical radiotherapy group	12 (86)	2 (14)	0 (0)	0 (0)		
Myelosuppression	Postoperative adjuvant radiotherapy group	12 (33)	9 (25)	10 (28)	5 (14)	3.691	0.297
	Radical radiotherapy group	3 (21)	4 (29)	2 (14)	5 (36)		

¹Fisher exact probability method.**Figure 1 Disease activity index score during volumetric modulated arc therapy in 50 patients with cervical cancer.**

Development and validation of the nomogram

Based on the multivariate logistic regression coefficients, the prediction model was visually represented as a nomogram (Figure 2B). The ROC curves for the anal bulge rating, DAI score and nomogram are shown in Figure 2C. ROC analysis showed that the AUC of the prediction model was 0.950 (95% CI: 0.891-1.000), which was much higher than that of each parameter alone (anal bulge rating: 0.805, 95% CI: 0.651-0.959; DAI score: 0.892, 95% CI: 0.873-1.000). The MAL thresholds for anal bulge rating and DAI score were 0.5 and 2.165 points. The nomogram had a high predictive efficiency (sensitivity: 80.0%, specificity: 91.4%). In addition, the calibration curve showed good agreement between the predicted severe acute RE and the actual observations (Figure 2D). In most threshold probabilities, DCA showed a positive net benefit with a satisfactory nomogram, indicating a good potential clinical effect of the model (Figure 2E).

Table 5 Comparison of general data between patients with cervical cancer with severe acute radiation enteritis or no severe acute radiation enteritis

Factor	SARE (n = X)	No SARE (n = X)	OR (95%CI)	P value
	Median (IQR)	Median (IQR)		
Clinical factors				
Elderly (> 60 yr)/young and middle-aged (20-60 yr)			0.123 (0.015-1.040)	0.054
Age	47.5 (37.0-54.0)	53.0 (46.0-63.0)	0.930 (0.872-0.992)	0.027
Weight	53.0 (51.0-60.9)	56.0 (50.0-61.8)	0.983 (0.918-1.054)	0.631
BMI/kg/m ²	20.5 (19.7-22.2)	21.9 (20.4-23.0)	0.936 (0.746-1.174)	0.566
ACCI (2-3/4-5)			0.123 (0.015-1.040)	0.054
Tumor type (squamous/non-squamous)			0.848 (0.139-5.179)	0.859
T (> 4 cm/≤ 4 cm)			0.510 (0.153-1.695)	0.272
N (positive/negative)			0.615 (0.165-2.298)	0.538
Metastatic pelvic lymph nodes (positive/negative)			0.694 (0.185-2.611)	0.589
Common iliac lymph node metastasis (positive/negative)			0.427 (0.046-3.980)	0.455
Para-aortic lymph nodes (positive/negative)			1.471 (1.216-1.779)	0.999
FIGO stage (IIB-IVA/IB-IIA)			1.108 (0.328-3.739)	0.869
LVSI (positive/negative)			0.548 (0.147-2.034)	0.368
Differentiation degree (low and medium differentiation/high differentiation)			1.011 (0.225-4.536)	0.988
Infiltration degree (shallow 1/3/medium depth 1/3)			0.837 (0.191-3.673)	0.813
Incisal margin (R1 + R2/R0)			0.427 (0.046-3.980)	0.455
Risk (low risk/medium high risk)			1.227 (0.340-4.242)	0.754
HPV (positive/negative)			1.324 (0.127-13.785)	0.815
SCC (abnormally elevated/not abnormal)			0.303 (0.074-1.245)	0.098
Surgery (performed/not performed)			2.080 (0.497-8.706)	0.316
CCRT (implemented/not implemented)			6.042 (1.681-21.718)	0.006
Myelosuppression (Grade 1-3/Grade 0)			0.617 (0.178-2.144)	0.446
Diarrhea (Grade 1-3/Grade 0)			NA	0.999
Abdominal pain (Grade 1-3/Grade 0)			25.375 (4.750-135.559)	< 0.001
Colitis (Grade 1-3/Grade 0)			38.500 (6.530-226.993)	< 0.001
Anal bulge (Grade 1-3/Grade 0)			5.185 (1.470-18.286)	0.010
Hematochezia (Grade 1-3/Grade 0)			NA	< 0.001
DAI score	2.8 (2.3-3.0)	1.7 (1.2-2.0)	152.546 (6.045-3849.771)	0.002
Physical dose parameters				
Small intestine				
V ₂₀ (%)	80.9 (68.6-93.7)	78.0 (66.8-82.5)	1.015 (0.975-1.056)	0.475
V ₂₅ (%)	58.2 (46.0-62.9)	52.0 (48.0-57.3)	1.014 (0.955-1.076)	0.659
V ₃₀ (%)	41.0 (35.1-47.6)	39.6 (31.1-42.1)	1.039 (0.961-1.122)	0.335
V ₃₅ (%)	29.6 (23.5-32.2)	26.9 (24.4-29.1)	1.032 (0.927-1.149)	0.569
V ₄₀ (%)	19.5 (16.8-23.3)	19.3 (13.3-20.7)	1.068 (0.928-1.229)	0.358
Rectum				
V ₂₀ (%)	80.9 (68.6-93.7)	78.0 (66.8-82.5)	1.015 (0.975-1.056)	0.475

