

Predictors of Quality of Life and Social Support as a Mediator between Defecation Function and Quality of Life among Rectal Cancer Patients



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Purpose: To identify the factors influencing Quality of Life (QoL) in rectal cancer patients and to examine whether perceived social support mediates the relationship between defecation function and QoL. **Methods:** Using self-report questionnaires, a descriptive survey was conducted with 131 rectal cancer patients who underwent anal sphincter-saving resection, in a large medical center in Korea. Data were collected between September and November, 2014, using research instruments validated in the Korean language. The data were analyzed using SPSS/WIN 21.0 and AMOS 21.0. **Results:** The mean age and post-surgical follow-up period of the participants were 59.0 ± 10.2 years and 15.9 ± 9.8 months, respectively. Multiple linear regression analysis revealed that, after adjusting for age and gender, defecation function, perceived social support, and employment status predicted QoL. Path analysis showed that perceived social support partially mediated the relationship between defecation function and QoL with a significant indirect effect. **Conclusion:** These results suggest that a positive perception of social support from family and healthcare providers is necessary to improve the QoL of patients experiencing defecation dysfunction after rectal cancer surgery. To this end, periodic counseling and education from outpatient healthcare providers, during the postoperative follow-up period, could encourage patients to perceive higher social support.

Key Words: Rectal neoplasms; Defecation; Social support; Quality of life

INTRODUCTION

Colorectal cancer is the third most common cancer, accounting for 12.4% of all cancers in Korea. Approximately 45% of colorectal cancer cases arise in the rectum, accounting for 5.4% of all cancer cases. Rates of colorectal cancer are increasing faster than those of other cancer types with high prevalence rates [1]. The increased incidence of colorectal cancer in recent years has largely been attributed to a westernized diet, increased alcohol consumption, and decreased physical activity [2]. Recent advances in treatment technology have resulted in improved survival of patients with colorectal cancer, and their Quality of Life (QoL) is an increasingly important aspect of long-term care for chronic illness. Approximately 80% of patients with rectal cancer undergo sphincter-saving surgery; the ultimate goal

of this surgical procedure is to minimize complications through radical resection, thereby preserving the anal sphincter muscle and improving QoL [3].

A systematic review reported that up to 60% of rectal cancer patients who underwent sphincter-saving resection reported urgency of defecation, frequent bowel movements, fecal incontinence, and changes in erectile function [4]. Defecation dysfunction is characterized by frequent passage of stools, feeling of defecation, urgency of defecation, loose feces, feeling of incomplete defecation, and constipation [5]. Particularly, patients with rectal cancer who undergo sphincter-saving resection persistently report defecation function problems as a long-term adverse effect over the course of four years after definitive treatment [6]. They are also often treated with a combination of radiation therapy and chemotherapy, which can cause symptoms of

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Received: May 23, 2019 / Revised: Jul 25, 2019 / Accepted: Aug 23, 2019

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defecation dysfunction and negatively affect physical, psychological, and social functioning during the treatment period, in addition to a marked decline in QoL [6]. Furthermore, patients with rectal cancer are reported to have lower QoL than patients with gastric or colon cancer due to the presence of symptoms of defecation dysfunction [7].

Previous studies have found that anxiety/depression [8], younger age, lower economic status, and higher nutritional risk are factors affecting QoL in patients with surgically treated colorectal cancer [9]. In addition, social support from the family and from healthcare professionals was found to predict lower stress and greater QoL in colorectal cancer patients [10], whereas symptom distress, such as defecation problems, made patients more depressed, and lowered perceived social support, as well as QoL [11]. Cohort studies report that colorectal cancer patients with high perceived social support and no psychological distress have better QoL one year after surgery [12], and about one-third of patients have decreasing levels of perceived social support two years after surgery, which is closely related to lower QoL [13]. Therefore, the support of healthcare professionals and the family seems to be very important for the patients, who need to follow up through outpatient clinic for a considerable period after surgery, and it is necessary to consider the role of social support in planning interventions for this population.

