Review began 08/08/2022 Review ended 08/12/2022 Published 08/17/2022

© Copyright 2022

Patel et al. This is an open access article distributed under the terms of the Creative Commons Attribution License CC-BY 4.0., which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited.

Chronic Pelvic Pain, Quality of Life, and Patient Satisfaction After Robotic Sacrocolpopexy for Pelvic Organ Prolapse

Nimesh Patel ¹, Priyansh Faldu ², Mohamed Fayed ³, Hannah Milad ⁴, Pradeep Nagaraju ⁵

1. Anesthesiology, Henry Ford Health System, Detroit, USA 2. Medicine, BJ Medical College, Ahmedabad, IND 3. Anesthesiology, Pain Management, and Perioperative Medicine, Henry Ford Health System, Detroit, USA 4. Anesthesia and Critical Care, Wayne State University, Detroit, USA 5. Urology, Beaumont Health, Troy, USA

Corresponding author: Mohamed Fayed, mfayed1@hfhs.org

Abstract

Background and objective

When evaluating repair outcomes in robotic sacrocolpopexy (RSC) for the treatment of pelvic organ prolapse (POP), it has become evident that surgeons usually focus on anatomical improvements and neglect equally important parameters of patient satisfaction and quality of life (QoL). Investigating these factors would aid in achieving a more patient-centered approach to treatment. This study aimed to examine QoL and satisfaction outcomes in women after RSC.

Methods

This study analyzed self-reported patient data regarding RSC for POP performed between October 2009 and February 2017 by fellowship-trained urologists in female pelvic medicine and reconstructive surgery. These patients participated in a survey to assess overall satisfaction and QoL, as well as contributing factors, such as changes in bladder and bowel function, vaginal bulge, and vaginal pain on a 7-point Likert scale (ranging from markedly worse to markedly improved). Data were examined using multivariate regression analysis. Positive treatment response was defined as scores of 6 or 7, whereas negative response was defined as scores of 1 to 5.

Results

The response rate was 41% (156/380), and the median age of the participants was 70 years [interquartile range (IQR): 63, 73]. Of note, 98.7% were Caucasian, with 73% currently in a significant relationship. The median duration since RSC was 2.12 years (IQR: 1.2, 3.7). Overall, 93 (66.9%), patients (23.0%), and 123 patients (88.5%) had a positive treatment response for bladder function, bowel function, and vaginal bulge, respectively. Furthermore, 66% of women had improved QoL, 84% reported improved overall satisfaction, and 91.4% stated that they would recommend RSC to a friend. After controlling for significant covariates, results of a multivariate analysis demonstrated positive treatment response for bladder function [odds ratio (OR): 14.6; p < 0.0001], bowel function (OR: 9.72; p = 0.003), and vaginal bulge (OR: 41.7; p < 0.0001), significantly associated with increased odds of having improved QoL, whereas positive treatment response for vaginal bulge (OR: 26.9; p = 0.023) and recommending RSC to a friend (OR: 175; p = 0.0009) were associated with positive overall satisfaction.

Conclusions

Our findings endorse using RSC surgery for patients with POP based on both QoL improvement and overall post-procedure satisfaction perspective. This study may help encourage surgeons and clinicians to employ a surgical modality that incorporates each patient's unique treatment desires and goals and provide patients with realistic post-procedure goals and expectations regarding treatment.

Categories: Obstetrics/Gynecology, Pain Management, Urology

Keywords: suburethral sling, obstetrics hysterectomy, clinical question research, bowel function, bladder function, patient's satisfaction, quality-of-life, pelvic organ prolapse (pop), chronic pelvic pain syndrome, pelvic pain

Introduction

Pelvic organ prolapse (POP) occurs when the uterus or vagina descends from its normal anatomical position, and it is associated with a lifetime risk of up to 12% for women by the age of 80 years [1-4]. Almost one-third of patients with POP complain about their quality of life (QoL) worsening due to the condition [5]. The POP incidence rate is expected to increase in the next 40 years, with as many as five million women estimated to be affected by 2050, with a reoperation rate of 30% [1,6]. Every year, approximately 200,000 procedures are performed for POP, which is calculated to increase by 46% by 2050 [7-9]. Age, parity, obesity, pelvic floor injury, connective tissue disorders, hysterectomy, estrogen deficiency, intestinal pathologies, chronic

pulmonary disease, and genetics have been previously shown to be risk factors for POP [2,10-15]. The treatment modality to repair POP can be either abdominal or vaginal surgery, with the former approach further sub-classified into open, laparoscopic, and robotic. Surgeons usually choose a treatment modality depending on their comfort level and the patient's comorbidities, age, type of reconstructive procedure, POP extent, the extent of obliterative procedures, and potential complications [16,17].