V ₂₅ (%)	58.2 (46.0-62.9)	52.0 (48.0-57.3)	1.014 (0.955-1.076)	0.659
V ₃₀ (%)	41.0 (35.1-47.6)	39.6 (31.1-42.1)	1.039 (0.961-1.122)	0.335
V ₃₅ (%)	29.6 (23.5-32.2)	26.9 (24.4-29.1)	1.032 (0.927-1.149)	0.569
V ₄₀ (%)	19.5 (16.8-23.3)	19.3 (13.3-20.7)	1.068 (0.928-1.229)	0.358
Colon				
V ₂₀ (%)	100.0 (100.0-100.0)	100.0 (100.0-100.0)	NA	NA
V ₂₅ (%)	100.0 (100.0-100.0)	100.0 (100.0-100.0)	NA	NA
V ₃₀ (%)	100.0 (100.0-100.0)	100.0 (100.0-100.0)	0.982 (0.775-1.246)	0.883
V ₃₅ (%)	100.0 (99.1-100.0)	100.0 (100.0-100.0)	0.981 (0.915-1.052)	0.981
V ₄₀ (%)	80.4 (71.3-89.2)	79.3 (72.1-87.8)	1.005 (0.956-1.056)	0.851
Femoral head				
V ₂₀ (%)	35.4 (19.7-57.3)	48.6 (22.9-65.2)	0.994 (0.971-1.017)	0.586
V ₂₅ (%)	24.8 (13.7-40.1)	31.6 (16.0-45.1)	0.996 (0.967-1.026)	0.797
V ₃₀ (%)	17.7 (9.8-29.6)	22.6 (11.4-32.6)	0.997 (0.957-1.039)	0.877
V ₃₅ (%)	13.3 (7.4-20.0)	16.9 (8.6-24.5)	0.989 (0.935-1.047)	0.711
V ₄₀ (%)	8.9 (4.9-14.3)	11.3 (5.7-16.3)	0.993 (0.916-1.076)	0.858
Total pelvic lymphatic drainage dose (Gy)	45.0 (45.0-45.0)	45.0 (45.0-48.5)	1.194 (0.877-1.625)	0.261
Para-aortic extended field radiotherapy (performed/not performed)			1.485 (1.222-1.804)	0.999

ACCI: Adjusted Charlson Comorbidity Index; CCRT: Concurrent chemoradiotherapy; SCC: Squamous cell carcinoma; HPV: Human papillomavirus; NA: Not available; DAI: Disease activity index.

Table 6 Multivariate analysis of influencing factors of severe radiation enteritis in 50 patients with cervical cancer treated with radiotherapy

	B	SE	Exp (B)	95%CI		Wald χ^2	P value
				Lower	Upper		
DAI score	4.106	1.371	60.705	4.134	891.506	9.463	0.003
Anal bulge rating	2.925	1.229	18.636	1.677	207.114	7.664	0.017

DAI: Disease activity index; 95%CI: 95% confidence interval.

DISCUSSION

Nearly 80% of patients receiving pelvic radiotherapy for cervical cancer have early acute toxicity and 20% have late toxicity[16]. Acute RE often occurs in the second week after the start of radiotherapy and reaches a peak in the fourth to fifth week. The clinical manifestations are nausea, vomiting, abdominal pain, bloody stool, mucous stool, diarrhea, tenesmus, and incontinence. In gynecological tumor radiotherapy, the clinical application of IMRT shows an increasing trend. This radiotherapy technology can not only obtain satisfactory target coverage dose, but also effectively reduces the radiation dose of OAR[17,18], such as V45 of rectum and small intestine[19,20], and reduces the toxic reaction of bone marrow[21,22]. Although the incidence and severity of RE have decreased under the current radiotherapy technology, these issues cannot be ignored. In addition, mild RE is often underestimated or ignored due to the lack of clinical records and inadequate assessment of toxicity[23].

In this study, we preliminarily demonstrated the close relationship between the anal bulge rating and the DAI score and SARE. More importantly, this study is the first to develop and validate an easy-to-understand visual nomogram prediction model for cervical cancer radiotherapy patients and doctors, providing more personalized and accurate SARE prediction for cervical cancer patients undergoing radiotherapy. Many studies have reported that the most important factor affecting radiation proctitis is the total dose received by the rectum, and it is also affected by the radiotherapy technique, fraction dose, total dose received by the rectum/total volume, dose volume ratio of rectum to radiation dose,

