# RESEARCH



# How does surgery influence female sexuality in patients with endometriosis compared to those with other benign gynecological conditions?

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### Abstract

**Background** Endometriosis is a chronic, estrogen-dependent, benign condition, affecting 10–15% of women of reproductive age. It is associated with a prevalence of sexual dysfunction that is nearly twice as high as that seen in women with other benign gynecological conditions. Our study aimed to assess the effect of surgical intervention on sexual function, as measured by the FSFI (Female Sexual Function Index) score, in women with endometriosis compared to those with other benign gynecological conditions, both before and after surgery.

**Methods** A comparative analysis was conducted at the Medical University of Vienna from 2015 to 2020. The study included patients suspected of having endometriosis, fibroids, adnexal cysts, and/or infertility. Based on findings during surgery, patients were categorized into two groups: those with endometriosis (n = 64) and control patients (n = 38). All participants completed the FSFI questionnaire before surgery and again 8 to 18 weeks after the operation.

**Results** No significant differences were observed in the preoperative FSFI scores between the endometriosis patients and the control group. Similarly, no significant differences were found between the two groups in postoperative scores. However, in women diagnosed with endometriosis, surgical removal of endometriotic lesions significantly increased their full-scale FSFI score, and resulted in a significant improvement in the areas "desire" and "satisfaction". Improvements were noted in all other areas as well, though they were not statistically significant.

**Conclusions** Our research indicates that the surgical removal of endometriotic lesions can lead to an improvement in sexual function, as measured by the FSFI, within 8 to 18 weeks post-surgery. This improvement was not observed in the control group, which underwent surgery for other benign gynecological issues.

Keywords Sexual function, FSFI, Endometriosis, Surgical excision

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## Background

Endometriosis is a chronic, benign, estrogen-dependent disease that affects 10–15% of women of reproductive age, resulting in a number of about 176 million women worldwide [1]. It is characterized by the implantation of endometrium-like tissue outside the uterus, which can cause a wide range of symptoms [2]. These include severe dysmenorrhea, dyspareunia, dyschezia, dysuria, chronic

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pelvic pain, infertility, and/or an impairment of the women's sexual function. These symptoms can have a negative impact on the social, mental, and physical well-being of affected women which can severely reduce their quality of life [1, 3–5]. It has been described that in women suffering from endometriosis, the prevalence of sexual dysfunction is almost double compared to women with other benign gynecologic diseases (i.e., 61% vs. 35%) [6–8]. Furthermore, endometriosis is one of the leading causes of deep dyspareunia. The risk of suffering from dyspareunia was described nine times higher in patients with endometriosis than in women without endometriosis [9]. Particularly deep-infiltrating endometriosis (DIE) seems to have a negative impact on sexual function including desire, the ability to reach orgasm, the frequency of sexual intercourse, and the global sexual satisfaction [10, 11].

In addition, hormonal medications such as combined contraceptive pills, progesterone-only pills, GnRH (gonadotropin-releasing hormone) analogues or hormonal intrauterine devices (IUDs) which are used as a first- and second-line therapeutic options for endometriosis-associated symptoms [12] can also affect female sexuality, potentially limiting desire, sexual arousal, lubrication, and/or the ability to reach an orgasm [13]. Furthermore, possible side effects of these medications include anxiety disorders and mood swings, which can consequently also lead to an impairment of the patient's sex life [14].

While some studies have shown that surgical excision of endometriotic lesions can result in a significant improvement of dyspareunia [15, 16], most trials that assessed the sexual function in women with endometriosis before and after surgery [17, 18] were limited by their small sample size and retrospective design. To our knowledge, there are no studies comparing pre- and postoperative sexual function in patients with endometriosis compared to controls.

In this prospective study, we compared the impact of surgical therapy on the sexual function measured by the Female Sexual Function Index (FSFI) score, in women suffering from endometriosis compared to controls. Furthermore, we aimed to compare pre- and postoperative sexual functions within each group.

#### Methods

Our study was conducted at the tertiary referral Endometriosis Centre of the Medical University of Vienna, certified by EuroEndoCert [19]. The study protocol was approved by the local Ethics Committee (EC no. 1907/2019). All patients gave their written and verbal informed consent at the time of inclusion. All patients had undergone laparoscopic surgery at our department between 2015 and 2020. Indications for surgery included suspected endometriosis, fibroids, adnexal cysts, or infertility. In our collective, patients with adnexal cysts had undergone cystectomy, patients with fibroids had undergone myomectomy, and patients suffering from infertility had undergone a diagnostic laparoscopy and in cases of endometriosis findings, excision of endometriotic lesions. These patients were otherwise healthy women without internal or autoimmune diseases. Depending on the intraoperative findings, patients were either classified as endometriosis patients (n, number = 64) or controls (n = 38). Endometriosis was staged intraoperatively, using the revised rASRM score (revised American Society of Reproductive Medicine Score) [20, 21] as well as the ENZIAN classification [20-22]. None of the patients included in this study suffered from intraand/or postoperative complications that might have changed their postoperative course.