Since its introduction in 2004, robotic sacrocolpopexy (RSC) has been used to treat POP. Over the years, it has become a widespread and popular treatment option for POP [18]. Compared to other modalities, RSC is less invasive and leads to less perioperative blood loss, lower postoperative pain, quicker return to a normal lifestyle, and reduced complications [19]. Past studies of RSC, which have focused on objective outcomes based on POP-Q stage 1 or less, estimate current anatomical cure rates to be 98.6% [17,20]. To date, research on RSC POP repair has primarily focused on anatomical outcomes; however, little research exists regarding the impact of RSC on functional outcomes [21]. Existing research has focused less on patient satisfaction and QoL and more on quantifiable measures, such as pad weight for incontinence and pelvic organ prolapse quantification (POP-Q) measurements for prolapse. Of late, there has been a greater interest in subjective outcomes due to the emergence of a more patient-centered approach to medicine. Conditions such as POP can limit a patient's interpersonal relationships socially, physically, and psychologically [22]. However, the subjective success of the treatment is not clearly defined. Several studies have used validated questionnaires, such as Pelvic Floor Distress Inventory (PFDI), the Pelvic Floor Impact Questionnaire (PFIQ), Pelvic Organ Prolapse/Urinary Incontinence Sexual Function Questionnaire (PISQ), and Urinary Distress Inventory with a view to quantify patient's symptoms and evaluate subjective success.

However, these validated questionnaires are also problematic in many ways. For instance, many questionnaires are too lengthy for patients to complete, and shorter versions often only identify one symptomatic domain. Clinicians may not accurately understand the patient's overall QoL without additional surveys focusing on various potential POP-related symptoms. Furthermore, the reliability of survey responses may become questionable when the patient population drastically changes or an untested surgical modality is used compared to the initially studied population [15].

This study aims to analyze women's overall satisfaction and QoL following RSC surgery to treat POP. In addition, the influence of symptoms on patients' overall QoL and post-surgery satisfaction will also be assessed.

This article was previously presented as a meeting abstract at the Neurology and Urodynamics Conference in February 2018 [23].

Materials And Methods

This retrospective questionnaire-based study was conducted in the Urology Department of Beaumont Hospital in association with the Michigan Institute of Urology in Royal Oak, Michigan. Institutional Review Board (IRB) approval through Beaumont Hospital was obtained before contacting patients who had undergone RSC treatment for POP (IRB approval number: 2016374). Patients with POP underwent RSC via the standard approach involving the Da Vinci surgical system, performed by fellowship-trained urologists in female pelvic medicine and reconstructive surgery. Concomitant robotic hysterectomy or surgical repair was performed when appropriate. All patients met the criteria for symptomatic vaginal bulge necessitating

Currently living patients who had been 18 years or older at the time of surgery and who underwent RSC between October 2009 and February 2017 were mailed an investigator-created questionnaire to assess self-reported improvements in QoL and overall satisfaction after RSC. A 7-point Likert Scale measured the degree of improvement in patients' subjective health status after undergoing RSC, including bladder function, bowel function, vaginal bulge, pelvic/vaginal pain, QoL, and overall satisfaction (Table 1).

7-Point Likert Scale	Treatment Response Categories	
1 = Markedly Worse		
2 = Moderately Worse		
3 = Mildly Worse	Non-Positive Treatment Response	
4 = Same		
5 = Slightly Improved		
6 = Moderately Improved	Desitive Teacher and Description	
7 = Markedly Improved	Positive Treatment Response	

TABLE 1: 7-Point Likert Scale Questionnaire

Patients were categorized as having a positive treatment response if they indicated that RSC moderately improved or markedly improved their subjective health status. In addition to assessing improvements in subjective health status, the questionnaire further asked patients about demographic information, subsequent surgical history, and whether the patient would recommend RSC to a friend.

Data are presented as medians and interquartile ranges (IQR) for continuous variables, while frequencies and percentages show categorical variables. Odds ratios (OR) with corresponding 95% confidence intervals and p-values were generated using univariate Firth logistic regressions to find significant predictors of positive treatment response in terms of QoL and overall satisfaction. Firth logistic regression was used instead of standard logistic regression due to a high prevalence of patients with positive treatment responses for the outcomes of this study. A p-value <0.05 was considered statistically significant. Data analysis was performed using SAS 9.4 (SAS Institute Inc., Cary, NC). The full questionnaire is available from the authors upon request.