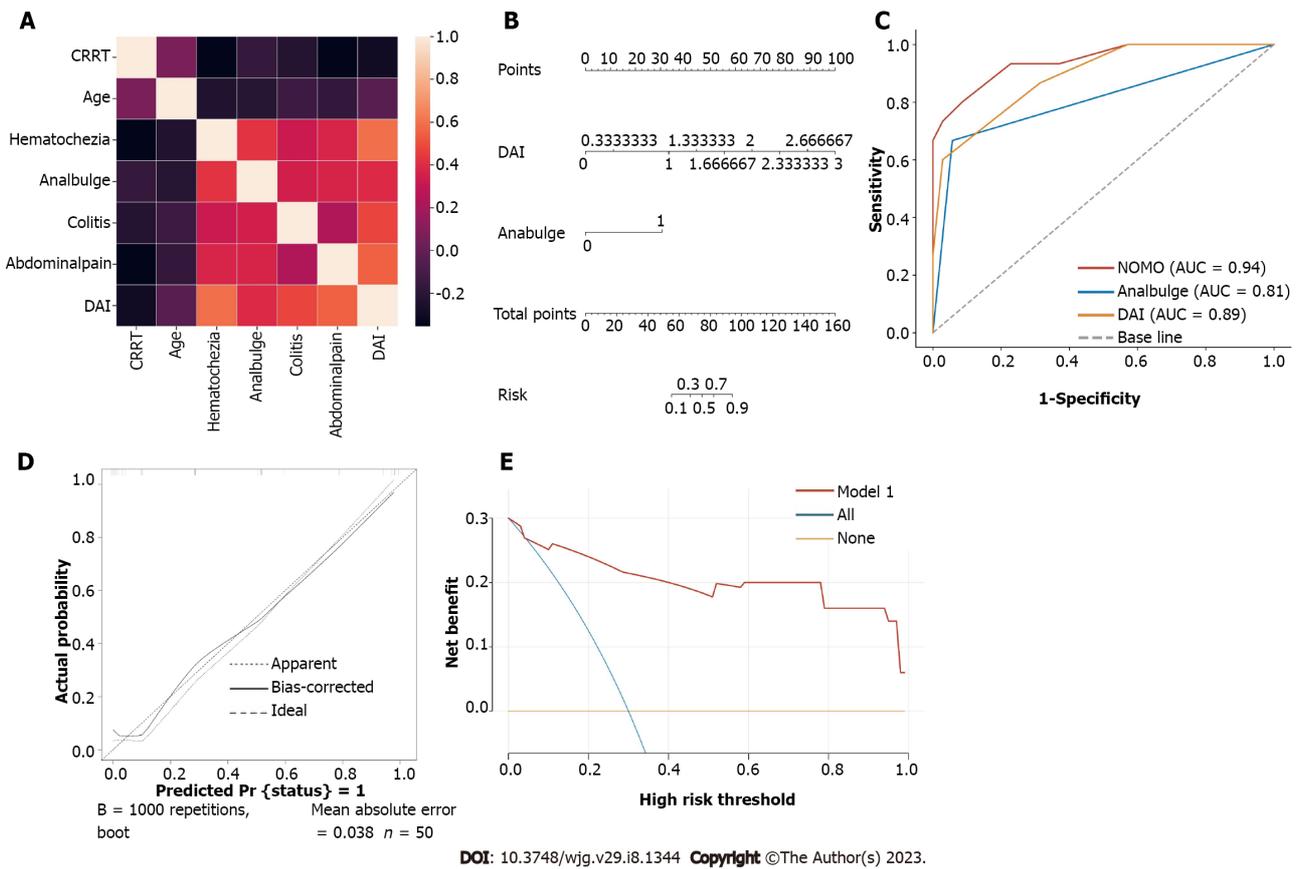


Figure 2 Development and validation of the nomogram. A: Kendall's rank correlation analyses among factors with $P < 0.05$ in univariate logistic regression; B: Nomogram predicting the occurrence of severe acute radiation enteritis (SARE). For each individual patient, two lines are drawn upward to determine the points received from the two variables in the nomogram; the sum of these points is located on the "Total Points" line, and a line is drawn downward to determine the likelihood of this patient to have SARE; C: Receiver operating characteristic curves of anal bulge rating. Disease activity index score vs the predictive model; D: Calibration curves of the nomogram predicting the occurrence of SARE. The x-axis and y-axis indicate the predicted and actual probabilities of having SARE, respectively; E: Decision curves of the nomogram predicting the occurrence of SARE. The x-axis shows the threshold probabilities. The y-axis measures the net benefit, which was calculated by adding the true positives and subtracting the false positives. NOMO: Nodal modulator; AUC: Area under the receiver operating characteristic curve; DAI: Disease activity index.