Perceived social support is likely to be higher if the patient feels that his or her needs are met by the healthcare professional or the family during the post-operative recovery period. However, a systematic review reported that colorectal cancer patients expressed diverse unmet needs after surgery, encompassing emotional support, information about diet and long-term self-management of symptoms, and complications at home [14]. They also experienced a lack of supportive care after discharge and wanted counseling received from healthcare professionals during the preoperative period to extend to the postoperative period, including instructions on the use of self-management skills for their functional complaints or symptoms [3, 15]. The information needs of patients can be met by providing social support in the form of counseling and education by nurses; such psychological interventions can help them overcome post-surgical symptoms and successfully cope with the disease, thereby improving their QoL [12].

To summarize, rectal cancer patients with defecation dysfunction are likely to be more satisfied if they receive social support from their families and health care professionals. In other words, perception of social support can be hypothesized to be an indirect variable to explain the relationship between defecation function and QoL in patients

with rectal cancer during the follow-up period. Social support has been found to mediate between symptom distress and QoL in gastric cancer patients [16]. Therefore, the purpose of the present study is two-fold: (1) to identify factors predicting QoL and, (2) to investigate whether perceived social support mediates the relationship between defecation function and QoL, in patients with rectal cancer, after anal sphincter-saving surgery.

METHODS

1. Study Design

The study adopted a descriptive, correlational design, to examine the relationship between defecation function, perceived social support, and QoL in patients with rectal cancer.

2. Study Participants

The study participants were patients from a large tertiary referral hospital with 2,700 beds in Seoul, Korea, who had been diagnosed with rectal cancer and were on outpatient treatment after undergoing sphincter-saving resection. Using the G*power program, the significance level (α) of .05, power ($1-\beta$) of .80, median effect size of .15 for regression analysis, and considering 13 independent variables (gender, age, marital status, living arrangement, education, employment status, monthly income, presence of caregiver, religion, surgical approach, time since surgery, social support and defecation dysfunction), the minimum required sample size was calculated to be 131. A convenience sample of 140 patients was recruited, taking into consideration dropouts due to possible exhaustion or inconvenience during treatment. Nine patients were excluded because they were unable to answer the questionnaire sufficiently, or experienced cancer recurrence after treatment. The remaining 131 patients were included in the final analysis.

Participants were included in this study if 1) they were aged 20 years or older, diagnosed with rectal cancer, regardless of cancer stage, 2) they were between three months to five years since undergoing a low anterior resection, an ultralow anterior resection, or an inter-sphincteric resection and, 3) they continued to follow-up at the outpatient department or the colorectal clinic. These patients were particularly targeted because complications frequently occur within three months after sphincter-saving surgery, while survival after five years indicates that the patient is completely cured [17]. The following criteria

were used to exclude participants from the study: 1) those who had temporary colostomy after low anterior resection for less than three months since stoma closure and, 2) those who were currently on chemotherapy owing to recurrence of cancer, did not undergo colostomy reversal after sphincter-saving surgery, or underwent a re-surgery to create an ostomy. All participants were able to communicate, read, and write, understand the purpose of the study, and provided consent to participate in the study.

3. Instruments

1) Defecation function

Defecation function was evaluated using the Memorial Sloan Kettering Cancer Center Bowel Function Instrument (MSKCC BFI) developed by Temple et al. [18], which is designed specifically for evaluating bowel function after sphincter-preserving surgery for rectal cancer. The Korean version of MSKCC BFI has been validated in patients with colorectal cancer [19] and was authorized for use in this study by the author. The MSKCC BFI consists of 18 questions and adopts a four-week recall period and equal-weighting scoring. The indices of clinically meaningful characteristics of defecation, such as frequent defecation, dietary characteristics, and fecal urgency/incontinence, are scored using a five-point scale ranging from “always” (1) to “never” (5). Five items are reverse scored. The total score ranges from 18 to 90, with higher scores indicating better defecation function. Cronbach’s α values of the original tool [18], of the Korean version [19] were .87 and .78, respectively, and as found in the current study was .81.

2) Social support

Perceived social support was evaluated using the Multi-dimensional Scale of Perceived Social Support (MSPSS) developed by Zimet et al. [20]. The MSPSS has been translated into Korean and validated in a study of unemployed people [21]. The Korean version was authorized for use in this study by the original author. The MSPSS contains 12 items that subjectively measure perceived social support using three subscales: family, friends, and significant others. In this study, items relating to support from “significant others” and a “special person” referred to the support from healthcare professionals such as a nurse.