Results

Of the 380 patients who were mailed the questionnaire, 156 responded and returned the completed questionnaire (response rate: 41%). Of the 156 returned questionnaires, 17 were excluded from the final data analysis due to incomplete data. The final sample size was 139 patients. The median age of the respondents was 70 years (IQR: 63, 73), while the median duration since RSC was 2.1 years (IQR: 1.2 years, 3.7 years). In addition, 16 patients (11.5%) had undergone at least one related surgery between RSC and responding to the survey (Table 2).

Surgeries	Number of Patients	
Hysterectomy	4	
Rectocele	3	
Enterocele	1	
Bowel Obstruction	1	
Urethral Sling	1	
Mesh Excision	3	
Other	3	
Total	16	

TABLE 2: Surgeries post-RSC

RSC: robotic sacrocolpopexy

One patient had an entangled bowel following RSC, requiring emergency surgery and ICU admission for nine days. Overall, 93 (66.9%), 32 (23.0%), and 123 patients (88.5%) had a positive treatment response for bladder

function, bowel function, and vaginal bulge, respectively. Of the 57 patients who self-reported pre-RSC pelvic/vaginal pain, 35 patients (61.4%) had a positive treatment response post-procedure. Most patients (91.4%) said they would recommend RSC to a friend. Self-reported patient demographics were stratified according to positive treatment response and non-positive treatment response for QoL and overall satisfaction (Table 3).

	Overall Cohort (n = 139)	Quality of Life Treatment Response				Overall Satisfaction Treatment Response			
		Positive Response (n = 95)	Non-Positive Response (n = 44)	OR (95% CI)	P- value	Positive Response (n = 116)	Non-Positive Response (n = 23)	OR (95% CI)	P- value
Age (Years)									
Median (IQR)	70.0 (63.0, 73.0)	70.0 (65.0, 74.0)	69.0 (61.5, 73.0)	1.02 (0.97, 1.06)	0.4658	70.0 (65.0, 74.0)	69.0 (59.0, 71.0)	1.04 (0.99, 1.09)	0.164
ears From Surgery to Survey									
Median (IQR)	2.1 (1.2, 3.7)	1.9 (1.2, 3.7)	2.8 (1.3, 3.9)	0.84 (0.68, 1.05)	0.1190	2.1 (1.2, 3.5)	2.0 (0.7, 4.6)	0.89 (0.68, 1.15)	0.357
Bladder Function									
Positive Treatment Response	93 (66.9%)	81 (85.3%)	12 (27.3%)	14.6 (6.15, 34.8)	< 0.0001	91 (78.5%)	2 (8.7%)	30.8 (7.69, 124)	< 0.000
Non-Positive Treatment Response	46 (33.1%)	14 (14.7%)	32 (72.7%)	Reference Group		25 (22.5%)	21 (91.3%)	Reference Group	
Bowel Function									
Positive Treatment Response	32 (23.0%)	30 (31.6%)	2 (4.6%)	7.92 (2.02, 31.0)	0.0030	32 (27.6%)	0 (0.0%)	18.1 (1.02, 320)	0.048
Non-Positive Treatment Response	107 (77.0%)	65 (68.4%)	42 (95.4%)	Reference Group		84 (72.4%)	23 (100.0%)	Reference Group	
/aginal Bulge									
Positive Treatment Response	123 (88.5%)	95 (100.0%)	28 (63.6%)	111 (3.90, 999)	0.0016	113 (97.4%)	10 (43.5%)	41.7 (10.6, 164)	< 0.000
Non-Positive Treatment Response	16 (11.5%)	0 (0.0%)	16 (36.4%)	Reference Group		3 (2.6%)	13 (56.5%)	Reference Group	
Pelvic/Vaginal Pain (n = 57)									
Positive Treatment Response	35 (61.4%)	29 (87.9%)	6 (25.0%)	18.7 (4.79, 72.7)	< 0.0001	33 (76.7%)	2 (14.3%)	16.0 (3.39, 75.1)	0.000
Non-Positive Treatment Response	22 (38.6%)	4 (12.1%)	18 (75.0%)	Reference Group		10 (23.3%)	12 (85.7%)	Reference Group	
Race of Patient									
White	138 (99.3%)	94 (98.9%)	44 (100.0%)	1.45 (0.07, 140)	0.8731	115 (99.1%)	23 (100.0%)	0.59 (0.01, 54.1)	0.820
Black	1 (0.7%)	1 (1.1%)	0 (0.0%)	Reference Group		1 (0.9%)	0 (0.0%)	Reference Group	
Ethnicity of Patient									