and uniformity of radiation dose distribution[4-6]. According to Seppenwoolde *et al*[24], rectal V40 ≥ 80% is a physical dose factor predicting VMAT-related rectal incontinence and diarrhea in patients with locally advanced cervical cancer. Ballari *et al*[25] suggested that small bowel V45 has predictive value for acute RE. In addition, individual factors such as IBD, collagen vascular disease, history of abdominal and pelvic surgery, history of pelvic inflammatory disease, diabetes mellitus, and acquired immunodeficiency syndrome affect the determination of radiation dose and volume[7-11]. In patients with cervical cancer, CCRT exerts radiosensitization and increases gastrointestinal toxicity by a factor of 3[18]. However, there is currently no consensus on the relative importance of these relevant predictors. In our study, the dose-volume of the small intestine receiving 20-40 Gy in the radical radiotherapy group of cervical cancer was significantly reduced compared with that in the postoperative adjuvant radiotherapy group, and the reduction of $V_{20\%}$, $V_{30\%}$ and V_{40} was statistically significant ($P < 0.05$), which was considered to be related to the anatomical displacement of organs after hysterectomy. After the removal of the tumor bed and uterus, part of the intestinal loop falls into the pelvic cavity[9], resulting in increased secondary toxicity of radiotherapy. In addition, the rectal $V_{30\%}$, $V_{35\%}$ and V_{40} in the radical radiotherapy group were significantly higher than those in the postoperative adjuvant radiotherapy group, and the increase of V_{40} was statistically significant ($P < 0.05$). The cervical cancer patients treated with radical radiotherapy in this study were FIGO IIB-IVA. CT or MRI showed obvious parametrial infiltration, and mesorectal invasion occurred in some patients, resulting in a relative increase in the volume of PTV to the left, right and dorsal side. Although we found dose-volume differences in the small intestine and rectum between groups, we were unable to demonstrate that a single physical parameter independently predicted RE or SARE. Therefore, more prospective, well-designed randomized controlled trials with larger sample sizes are needed for validation.

Acute RE is self-limited, and not all acute RE will be classified as chronic RE. The prevalence of chronic RE after 10 years of radiotherapy is 10%-20%[26]. In terms of pathogenesis, inflammation and cell death occur in the intestinal mucosa during the acute phase of RE, and sustained cytokine activation

in the submucosa leads to progressive ischemia, fibrosis, and stem cell loss[27]. In contrast, chronic RE is associated with progressive endarteritis obliterans, secondary to capillary fibrosis, leading to chronic ischemia and fibrosis of the affected bowel segments[28]. The outcome of chronic RE often induces intestinal obstruction, which is similar to the Koenig syndrome observed in Crohn's disease and is considered to be related to ileal stenosis. In view of the similarities between RE and IBD, we innovatively adopted the IBD evaluation index-DAI score. Univariate and multivariate logistic regression analysis confirmed that DAI score was independent of other physical parameters and clinical factors and could be used as a new predictor of SARE. In univariate analysis, abdominal pain, colitis, anal distension, hematochezia, DAI score, age, and CCRT were statistically correlated with the occurrence of severe acute RE ($P < 0.05$). Kendall analysis showed a relatively strong correlation between abdominal pain, hematochezia, and DAI scores. Therefore, DAI score is a convenient, non-invasive and highly beneficial diagnostic system for SARE.

The clinical manifestations of SARE are complex, and it is difficult for doctors, even with extensive clinical experience, to predict and diagnose the disease early, comprehensively and accurately. If cervical cancer can be identified and treated early before, during or after radiotherapy, the toxic and side effects of pelvic radiotherapy may be improved, and the probability of radiotherapy intolerance or even termination of radiotherapy may be reduced. The multi-factor prediction model is helpful to further improve the prediction of RE. Several prediction models have been described in previous studies, but they are still not used in clinic due to the low recognition ability of RE and the neglect of heterogeneity among different patients[29,30]. In this study, we established a SARE-SS evaluation system and found that anal bulge rating and DAI score were independent predictors of SARE through multivariate statistical analysis. The above predictors were further integrated into the nomogram to allow precise individualized SARE risk assessment for each patient. The AUC of the predictive model was 0.950, which was much higher than the two independent factors of DAI score and anal bulge score. The MAL thresholds for anal bulge rating and DAI score were 0.5 grade and 2.165 points, respectively, indicating an advantage over previous models in predicting RE. We adopted the Bootstrap validation method (including drawing the calibration curve and DCA curve), which showed a statistical advantage in a relatively small sample size statistic[31,32]. In later studies, we will expand the sample size, carry out prospective cohort studies, and use external validation methods to reduce data selection bias and increase test power.

Although our study has many advantages, there are still several limitations to be addressed. First, our sample size is small, so a large cohort is needed to further develop and validate the nomogram for predicting RE. Second, since this was a retrospective study, our study may have had a selection bias. Third, we only integrated the most important and commonly used parameters in clinic. From the perspective of precision medicine, we need to combine biological factors in the future including individual genomics, proteomics, metabolomics, microbiomics, real-time dosimetry and a wider range of clinical parameters to establish a comprehensive prediction model to improve the accuracy of RE prediction. Therefore, we will add biological factors to the analysis of related studies in later studies and need to establish a more comprehensive and specific prediction model of RE.