Exclusion criteria were defined as history of breast cancer in the past 10 years or other malignant diseases in the past 5 years, a history of infectious diseases such as HIV, hepatitis, tuberculosis, or systemic autoimmune diseases. All patients included in our study had regular menstrual cycles and had not taken any hormonal medication in the 3 months preceding study inclusion nor in the 5 months following the surgical intervention.

Preoperatively, an exact patient history was taken regarding demographic and clinical data, age, body mass index (BMI), gravidity, parity, menarche as well as the pain intensity of dysmenorrhea, dyspareunia, and dysuria, measured using the visual analogue scale (VAS) and the extent of the influence of dyspareunia on the patient's sex life. Patients' sexual function was assessed using the German-validated version of the FSFI questionnaire [23], which was filled in preoperatively, as well as 8 to 18 weeks postoperatively. This whole procedure took 25 min per patient. Regarding postoperative advice, depending on the extent of the surgical intervention, patients were recommended to abstain from sexual intercourse for a period of 2 up to 4 weeks after surgery.

While several validated questionnaires have been developed to evaluate the quality of female sexuality [24–27], the most commonly used is the Female Sexual Function Index (FSFI). This 19-item questionnaire records the sexual function based on the subjective perception of affected women [28]. It has already been in use for two decades in women suffering from a wide range of different diseases and enables the comparison of different groups [27]. Reliability and validity were confirmed by Rosen et al. in 2000 [25].

The FSFI questionnaire is split into six domains: desire (two items), arousal (four items), lubrication (four items), orgasm (three items), satisfaction (three items), and pain (three items). Each item in the questionnaire is scored from 0 to 5. The sum of each domain is then multiplied by a factor that takes into account the amount of influence each domain potentially has on the overall score. A total score of  $\leq 26.55$  seems to represent a higher risk of sexual dysfunction [25].

#### **Statistics and power analysis**

Our data is reported via median and interquartile range (numerical variables). Statistical tests were done with the R software package (version 4.2.1) [29]. Data were plotted with the ggplot2 package (version 3.3.6) [30]; power analysis was performed with the package pwr (version 1.3–0) [31].

The difference in parameters between groups was examined via Wilcoxon rank-sum tests, respectively via Wilcoxon signed rank test with continuity correction in the case of paired groups.

Since it is not possible to directly conduct a power analysis for a non-parametric test, it is necessary to employ Pitman asymptotic relative efficiency (ARE) to at least estimate the effect sizes one is able to detect [32]. Since, at worst, the ARE of the Wilcoxon rank-sum test was 0.864 [33] in comparison to the *t*-test, we performed a power analysis for the two-sided unpaired *t*-test under the assumptions of: sample sizes n1=33 (~38\*0.864), n2=55 (~64\*0.864), significance level=0.05, power=0.8. We calculated a detectable effect size of at least ~ 0.62 which corresponds to effects of size medium to large [34].

For our comparisons of the paired groups, an analogue approach revealed that we were able to observe effects of sizes 0.38 respectively 0.72, which corresponds to medium respectively large effects [34]. A *p*-value of < 0.05 was considered as statistically significant.

#### Results

Our study collective comprised 102 premenopausal women (n, number = 102); 64 women had laparoscopically and histologically confirmed endometriosis. Endometriosis was surgically excluded in 38 women who were therefore included in our control group.

Patient characteristics are presented in Table 1.

There were no statistically significant differences between the two groups in median age, BMI, gravidity, parity, and menarche, nor in the pain categories "overall pain score" and "dyspareunia". However, patients suffering from endometriosis had a significantly higher intensity of "dysmenorrhea" than controls (p = 0.0158). There were no significant differences between FSFI full-scale score of controls and patients with endometriosis in preoperatively assessed total FSFI score and in the six domains (Table 2).

In addition, no significant differences between the groups were found 8 to 18 weeks postoperatively (Table 3).

Furthermore, we compared pre- and postoperative sexual function within the groups. In women suffering from endometriosis, surgical excision of endometriotic lesions resulted in a significant improvement of the FSFI full-scale score (23.8 vs. 27.9, p=0.0499) as well as of the domains "desire" (3.0 vs. 3.6, p=0.0008), and "satisfaction" (4.4 vs. 5.2, p=0.0024). Although an improvement was found in all other domains, this showed no statistical significance (Table 4).

Comparing pre- and postsurgical assessment, women included in the control group showed a statistically significant improvement in the domain "desire" (3.0 vs. 3.6, p=0.0254). All other domains as well as the full-scale score showed no statistically significant improvement after surgery. Details are presented in Table 5.