Hispanic/Latino	3 (2.2%)	3 (3.2%)	0 (0.0%)	(0.11, 105)	0.4893	3 (2.6%)	0 (0.0%)	(0.05, 45.8)	0.8327
Non-Hispanic/Latino	136 (97.8%)	92 (96.8%)	44 (100.0%)	Reference Group		113 (97.4%)	23 (100.0%)	Reference Group	
Current Relationship Status									
In a Significant Relationship but Not Living Together	3 (2.1%)	3 (3.2%)	0 (0.0%)	3.80 (0.12, 119)	0.4477	3 (2.6%)	0 (0.0%)	1.42 (0.04, 45.3)	0.8432
Not in a Significant Relationship	39 (28.1%)	29 (30.5%)	10 (22.7%)	1.53 (0.67, 3.49)	0.3155	32 (27.6%)	7 (30.4%)	0.88 (0.34, 2.30)	0.7901
Living With Spouse/Partner	97 (69.8%)	63 (66.3%)	34 (77.3%)	Reference Group		81 (69.8%)	16 (69.6%)	Reference Group	
Education Level									
High School or Technical School Graduate	47 (33.8%)	33 (34.7%)	14 (31.8%)	2.31 (0.14, 39.6)	0.5632	39 (33.6%)	8 (34.8%)	4.65 (0.26, 82.1)	0.2941
Some College	47 (33.8%)	32 (33.7%)	15 (34.1%)	2.10 (0.12, 35.9)	0.6089	38 (32.8%)	9 (39.0%)	4.06 (0.23, 71.0)	0.3379
College Graduate	22 (15.8%)	17 (17.9%)	5 (11.4%)	3.18 (0.17, 60.2)	0.4401	21 (18.1%)	1 (4.4%)	14.3 (0.56, 369)	0.1080
Graduate or Professional School	21 (15.1%)	12 (12.6%)	9 (20.4%)	1.32 (0.07, 24.0)	0.8527	17 (14.7%)	4 (17.4%)	3.89 (0.20, 75.7)	0.3695
Less than High School	2 (1.5%)	1 (1.1%)	1 (2.3%)	Reference Group		1 (0.8%)	1 (4.4%)	Reference Group	
Current Employment Status									
Employed Full Time	23 (16.6%)	17 (17.9%)	6 (13.6%)	1.18 (1.16, 1.21)	< 0.0001	17 (14.7%)	6 (26.1%)	0.47 (0.16, 1.42)	0.1796
Employed Part Time	18 (13.0%)	11 (11.6%)	7 (15.9%)	0.67 (0.66, 0.69)	< 0.0001	15 (12.9%)	3 (13.0%)	0.77 (0.20, 2.95)	0.7075
Home Maker	12 (8.6%)	9 (9.5%)		1.19	<	10 (8.6%)	2 (8.7%)	0.73	0.0070
		0 (0.070)	3 (6.8%)	(1.16, 1.23)	0.0001			(0.16, 3.49)	0.6976
Unemployed	1 (0.7%)	0 (0.0%)	1 (2.3%)		0.0001	1 (0.9%)	0 (0.0%)		0.7350
Unemployed Disabled	1 (0.7%)			1.23) 0.01 (0.01,		1 (0.9%)	0 (0.0%)	3.49) 0.47 (0.01,	
		0 (0.0%)	1 (2.3%)	1.23) 0.01 (0.01, 999) 0.09 (0.08,	0.3140			3.49) 0.47 (0.01, 39.3) 0.87 (0.02,	0.7350
Disabled	2 (1.4%)	0 (0.0%)	1 (2.3%) 2 (4.6%)	1.23) 0.01 (0.01, 999) 0.09 (0.08, 0.10) Reference	0.3140	2 (1.7%)	0 (0.0%)	3.49) 0.47 (0.01, 39.3) 0.87 (0.02, 37.8) Reference	0.7350
Disabled Retired	2 (1.4%)	0 (0.0%)	1 (2.3%) 2 (4.6%)	1.23) 0.01 (0.01, 999) 0.09 (0.08, 0.10) Reference	0.3140	2 (1.7%)	0 (0.0%)	3.49) 0.47 (0.01, 39.3) 0.87 (0.02, 37.8) Reference	0.7350
Disabled Retired Recommend to Friend?	2 (1.4%) 83 (59.7%)	0 (0.0%) 0 (0.0%) 58 (61.0%)	1 (2.3%) 2 (4.6%) 25 (56.8%)	1.23) 0.01 (0.01, 999) 0.09 (0.08, 0.10) Reference Group 51.5 (2.51,	0.3140 < 0.0001	2 (1.7%) 71 (61.2%)	0 (0.0%)	3.49) 0.47 (0.01, 39.3) 0.87 (0.02, 37.8) Reference Group	0.7350