CONCLUSION

Our findings showed that during VMAT for cervical cancer, gastrointestinal and hematological system toxicity gradually increased with radiotherapy treatment and reached the peak at the end of radiotherapy. The main adverse reactions were diarrhea, abdominal pain, colitis, anal bulging and hematochezia. There were significant differences in V_{20} , V_{30} , V_{40} of small intestine and V_{40} of rectum between postoperative adjuvant radiotherapy and radical radiotherapy, which may be related to intestinal anatomical displacement caused by surgery and primary disease. However, there were no significant differences in the cumulative incidence of adverse reactions between the two groups. Anal bulge rating (> 0.500) and DAI score (> 2.165) were identified as independent predictors of SARE by multivariate analysis. The validity of our established nomogram was verified by the internal verification method. Although further external validation is needed, our results indicate that the nomogram has clinical value for the prediction and evaluation of SARE.

ARTICLE HIGHLIGHTS

Research background

Radiation enteritis (RE) not only seriously affects the quality of life of patients, but it also leads to radiotherapy intolerance or termination of radiotherapy. However, data on the clinical efficacy and side effects of volumetric modulated arc therapy (VMAT) for cervical cancer are limited.

Research motivation

If the incidence of RE in patients can be predicted in advance, and targeted clinical preventive treatment can be carried out, the side effects of radiotherapy in cervical cancer patients can be significantly reduced. Furthermore, accurate prediction of RE is essential for the selection of individualized radiation dose and the optimization of the radiotherapy plan.

Research objectives

To analyze the relationships between severe acute RE (SARE) of cervical cancer radiotherapy and clinical factors and dose-volume parameters retrospectively.

Research methods

We included 50 cervical cancer patients who received VMAT from September 2017 to June 2018 in the Department of Radiotherapy at The First Affiliated Hospital Soochow University. Clinical and dose-volume histogram factors of patients were collected. Logistic regression analysis was used to evaluate the predictive value of each factor for SARE. A nomogram to predict SARE was developed (SARE scoring system ≥ 3 points) based on the multiple regression coefficients; validity was verified by an internal verification method.

Research results

Gastrointestinal and hematological toxicity of cervical cancer VMAT gradually increased with radiotherapy and reached the peak at the end of radiotherapy. The main adverse reactions were diarrhea, abdominal pain, colitis, anal swelling, and blood in the stool. There was no significant difference in the incidence of gastrointestinal toxicity between the radical and postoperative adjuvant radiotherapy groups ($P > 0.05$). There were significant differences in the small intestine V20, V30, V40, and rectal V40 between adjuvant radiotherapy and radical radiotherapy after surgery ($P < 0.05$). Univariate and multivariate analysis revealed anal bulge rating (OR: 14.779, 95%CI: 1.281-170.547, $P = 0.031$) and disease activity index (DAI) score (OR: 53.928, 95%CI: 3.822-760.948, $P = 0.003$) as independent predictors of SARE.

Research conclusions

Anal bulge rating (> 0.500 grade) and DAI score (> 2.165 points) can predict SARE. The nomogram shows potential value in clinical practice.

Research perspectives

From the perspective of precision medicine, it will also be necessary to combine biological factors, such as individual genomics, proteomics, metabolomics, microbiomics, real-time dosimetry, and a wider range of clinical parameters to establish comprehensive predictive models. This study is a prospective study with a small sample size. In the later stage, we will expand the sample size, conduct prospective cohort studies, and use external validation methods to reduce data selection bias and increase test efficiency.

FOOTNOTES

Author contributions: Ma CY and Zhao J contributed to conceptualization and methodology; He XL and Gan GH contributed to data curation; Ma CY contributed to formal analysis; Ma CY, Zhou JY contributed to funding acquisition; Zhou JY, Xu XT, Qin SB contributed to project administration; Zhou JY, Xu XT, Qin SB, Wang LL, and Li L contributed to resources; Xu XT and Qin SB contributed to supervision; Ma CY contributed to validation and visualization; Ma CY and Zhao J contributed to roles/writing-original draft; Ma CY contributed to writing-review and editing.

Supported by the National Natural Science Foundation of China, No. 81602792 and No. 81602802; Project of State Key Laboratory of Radiation Medicine and Protection, Soochow University, No. GZK1202101; Suzhou Science and Technology Development Plan Project, No. KJXW2020008; and BOXI Natural Science Cultivation Foundation of China of the First Affiliated Hospital of Soochow University, No. BXQN202107.

Institutional review board statement: The study was reviewed and approved by the Ethics Committee of The First Affiliated Hospital of Soochow University Medical Ethics Committee [approval No. 2016(100)].

Informed consent statement: All enrolled patients provided signed informed consent.

Conflict-of-interest statement: The authors declare that they have no conflict of interest.

Data sharing statement: The datasets generated and analyzed during the present study are available from the corresponding author on reasonable request.

STROBE statement: The authors have read the STROBE Statement – checklist of items, and the manuscript was prepared and revised according to the STROBE Statement – checklist of items.

Open-Access: This article is an open-access article that was selected by an in-house editor and fully peer-reviewed by external reviewers. It is distributed in accordance with the Creative Commons Attribution NonCommercial (CC BY-NC 4.0) license, which permits others to distribute, remix, adapt, build upon this work non-commercially, and license their derivative works on different terms, provided the original work is properly cited and the use is non-commercial. See: <https://creativecommons.org/licenses/by-nc/4.0/>

Country/Territory of origin: China

ORCID number: Ju-Ying Zhou [0000-0002-2796-6388](https://orcid.org/0000-0002-2796-6388).