In the original MSPSS, the 12 items are rated on a seven-point Likert-type scale. In this study, the 12 items were rated on a five-point Likert-type scale, with answers ranging from “strongly disagree” (1) to “strongly agree” (5). Higher scores indicate higher perceived social support. Cronbach’s α values of the original study [20], the

Korean version [21], and in the current study were .85, 0.89, and .82, respectively. Cronbach’s α values of the subscales were .91 for family support, .91 for friends’ support, and .92 for significant others’ support.

3) Quality of life

QoL was evaluated using the Functional Assessment of Cancer Therapy-Colorectal (FACT-C) instrument developed by Ward et al. [22]. The FACT-C has been translated into Korean, and its reliability and validity have been tested [23]. Authorization from FACIT (www.facit.org) was received prior to the use of the FACT-C in this study. The instrument is divided into five domains: physical (7 items), social/family (7 items), emotional (6 items), functional (7 items), and colorectal cancer-specific (7 items). The 34 items are rated on a five-point scale ranging from “never” (0) to “always” (4). Negative items are reverse scored. Higher scores indicate higher perceived QoL. Cronbach’s α values of the original tool [22], the tool validated in Korean [23], and as found in the current study were .87, .86, and .92, respectively. Cronbach’s α values of the FACT-C subscales were .83 for physical well-being, .88 for social/family well-being, .74 for emotional well-being, .92 for functional well-being, and .63 for colorectal cancer-specific items.

4. Data Collection

Data were collected from September to November 2014. All participants were receiving outpatient follow-up treatment at the colorectal cancer clinic after surgery for rectal cancer. The first author of this study, who consults patients as a coordinator at the colon cancer clinic, explained the purpose of the study to the participants and collected data through one-on-one interviews. Participants were asked to respond to a self-report questionnaire. If they had difficulty understanding the text, or if they had poor vision, the researcher read the questions aloud and recorded their responses. We provided a gift to all research participants for participating in the survey. Information about disease- and surgery-related characteristics were collected from electronic medical records.

5. Ethical Consideration

This study was approved by the institute ethics committee of the hospital (AMC #2014-0951). Participants were informed of the purpose of the study, and possible risks and benefits related to their participation. Moreover, they were told that their participation was voluntary and that

they had the right to withdraw from the study at any time. All participants were asked to sign a written consent form prior to participating in the survey.

6. Data Analysis

The collected data were analyzed using SPSS/WIN version 21.0 and AMOS version 21.0 (IBM Corp., Armonk, NY, USA). Descriptive statistics were used for the general characteristics of the participants, and means and standard deviations were used to describe defecation function (MSKCC BFI score), social support (MSPSS score), and QoL (FACT-C score). Student's *t* test and one-way Analysis of Variance (ANOVA) were used to examine the differences between the two groups according to the general characteristics of the participants. Scheffé's post hoc test was used to compare significance among multiple categories. Pearson's correlation coefficient was used to evaluate the correlations between the research variables, and multiple linear regression analysis was used to determine factors associated with QoL.

Path analysis was used to verify causal relationships between direct, indirect, and total effects of the three variables. Regression coefficient, standardized coefficient, standard error, critical ratio, and *p* values were used to test the significance of paths in the structural equation model. Bootstrapping method (1:1,000) was used in examining the statistical significance of the model's direct, indirect, and total effects.

RESULTS

1. Descriptive Statistics of Study Participants and Measured Variables

Among the study participants, 61.8% were male and 87% were married. The mean age of the participants was 59.0 ± 10.2 years, and 68.7% had a high school diploma or college degree. As to the characteristics of their surgery, 46.6% had undergone robotic surgery, and 59.6% a lower anterior rectal resection. Additionally, 51.9% had cancer in the stages II and III, 40.5% had a temporary stoma, and 56.5% underwent chemotherapy. Among the patients, 47.3% had undergone surgery in the previous 12 months, and the average time elapsed since surgery was 15.93 ± 9.82 months (Table 1). The average scores for defecation function (MSKCC BFI), social support (MSPSS), and QoL (FACT-C) were 65.31 ± 8.33 , 43.14 ± 9.71 , and 98.80 ± 19.62 , respectively.