				999)				510)	
No	9 (6.5%)	0 (0.0%)	9 (20.5%)	Reference Group		1 (0.9%)	2 (8.7%)	Reference Group	
Other Subsequent Surgeries									
Yes	16 (11.5%)	10 (10.5%)	6 (13.6%)	0.73 (0.25, 2.14)	0.5627	14 (12.1%)	2 (8.7%)	1.22 (0.28, 5.24)	0.7925
No	123 (88.5%)	85 (89.5%)	38 (86.4%)	Reference Group		102 (87.9%)	21 (91.3%)	Reference Group	

TABLE 3: Study Variables Stratified by Quality of Life and Overall Patient Satisfaction

IQR: interquartile range

Among patients with positive treatment responses for QoL, 85.3%, 31.6%, and 100.0% of patients also indicated positive treatment responses for bladder function, bowel function, and vaginal bulge, respectively. Univariate logistic regression demonstrated that positive treatment response for bladder function (OR: 14.6; p = < 0.0001), bowel function (OR: 7.92; p = 0.0030), and vaginal bulge (OR: 111; p = 0.0160) were all significantly associated with increased odds of positive treatment response for QoL (Figure 1).

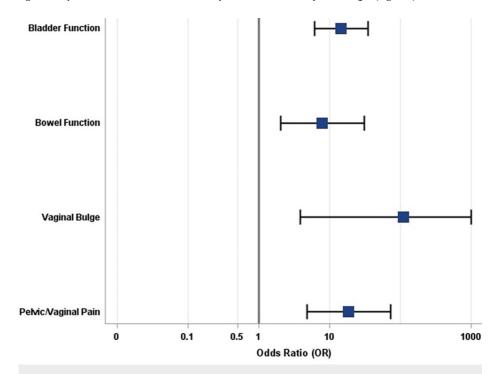


FIGURE 1: Forest Plot for Positive Treatment Response for Quality of Life

Among patients with pelvic/vaginal pain, 87.9% had positive treatment responses for QoL, and this association was also statistically significant (OR: 18.7; p = < 0.0001) (Figure 2).

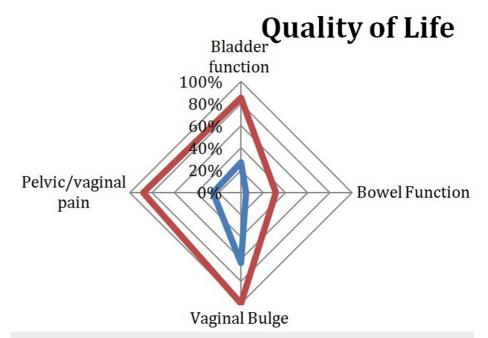


FIGURE 2: Positive Treatment Response and Non-Positive Treatment Response for Quality of Life

Red line: positive treatment response; blue line: non-positive treatment response

Similarly, 78.5%, 27.6%, and 97.4% of patients with positive treatment responses to bladder function, bowel function, and vaginal bulge, respectively, also had positive treatment responses in terms of overall satisfaction. Positive treatment response for bladder function (OR: 30.8; p = < 0.0001), bowel function (OR: 18.1; p = 0.0483), and vaginal bulge (OR: 41.7; p = 0.0005) were all significantly associated with increased odds of having positive treatment responses in terms of overall satisfaction (Figure 3).

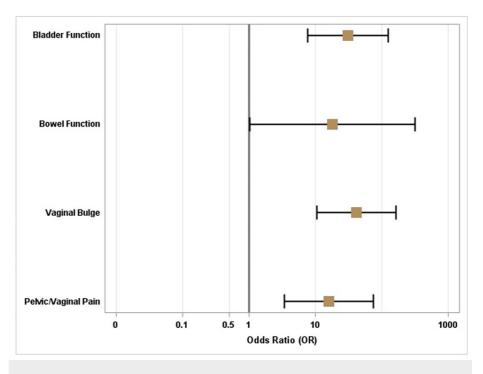


FIGURE 3: Forest Plot for Positive Treatment Response for Overall Satisfaction

For patients who self-reported pelvic/vaginal pain prior to RSC, positive treatment response for pelvic/vaginal pain was also significantly associated with positive treatment response regarding overall satisfaction (OR: 16.0; p = 0.0005), with 76.7% of patients having positive treatment responses for both overall satisfaction and pelvic/vaginal pain (Figure 4).