S-Editor: Chen YL

L-Editor: A

P-Editor: Chen YL

REFERENCES

- 1 Erratum: Global cancer statistics 2018: GLOBOCAN estimates of incidence and mortality worldwide for 36 cancers in 185 countries. *CA Cancer J Clin* 2020; **70**: 313 [PMID: [32767693](https://pubmed.ncbi.nlm.nih.gov/32767693/) DOI: [10.3322/caac.21609](https://doi.org/10.3322/caac.21609)]
- 2 Hyun Kim T, Choi J, Park SY, Lee SH, Lee KC, Yang DS, Shin KH, Cho KH, Lim HS, Kim JY. Dosimetric parameters that predict late rectal complications after curative radiotherapy in patients with uterine cervical carcinoma. *Cancer* 2005; **104**: 1304-1311 [PMID: [16078262](https://pubmed.ncbi.nlm.nih.gov/16078262/) DOI: [10.1002/ncr.21292](https://doi.org/10.1002/ncr.21292)]
- 3 van de Bunt L, van der Heide UA, Ketelaars M, de Kort GA, Jürgenliemk-Schulz IM. Conventional, conformal, and intensity-modulated radiation therapy treatment planning of external beam radiotherapy for cervical cancer: The impact of tumor regression. *Int J Radiat Oncol Biol Phys* 2006; **64**: 189-196 [PMID: [15978745](https://pubmed.ncbi.nlm.nih.gov/15978745/) DOI: [10.1016/j.ijrobp.2005.04.025](https://doi.org/10.1016/j.ijrobp.2005.04.025)]
- 4 Forrest JL, Ackerman I, Barbera L, Barnes EA, Davidson M, Kiss A, Thomas G. Patient outcome study of concurrent chemoradiation, external beam radiotherapy, and high-dose rate brachytherapy in locally advanced carcinoma of the cervix. *Int J Gynecol Cancer* 2010; **20**: 1074-1078 [PMID: [20683420](https://pubmed.ncbi.nlm.nih.gov/20683420/) DOI: [10.1111/IGC.0b013e3181e6f321](https://doi.org/10.1111/IGC.0b013e3181e6f321)]
- 5 Kozak MM, Koenig JL, von Eyben R, Kidd EA. Less Than Whole Uterus Irradiation for Locally Advanced Cervical Cancer Maintains Locoregional Control and Decreases Radiation Dose to Bowel. *Pract Radiat Oncol* 2019; **9**: e164-e171 [PMID: [30395974](https://pubmed.ncbi.nlm.nih.gov/30395974/) DOI: [10.1016/j.prr.2018.10.009](https://doi.org/10.1016/j.prr.2018.10.009)]
- 6 Ray A, Sarkar B. Small bowel toxicity in pelvic radiotherapy for postoperative gynecological cancer: comparison between conformal radiotherapy and intensity modulated radiotherapy. *Asia Pac J Clin Oncol* 2013; **9**: 280-284 [PMID: [23279854](https://pubmed.ncbi.nlm.nih.gov/23279854/) DOI: [10.1111/ajco.12049](https://doi.org/10.1111/ajco.12049)]
- 7 Willett CG, Ooi CJ, Zietman AL, Menon V, Goldberg S, Sands BE, Podolsky DK. Acute and late toxicity of patients with inflammatory bowel disease undergoing irradiation for abdominal and pelvic neoplasms. *Int J Radiat Oncol Biol Phys* 2000; **46**: 995-998 [PMID: [10705022](https://pubmed.ncbi.nlm.nih.gov/10705022/) DOI: [10.1016/s0360-3016\(99\)00374-0](https://doi.org/10.1016/s0360-3016(99)00374-0)]
- 8 Lin A, Abu-Isa E, Griffith KA, Ben-Josef E. Toxicity of radiotherapy in patients with collagen vascular disease. *Cancer* 2008; **113**: 648-653 [PMID: [18506734](https://pubmed.ncbi.nlm.nih.gov/18506734/) DOI: [10.1002/ncr.23591](https://doi.org/10.1002/ncr.23591)]