Table 1. Demographic and Clinical Characteristics of Study Participants (N=131)

Characteristics	Categories	n (%) or M \pm SD
Gender	Men	81 (61.8)
	Women	50 (38.2)
Age (year)	30~59	67 (51.2)
	60~82	64 (48.8)
		59.03 \pm 10.21
Marital status	Married	114 (87.0)
	Single or widowed	17 (13.0)
Living with	Family	121 (92.4)
	Alone	10 (7.6)
Education	\leq Middle school	41 (31.3)
	High school	56 (42.7)
	\geq College	34 (26.0)
Employment status	Employed	63 (48.1)
	Unemployed	68 (51.9)
Monthly income (USD)	< 2,000	54 (41.2)
	2,000~4,000	48 (36.6)
	> 4,000	29 (22.2)
Presence of caregiver	Yes	92 (70.2)
	No	39 (29.8)
Surgical approach	Robotic	61 (46.6)
	Laparoscopy	17 (13.0)
	Open	53 (40.4)
Operative method	LAR	78 (59.6)
	uLAR	40 (30.5)
	uLAR/ISR	13 (9.9)
Cancer stage	0	14 (10.7)
	I	49 (37.4)
	II	32 (24.4)
	III	36 (27.5)
Anastomosis (cm from AV)	≤ 5	97 (74.0)
	≥ 6	34 (26.0)
		4.01 \pm 2.53
Temporary stoma	Yes	53 (40.5)
	No	78 (59.5)
Experience of radiotherapy	Yes	43 (32.8)
	No	88 (67.2)
Experience of chemotherapy	Yes	74 (56.5)
	No	57 (43.5)
Time since surgery (month)	3~12	62 (47.3)
	13~24	32 (24.5)
	25~49	37 (28.2)
		15.93 \pm 9.82

AV=anal verge; ISR=intersphincteric resection; LAR=low anterior resection; uLAR=ultralow anterior resection.

2. Relationships between the Variables and Factors Affecting QoL

Pearson correlation coefficients showed significant relationships between defecation function and social support ($r=.20, p=.025$), between defecation function and QoL ($r=.31, p=.001$), and between social support and QoL ($r=.38, p=.001$). Bivariate analyses showed that QoL was significantly associated with education level ($F=3.51, p=.033$) and employment status ($t=2.87, p=.005$). High school graduates had a lower QoL than those with middle school or lower education, and those who graduated from college. Additionally, QoL was significantly higher in the employed group than in the unemployed group (Table 2).

In order to confirm the predictors of QoL among the patients, multiple linear regression analysis was conducted using forward selection. Among the independent variables, variance inflation factor (1.02~1.05) and tolerance (0.95~0.98) were within the normal range, satisfying the basic assumptions of multicollinearity ($p < .050$). In addition to defecation function and social support, education level and employment status, which were statistically sig-

nificant in the bivariate analysis, were included in the regression analysis. When adjusted for age and gender, the final regression model including defecation function, social support and employment status was statistically significant ($F=7.14, p < .001$) and accounted for the 25% of the variance (adjusted $R^2=.25$) (Table 3).

3. Path Analysis for the Mediating Effect of Social Support

The path model was established based on previous studies and tested for the mediating effect of social support on the relationship between defecation function and QoL (Figure 1). The initial model was confirmed to be a saturated model with zero degrees of freedom and $\chi^2=0.00$, implying a perfect fit for the data, and therefore, it was not necessary to confirm the fitness index of the model. Standardized coefficients for the path model were all statistically significant ($p < .050$). All three paths were statistically significant: defecation function affected social support ($\beta=.19, p=.031$), social support affected QoL ($\beta=.33, p=.005$), and defecation function affected QoL ($\beta=.25, p=.021$).

Defecation function had a positive direct effect on social

Table 2. Relationships between Participant Characteristics and Quality of Life (N=131)

Characteristics	Categories	M±SD	t or F	p	Scheffé
Gender	Men	100.53±19.50	1.27	.205	
	Women	96.21±19.82			
Age (year)	< 60	101.73±19.41	1.85	.066	
	≥ 60	95.44±19.50			
Marital status	Married	98.74±20.01	-0.12	.901	
	Single or widowed	99.32±8.13			
Living arrangements	With family	98.82±19.90	0.02	.978	
	Alone	99.00±17.02			
Education	≤ Middle school ^a	103.91±19.33	3.51	.033	b < a, c
	High school ^b	93.82±18.13			
	≥ College ^c	101.01±21.14			
Employment status	Employed	103.80±17.61	2.87	.005	
	Unemployed	94.23±20.42			
Religion	Yes	98.31±18.90	0.33	.739	
	No	99.52±20.81			
Monthly income (USD)	< 2,000	96.43±19.54	0.85	.426	
	2,000~4,000	99.51±18.80			
	> 4,000	102.21±21.42			
Presence of caregiver	Yes	100.34±19.42	1.31	.191	
	No	95.40±20.10			
Time since surgery (month)	3~12	100.52±19.21	0.56	.570	
	13~24	96.00±22.43			
	25~49	98.31±18.14			