Overall Satisfaction

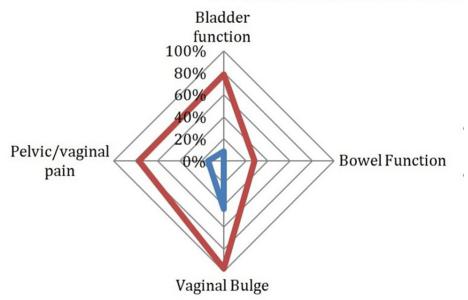


FIGURE 4: Positive Treatment Response and Non-Positive Treatment Response for Overall Satisfaction

Red line: positive treatment response; blue line: non-positive treatment response

Current relationship status, education level, and other subsequent surgeries were not associated with positive treatment response for either QoL or overall satisfaction (all p > 0.05). Current employment status was significantly associated with positive treatment response for QoL but not with positive treatment response for overall satisfaction. Patients recommending RSC to friends had significantly increased odds of having positive treatment responses for both QoL and overall satisfaction (p < 0.05).

Discussion

RSC has come to be routinely used for POP; however, subjective measures of surgical satisfaction, including QoL and overall satisfaction, have yet to be evaluated as a primary aim. Our questionnaire was developed to ascertain patients' subjective symptoms after RSC and its influence on their QoL and overall satisfaction. POP affects women's QoL, and it is thus essential to measure QoL when considering a treatment option [24]. QoL was defined as a patient's overall wellbeing with or without distress from their POP, whereas overall satisfaction pertains to how content patients were with RSC surgery when completing their questionnaire.

One important finding was that positive treatment response for bowel function was significantly associated with positive treatment response for QoL and overall satisfaction. Our results show that positive treatment response for bowel function continues for a median duration of 2.1 years, whereas previously published literature had a median one-year follow-up after RSC surgery [25-27]. In addition, our results show that only 23% of patients reported positive treatment responses for bowel function, whereas the majority had no changes in bowel function. This suggests that RSC surgery dramatically affects the QoL of a minority of patients, who are symptomatic of bowel dysfunction caused by their POP.

Positive treatment response for vaginal bulge significantly influences QoL and overall satisfaction. This finding supports a previous study, which showed that subjective cure, the absence of bulge symptoms, occurred in 92.1% of patients, compared to 88.5% [28]. The improvement in vaginal bulge symptoms is essential to women's QoL and overall satisfaction, as nearly every patient with positive treatment response in our study also had a positive treatment response for QoL. Likewise, the patient's pelvic/vaginal pain significantly affects her QoL and overall surgical satisfaction. Previous publications have elucidated that RSC is associated with greater inter- and postoperative pain compared to other surgical modalities [29,30]. However, these findings should be interpreted with caution, since our study indicates that only 38.6%

(22/57) of patients did not have a positive treatment response after their RSC surgery, whereas 83% (29/35) with positive treatment response for pelvic pain had improved QoL.

It is well supported within the literature that urinary retention, urinary tract infection, and bladder injury are frequent complications of RSC surgery [31]. Our study shows that these urinary adverse effects should be treated immediately, as bladder function is significantly associated with improving patients' QoL and overall satisfaction. This is reinforced further by the difference between positive and non-positive treatment responses of 58% and 70% for QoL and overall satisfaction, respectively. This suggests that patients can immediately recognize any bladder problems that negatively affect their QoL or, conversely, improvements in bladder function post-RSC surgery increases QoL and RSC satisfaction.

This study has several limitations that may affect its results and conclusions. Firstly, the investigator-created questionnaire was not validated; however, the investigators could not locate a previously validated survey that adequately examines improvements in subjective health status following this unique procedure. Moreover, the response rate of 41% was low, and we did not draw any conclusions in terms of a comparison between survey responders and non-responders. There is a possibility that patients with favorable outcomes were more likely to complete and return the questionnaire. Finally, due to the high prevalence of positive treatment responses for QoL and overall satisfaction, multivariate logistic regression results were subject to bias from over-fitting and hence were excluded from this analysis.