- 9 Nuytens JJ, Robertson JM, Yan D, Martinez A. The influence of small bowel motion on both a conventional three-field and intensity modulated radiation therapy (IMRT) for rectal cancer. *Cancer Radiother* 2004; **8**: 297-304 [PMID: [15561595](https://pubmed.ncbi.nlm.nih.gov/15561595/) DOI: [10.1016/j.canrad.2004.08.001](https://doi.org/10.1016/j.canrad.2004.08.001)]
- 10 Andreyev HJ, Wotherspoon A, Denham JW, Hauer-Jensen M. "Pelvic radiation disease": new understanding and new solutions for a new disease in the era of cancer survivorship. *Scand J Gastroenterol* 2011; **46**: 389-397 [PMID: [21189094](https://pubmed.ncbi.nlm.nih.gov/21189094/) DOI: [10.3109/00365521.2010.545832](https://doi.org/10.3109/00365521.2010.545832)]
- 11 Gichangi P, Bwayo J, Estambale B, Rogo K, Njuguna E, Ojwang S, Temmerman M. HIV impact on acute morbidity and pelvic tumor control following radiotherapy for cervical cancer. *Gynecol Oncol* 2006; **100**: 405-411 [PMID: [16274737](https://pubmed.ncbi.nlm.nih.gov/16274737/) DOI: [10.1016/j.ygyno.2005.10.006](https://doi.org/10.1016/j.ygyno.2005.10.006)]
- 12 Dröge LH, von Sivers FF, Schirmer MA, Wolff HA. Conventional 3D conformal radiotherapy and volumetric modulated arc therapy for cervical cancer: Comparison of clinical results with special consideration of the influence of patient- and treatment-related parameters. *Strahlenther Onkol* 2021; **197**: 520-527 [PMID: [33938967](https://pubmed.ncbi.nlm.nih.gov/33938967/) DOI: [10.1007/s00066-021-01782-5](https://doi.org/10.1007/s00066-021-01782-5)]
- 13 Andreyev HJ, Davidson SE, Gillespie C, Allum WH, Swarbrick E; British Society of Gastroenterology; Association of Colo-Proctology of Great Britain and Ireland; Association of Upper Gastrointestinal Surgeons; Faculty of Clinical Oncology Section of the Royal College of Radiologists. Practice guidance on the management of acute and chronic gastrointestinal problems arising as a result of treatment for cancer. *Gut* 2012; **61**: 179-192 [PMID: [22057051](https://pubmed.ncbi.nlm.nih.gov/22057051/) DOI: [10.1136/gutjnl-2011-300563](https://doi.org/10.1136/gutjnl-2011-300563)]
- 14 Abu-Rustum NR, Yashar CM, Bean S, Bradley K, Campos SM, Chon HS, Chu C, Cohn D, Crispens MA, Damast S, Fisher CM, Frederick P, Gaffney DK, Giuntoli R, Han E, Huh WK, Lurain Iii JR, Mariani A, Mutch D, Nagel C, Nekhlyudov L, Fader AN, Remmenga SW, Reynolds RK, Sisodia R, Tillmanns T, Ueda S, Urban R, Wyse E, McMillian NR, Motter AD. NCCN Guidelines Insights: Cervical Cancer, Version 1.2020. *J Natl Compr Canc Netw* 2020; **18**: 660-666 [PMID: [32502976](https://pubmed.ncbi.nlm.nih.gov/32502976/) DOI: [10.6004/jnccn.2020.0027](https://doi.org/10.6004/jnccn.2020.0027)]
- 15 Eifel PJ, Jhingran A, Bodurka DC, Levenback C, Thames H. Correlation of smoking history and other patient