support ($\beta = .38, p = .011$), social support had a positive direct effect ($\beta = .49, p = .005$) on QoL, and defecation function had a positive direct effect ($\beta = .38, p = .021$) and a positive indirect effect through social support ($\beta = .16, p = .007$) on QoL (total effect = $0.47, p = .009$). The bootstrapping test for the indirect effect of social support on the relationship between defecation function and QoL showed a statistically significant relationship ($\beta = .06$, standard error = $.19, p = .005$, bias-corrected 95% confidence interval = $.02 \sim .06$). Thus, social support mediated the relationship between defecation function and QoL (Table 4).

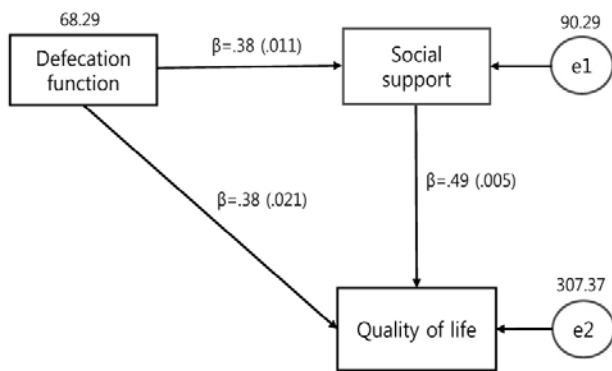


Figure 1. Path diagram of the model.

DISCUSSION

This study was conducted to identify the factors affecting QoL in patients with rectal cancer and to clarify the role of social support in relation to defecation function and QoL. Multiple regression analysis revealed that post-operative defecation function in rectal cancer patients significantly predicted QoL, which means that better defecation function was associated with higher QoL. This finding supports previous studies showing that rectal cancer patients with fecal incontinence and urgency after sphincter-saving resection surgery had significantly lower QoL than those without such problems [6,24]. In addition, this finding is consistent with the finding that defecation dysfunction is a major predictor of QoL following sphincter-preserving resection [25]. QoL of patients with rectal cancer was also closely associated with the severity of the low anterior resection syndrome [26] and cancer symptoms [27]. Changes in defecation function in rectal cancer patients was most pronounced during the first months after surgery, improved during the first year, and reached a steady state in one-two years after surgery [25]. However, since the time elapsed since surgery ranged from three to forty-nine months for participants in the study, additional longitudinal studies are needed to confirm the effects of

Table 3. Multiple Linear Regression Analysis for the Factors on Quality of Life (N=131)

Variables	B	SE	β	t	p	95% CI	
(Constant)	27.43	18.43		1.49	.139		
Social support	0.63	0.16	.32	3.04	< .001	0.27~0.92	
Defecation function	0.56	0.19	.25	3.04	.003	0.13~0.87	
Employment status (ref.: unemployed)	Employed	10.14	3.26	.26	3.11	.002	-18.60~-3.56
Education (ref.: ≤ middle school)	High school	-9.19	3.79	-.23	-1.93	.057	-9.20~0.23
	≥ College	-7.66	4.34	-.17	-1.76	.080	-11.89~0.79
$R^2 = .29, \text{Adj. } R^2 = .25, F = 7.14, p < .001$							

† Adjusted for age and gender; CI=confidence interval; SE=standard error.

Table 4. Path Coefficients between Endogenous and Exogenous Variables in the Model

Endogenous variables	Exogenous variables	B	β	SE	CR (p)	Standardized direct effect (p)	Standardized indirect effect (p)	Standardized total effect (p)
Social support	Defecation function	0.23	.19	.10	2.27 (.031)	.38 (.011)		.38 (.011)
Quality of life	Social support	0.67	.33	.16	4.07 (.005)	.49 (.005)		.49 (.005)
Quality of life	Defecation function	0.58	.25	.19	3.05 (.021)	.38 (.021)	.16 (.007)	.47 (.009)

CR=critical ratio; SE=standard error.

postoperative duration on bowel function.