Conclusions

Based on our findings, this study endorses using RSC surgery specifically concerning patient QoL improvement and overall satisfaction. An improvement in bladder and bowel function, vaginal bulge, and pelvic pain underscores its contribution to improving patients' QoL. This study may help surgeons to employ a surgical modality that incorporates the patient's treatment desires. The results of this study could also help surgeons better understand the factors contributing to satisfaction and QoL to help patients set realistic goals for treatment.

Additional Information

Disclosures

Human subjects: Consent was obtained or waived by all participants in this study. Institutional Review Board, Beaumont Hospital issued approval 2016-374. **Animal subjects:** All authors have confirmed that this study did not involve animal subjects or tissue. **Conflicts of interest:** In compliance with the ICMJE uniform disclosure form, all authors declare the following: **Payment/services info:** All authors have declared that no financial support was received from any organization for the submitted work. **Financial relationships:** All authors have declared that they have no financial relationships at present or within the previous three years with any organizations that might have an interest in the submitted work. **Other relationships:** All authors have declared that there are no other relationships or activities that could appear to have influenced the submitted work.

References

- Al-Badr A: Quality of life questionnaires for the assessment of pelvic organ prolapse: use in clinical practice. Low Urin Tract Symptoms. 2013, 5:121-8. 10.1111/luts.12006
- Alas AN, Anger JT: Management of apical pelvic organ prolapse. Curr Urol Rep. 2015, 16:33. 10.1007/s11934-015-0498-6
- Wu JM, Matthews CA, Conover MM, Pate V, Jonsson Funk M: Lifetime risk of stress urinary incontinence or pelvic organ prolapse surgery. Obstet Gynecol. 2014, 123:1201-6. 10.1097/AOG.0000000000000286
- Buca DI, Leombroni M, Falò E, et al.: A 2-year evaluation of quality of life outcomes of patients with pelvic organ prolapse treated with an elevate prolapse repair system. Female Pelvic Med Reconstr Surg. 2016, 22:410-4. 10.1097/SPV.0000000000000310
- Nygaard IE, McCreery R, Brubaker L, Connolly A, Cundiff G, Weber AM, Zyczynski H: Abdominal sacrocolpopexy: a comprehensive review. Obstet Gynecol. 2004, 104:805-23. 10.1097/01.AOG.0000139514.90897.07
- Wu JM, Hundley AF, Fulton RG, Myers ER: Forecasting the prevalence of pelvic floor disorders in U.S. Women: 2010 to 2050. Obstet Gynecol. 2009, 114:1278-83. 10.1097/AOG.0b013e3181c2ce96
- Jones KA, Shepherd JP, Oliphant SS, Wang L, Bunker CH, Lowder JL: Trends in inpatient prolapse procedures in the United States, 1979-2006. Am J Obstet Gynecol. 2010, 202:501.e1-7. 10.1016/j.ajog.2010.01.017
- Brown JS, Waetjen LE, Subak LL, Thom DH, Van den Eeden S, Vittinghoff E: Pelvic organ prolapse surgery in the United States, 1997. Am J Obstet Gynecol. 2002, 186:712-6. 10.1067/mob.2002.121897
- Dannecker C, Anthuber C: The effects of childbirth on the pelvic-floor . J Perinat Med. 2000, 28:175-84.
 10.1515/[PM.2000.025
- Fritel X, Varnoux N, Zins M, Breart G, Ringa V: Symptomatic pelvic organ prolapse at midlife, quality of life, and risk factors. Obstet Gynecol. 2009, 113:609-16. 10.1097/AOG.0b013e3181985312
- 12. Whitcomb EL, Lukacz ES, Lawrence JM, Nager CW, Luber KM: Prevalence and degree of bother from pelvic