- characteristics with major complications of pelvic radiation therapy for cervical cancer. *J Clin Oncol* 2002; **20**: 3651-3657 [PMID: 12202666 DOI: 10.1200/JCO.2002.10.128]
- 16 **Shadad AK**, Sullivan FJ, Martin JD, Egan LJ. Gastrointestinal radiation injury: symptoms, risk factors and mechanisms. *World J Gastroenterol* 2013; **19**: 185-198 [PMID: 23345941 DOI: 10.3748/wjg.v19.i2.185]
 - 17 **Mell LK**, Sirák I, Wei L, Tarnawski R, Mahantshetty U, Yashar CM, McHale MT, Xu R, Honerkamp-Smith G, Carmona R, Wright M, Williamson CW, Kasaová L, Li N, Kry S, Michalski J, Bosch W, Straube W, Schwarz J, Lowenstein J, Jiang SB, Saenz CC, Plaxe S, Einck J, Khorprasert C, Koonings P, Harrison T, Shi M, Mundt AJ; INTERTECC Study Group. Bone Marrow-sparing Intensity Modulated Radiation Therapy With Concurrent Cisplatin For Stage IB-IVA Cervical Cancer: An International Multicenter Phase II Clinical Trial (INTERTECC-2). *Int J Radiat Oncol Biol Phys* 2017; **97**: 536-545 [PMID: 28126303 DOI: 10.1016/j.ijrobp.2016.11.027]
 - 18 **Zhang K**, Wang H, Wang Z, Li F, Cui Y, Ma S, Chen R, Wang Y, Guo S, Wei Y. Intensity-modulated radiation therapy (IMRT)-based concurrent chemoradiotherapy (CCRT) with Endostar in patients with pelvic locoregional recurrence of cervical cancer: Results from a hospital in the Qinghai-Tibet Plateau. *Medicine (Baltimore)* 2020; **99**: e21966 [PMID: 33285664 DOI: 10.1097/MD.00000000000021966]
 - 19 **Chino J**, Annunziata CM, Beriwal S, Bradfield L, Erickson BA, Fields EC, Fitch K, Harkenrider MM, Holschneider CH, Kamrava M, Leung E, Lin LL, Mayadev JS, Morcos M, Nwachukwu C, Petereit D, Viswanathan AN. Radiation Therapy for Cervical Cancer: Executive Summary of an ASTRO Clinical Practice Guideline. *Pract Radiat Oncol* 2020; **10**: 220-234 [PMID: 32473857 DOI: 10.1016/j.prro.2020.04.002]
 - 20 **Yang B**, Zhu L, Cheng H, Li Q, Zhang Y, Zhao Y. Dosimetric comparison of intensity modulated radiotherapy and three-dimensional conformal radiotherapy in patients with gynecologic malignancies: a systematic review and meta-analysis. *Radiat Oncol* 2012; **7**: 197 [PMID: 23176540 DOI: 10.1186/1748-717X-7-197]
 - 21 **Contreras J**, Srivastava A, Chundury A, Schwarz JK, Markovina S, Thaker PH, Massad LS, Mutch DG, Powell MA, Grigsby PW, Lin AJ. Long-term outcomes of intensity-modulated radiation therapy (IMRT) and high dose rate brachytherapy as adjuvant therapy after radical hysterectomy for cervical cancer. *Int J Gynecol Cancer* 2020; **30**: 1157-1161 [PMID: 32527770 DOI: 10.1136/ijgc-2020-001412]
 - 22 **Liu Y**, Yu J, Qian L, Zhang H, Ma J. Extended field intensity-modulated radiotherapy plus concurrent nedaplatin treatment in cervical cancer. *Oncol Lett* 2016; **11**: 3421-3427 [PMID: 27123128 DOI: 10.3892/ol.2016.4416]
 - 23 **Wang W**, Zhang F, Hu K, Hou X. Image-guided, intensity-modulated radiation therapy in definitive radiotherapy for 1433 patients with cervical cancer. *Gynecol Oncol* 2018; **151**: 444-448 [PMID: 30266261 DOI: 10.1016/j.ygyno.2018.09.024]
 - 24 **Seppenwoolde Y**, Majercakova K, Buschmann M, Dörr E, Sturdza AE, Schmid MP, Pötter R, Georg D. Early morbidity and dose-volume effects in definitive radiochemotherapy for locally advanced cervical cancer: a prospective cohort study covering modern treatment techniques. *Strahlenther Onkol* 2021; **197**: 505-519 [PMID: 33929558 DOI: 10.1007/s00066-021-01781-6]
 - 25 **Ballari N**, Rai B, Bahl A, Mittal BR, Ghoshal S. Prospective observational study evaluating acute and delayed treatment related toxicities of prophylactic extended field volumetric modulated arc therapy with concurrent cisplatin in cervical cancer patients with pelvic lymph node metastasis. *Tech Innov Patient Support Radiat Oncol* 2021; **17**: 48-56 [PMID: 33748442 DOI: 10.1016/j.tipsro.2021.02.009]
 - 26 **Shen XJ**, Liu L, Zhu JY. Radiofrequency ablation in a patient with radiation enteritis: A case report. *Medicine (Baltimore)* 2018; **97**: e13328 [PMID: 30461649 DOI: 10.1097/MD.00000000000013328]
 - 27 **Denham JW**, Hauer-Jensen M. The radiotherapeutic injury--a complex 'wound'. *Radiother Oncol* 2002; **63**: 129-145 [PMID: 12063002 DOI: 10.1016/s0167-8140(02)00060-9]
 - 28 **Yarnold J**, Brotons MC. Pathogenetic mechanisms in radiation fibrosis. *Radiother Oncol* 2010; **97**: 149-161 [PMID: 20888056 DOI: 10.1016/j.radonc.2010.09.002]
 - 29 **Loge L**, Florescu C, Alves A, Menahem B. Radiation enteritis: Diagnostic and therapeutic issues. *J Visc Surg* 2020; **157**: 475-485 [PMID: 32883650 DOI: 10.1016/j.jvisurg.2020.08.012]
 - 30 **Hale MF**. Radiation enteritis: from diagnosis to management. *Curr Opin Gastroenterol* 2020; **36**: 208-214 [PMID: 32141897 DOI: 10.1097/MOG.0000000000000632]
 - 31 **Halabi S**, Li C, Luo S. Developing and Validating Risk Assessment Models of Clinical Outcomes in Modern Oncology. *JCO Precis Oncol* 2019; **3** [PMID: 31840130 DOI: 10.1200/PO.19.00068]
 - 32 **Steyerberg EW**, Harrell FE Jr. Prediction models need appropriate internal, internal-external, and external validation. *J Clin Epidemiol* 2016; **69**: 245-247 [PMID: 25981519 DOI: 10.1016/j.jclinepi.2015.04.005]



Published by **Baishideng Publishing Group Inc**
7041 Koll Center Parkway, Suite 160, Pleasanton, CA 94566, USA
Telephone: +1-925-3991568
E-mail: bpgoffice@wjgnet.com
Help Desk: <https://www.f6publishing.com/helpdesk>
<https://www.wjgnet.com>