Some studies suggest that defecation function in rectal cancer patients can be restored in approximately 6~12 months after surgery [28]. However, it typically cannot be restored to pre-surgical levels, and QoL is therefore impacted. A qualitative study found that rectal cancer patients who underwent sphincter saving resection did not receive care in relation to bowel problems over time, and did not know whom to consult [18]. This finding suggests that counseling and education are necessary for the patient even after discharge, and that nurses should follow-up on changes in defecation function of the patient for a considerable period of time after discharge.

In this study, social support was identified as a predictor of QoL. Path analysis also showed that there was a significant indirect effect of social support in the relationship between defecation function and QoL. This supports the findings of a previous study that colorectal cancer patients with more social support had better QoL at one year after surgery, when adjusted for age and gender, and that social support had a strong and positive direct effect on QoL [9]. In addition, this finding supports a nationwide study conducted in Korea showing that perceived social support played an important protective role in improving mental health and QoL in cancer patients [29]. Previous studies show that a strong perception of social support increase resilience and QoL, and decreased stress and anxiety in patients with colon cancer [10,30]. These findings highlight the need for healthcare providers to take steps to provide and strengthen timely social support in order help colorectal cancer survivors manage their defecation problems and improve their QoL [10]. With regard to persistent, long-term bowel dysfunction in rectal cancer patients, we therefore suggest that experienced nurses, who are ideally placed to carry out advanced assessments, play an active role in supportive care [6] through counseling and providing information on future progress and health-maintenance behavior.

Furthermore, social support was directly influenced by defecation function in the path analysis of this study. This finding is supported by a systematic review showing that colorectal cancer patients at a more advanced stage were more likely to be anxious and depressed, and have higher person-centered, supportive care needs than those at a less advanced stage [14]. Since perceived social support is an important factor influencing QoL, it is necessary to assess it regularly in patients with defecation problems during follow-up outpatient visits after surgery, and to provide targeted interventions to them. We encourage future studies to collect data from a more diverse sample of colorectal

cancer patients and design a longitudinal study to examine the role of social support in alleviating the influence of defecation function on QoL.

We found that employment status was associated with QoL; in other words, QoL was better among those who were employed. This is consistent with a previous report that job type and employment status contributed to health-related QoL in cancer survivors [31]. This result is also similar to previous studies that defecation function affects employment status, which in turn influences QoL in rectal cancer patients [6, 32]. These findings highlight the importance of employment status for QoL and further research is needed on returning to work after surgery for rectal cancer. Therefore, when patients with colorectal cancer requiring a long recovery period after surgery, and have a job, nurses and healthcare professionals should assess their return to work and job characteristics, and provide counseling to help them practice self-management at work.

The current study has some limitations. First, the study sample was obtained from a single medical center. Although numerous surgeries are performed in this hospital on a large number of rectal cancer patients, the study population cannot be considered to be representative of all rectal cancer survivors in Korea. Another limitation is the possible recall bias due to the use of a self-report questionnaire survey, and the fact that the data are not recent and were collected in 2014.

CONCLUSION

Perceived social support from family and healthcare professionals, defecation function, and employment status were found to be independent predictors of QoL in rectal cancer patients who underwent anal sphincter-saving surgery. Patients with better defecation function had higher perceived social support and higher QoL. In particular, social support partially mediated the relationship between defecation function and QoL. These results suggest that to improve the QoL of rectal cancer patients with defecation dysfunction, it is necessary to improve their perceived social support, and to periodically assess the degree of social support they perceive from family and healthcare providers. In order to do this, it is necessary to for outpatient clinics to provide periodic counseling and education for rectal cancer patients during the postoperative, follow-up period.

CONFLICTS OF INTEREST

The authors declared no conflict of interest.

AUTHORSHIP

Study conception and design - KJR and HSY; Data collection - KJR; Analysis and interpretation of data - KJR and HSY; Drafting and critical revision of the manuscript - KJR and HSY.

ACKNOWLEDGEMENT

This article is a revision of the first author's master's thesis from Hanyang University.

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