- floor disorders in obese women. Int Urogynecol J Pelvic Floor Dysfunct. 2009, 20:289-94. 10.1007/s00192-008-0765-x
- Swift S, Woodman P, O'Boyle A, et al.: Pelvic Organ Support Study (POSST): the distribution, clinical definition, and epidemiologic condition of pelvic organ support defects. Am J Obstet Gynecol. 2005, 192:795-806. 10.1016/j.ajog.2004.10.602
- Kudish BI, Iglesia CB, Sokol RJ, et al.: Effect of weight change on natural history of pelvic organ prolapse .
 Obstet Gynecol. 2009, 113:81-8. 10.1097/AOG.0b013e318190a0dd
- Digesu GA, Khullar V, Cardozo L, Robinson D, Salvatore S: P-QOL: a validated questionnaire to assess the symptoms and quality of life of women with urogenital prolapse. Int Urogynecol J Pelvic Floor Dysfunct. 2005, 16:176-81. 10.1007/s00192-004-1225-x
- Gilleran JP, Johnson M, Hundley A: Robotic-assisted laparoscopic mesh sacrocolpopexy. Ther Adv Urol. 2010, 2:195-208. 10.1177/1756287210379120
- Daneshgari F, Kefer JC, Moore C, Kaouk J: Robotic abdominal sacrocolpopexy/sacrouteropexy repair of advanced female pelvic organ prolaspe (POP): utilizing POP-quantification-based staging and outcomes. BJU Int. 2007, 100:875-9. 10.1111/j.1464-410X.2007.07109.x
- Sajadi KP, Goldman HB: Robotic pelvic organ prolapse surgery. Nat Rev Urol. 2015, 12:216-24.
 10.1038/nrurol.2015.51
- Elliott DS, Siddiqui SA, Chow GK: Assessment of the durability of robot-assisted laparoscopic sacrocolpopexy for treatment of vaginal vault prolapse. J Robot Surg. 2007, 1:163-8. 10.1007/s11701-007-0028-8
- Lee RK, Mottrie A, Payne CK, Waltregny D: A review of the current status of laparoscopic and robot-assisted sacrocolpopexy for pelvic organ prolapse. Eur Urol. 2014, 65:1128-37. 10.1016/j.eururo.2013.12.064
- Rogers RG: To mesh or not to mesh: current debates in prolapse repair fueled by the U.S. Food and Drug Administration Safety Notification. Obstet Gynecol. 2011, 118:771-3. 10.1097/AOG.0b013e318230b1f4
- Digesu GA, Santamato S, Khullar V, et al.: Validation of an Italian version of the prolapse quality of life questionnaire. Eur J Obstet Gynecol Reprod Biol. 2003, 106:184-92. 10.1016/s0301-2115(02)00229-4
- Patel N, Killinger K, Thapar K, Karabon P, Nagaraju P: Patient satisfaction and quality of life after robotic sacrocolpopexy for pelvic organ prolapse. Neurourol Urodyn. 2018, 37:S679-80.
- Mourik SL, Martens JE, Aktas M: Uterine preservation in pelvic organ prolapse using robot assisted laparoscopic sacrohysteropexy: quality of life and technique. Eur J Obstet Gynecol Reprod Biol. 2012, 165:122-7. 10.1016/j.ejogrb.2012.07.025
- Lewis C, Salamon C, Priestley JL, Gurshumov E, Culligan P: Prospective cohort study of bowel function after robotic sacrocolpopexy. Female Pelvic Med Reconstr Surg. 2014, 20:87-9. 10.1097/SPV.000000000000000049
- Perez T, Crochet P, Descargues G, et al.: Laparoscopic sacrocolpopexy for management of pelvic organ prolapse enhances quality of life at one year: a prospective observational study. J Minim Invasive Gynecol. 2011, 18:747-54. 10.1016/j.jmig.2011.08.002
- Geller EJ, Parnell BA, Dunivan GC: Pelvic floor function before and after robotic sacrocolpopexy: one-year outcomes. J Minim Invasive Gynecol. 2011, 18:322-7. 10.1016/j.jmig.2011.01.008
- Barber MD, Brubaker L, Nygaard I, Wheeler TL 2nd, Schaffer J, Chen Z, Spino C: Defining success after surgery for pelvic organ prolapse. Obstet Gynecol. 2009. 114:600-9. 10.1097/AOG.0b013e3181b2b1ae
- Callewaert G, Bosteels J, Housmans S, et al.: Laparoscopic versus robotic-assisted sacrocolpopexy for pelvic organ prolapse: a systematic review. Gynecol Surg. 2016, 13:115-23. 10.1007/s10397-016-0930-z
- De Gouveia De Sa M, Claydon LS, Whitlow B, Dolcet Artahona MA: Robotic versus laparoscopic sacrocolpopexy for treatment of prolapse of the apical segment of the vagina: a systematic review and meta-analysis. Int Urogynecol J. 2016, 27:355-66. 10.1007/s00192-015-2763-0
- Danforth TL, Aron M, Ginsberg DA: Robotic sacrocolpopexy. Indian J Urol. 2014, 30:318-25. 10.4103/0970-1591.128502