The changes of the full-scale score between the groups and between the two different time points within each group are shown in Fig. 1.

#### Discussion

Endometriosis is often associated with dyspareunia, which can strongly affect the patient's overall sexual function.

Pain in itself is a very complex phenomenon influenced by a multitude of factors. The data regarding endometriosis-associated pain is therefore not consistent throughout the literature. While our findings confirmed results from some previous studies [35-37] regarding dysmenorrhea (VAS 8 in the endometriosis vs. VAS 6 in controls, p=0.0158), one other study showed no difference in intensity of menstrual pain between patients with endometriosis and controls [28]. Regarding dyspareunia however, the systematic review of Shi et al. (2022) described a significant difference in dyspareunia intensity [38], whereas other previous papers [39, 40] and our current study showed no difference between the groups.

These results are mirrored in our data regarding the FSFI: while the domain "pain" was increased in endometriosis patients compared to controls, the difference between the two groups, in contrast to other studies [27, 28, 36, 41], showed no statistical significance.

When examining the effect of surgical therapy on dyspareunia, we found no significant improvement. Another cause for dyspareunia has been found in the pelvic floor muscles. Recently published data show the importance of a pelvic floor examination in patients with dyspareunia in order to evaluate myofascial components, which are known to play a potentially important role in this type of chronic pelvic pain [42]. This examination is however lacking in our and other previously mentioned endometriosis studies [38].

Regarding the full-scale score of FSFI, we noticed a lower preoperative score (median 23.8) in the

#### Table 1 Patient characteristics

Patient characteristics	All ( <i>n</i> = 102)	Control group (n = 38)	Endometriosis group ( <i>n</i> =64)	<i>p</i> -value
General information <sup>a</sup>				
Age (years)	34 (31—40)	35 (31—41.5)	33 (30.75—39.25)	0.2556
BMI (kg/m <sup>2</sup> )	22.54 (20.70—26.85)	21.96 (20.70—26.19)	22.60 (20.71—26.96)	0.9228
Gravidity	0.5 (0—2)	1 (0—2)	0 (0—1)	0.3486
Parity	0 (0—0)	0 (0—0)	0 (0—1)	0.4614
Menarche	13 (11—14)	13 (11.25—14)	12 (11—14)	0.9972
Smoking <sup>b</sup>				0.2348
No	82	33	49	
Yes	20	5	15	
In a relationship <sup>b</sup>				0.1186
No	22	5	17	
Yes	80	33	47	
Completed family planning <sup>b</sup>				0.2852
No	66	22	44	
Yes	36	16	20	
Desire to have children <sup>b</sup>				0.7904
No	64	23	41	
Yes	38	15	23	
Preoperative pain symptoms				
Overall non-menstrual pain score	6.5 (3—9)	6 (3—9)	7 (3.75—9)	0.5860
Dysmenorrhea intensity (VAS)	8 (5—9)	6 (2—8)	8 (5—10)	0.0158
Dyspareunia intensity (VAS)	1 (0—6)	0 (0—5)	3 (0—6)	0.1197
Influence on sex life (VAS)	0 (0—6)	0 (0—4)	0 (0—7)	0.0877
Dysuria intensity (VAS)	0 (0—0)	0 (0—0)	0 (0—0)	0.3442
rASRM score ( <i>n</i> )				
I		NA	13	
II		NA	14	
III		NA	14	
IV		NA	16	
NA		NA	7	
ENZIAN score ( <i>n</i> )				
A 1–3		NA	9	
B 1–3		NA	16	
C 1–3		NA	7	
FA		NA	11	
FB		NA	3	
FI		NA	2	
FO		NA	5	

<sup>a</sup> Wilcoxon rank-sum test with continuity correction

Values are expressed in median and interquartile range. Categories are presented in absolute values

<sup>b</sup> Barnard's unconditional test

endometriosis group compared to controls (median 27.7). Although this difference was not statistically significant, patients suffering from endometriosis had a higher potential to benefit from surgery, confirmed by a statistically significant post-surgical increase of the full-scale score in this group (from 23.8 to 27.9, p < 0.05).

While many gynecological disorders seen in our patient collectives can lead to a reduction in the FSFI score, endometriosis patients seem to benefit more from surgical therapy than our controls. The same can be said for the domain "satisfaction" which improved in both groups, yet significantly only in patients with endometriosis (p = 0.0024).

**Table 2**Preoperative FSFI scores: controls vs. endometriosispatients

Variable	Controls (n = 38)	Endometriosis (n=64)	<i>p</i> -value
Desire	3.0 (2.4–3.6)	3.0 (2.4–3.8)	0.4599
Arousal	3.9 (2.5–5.0)	3.9 (2.0–4.8)	0.6346
Lubrication	5.1 (3.2–5.9)	4.4 (3.3–5.7)	0.5742
Orgasm	4.8 (2.1–5.9)	4.4 (2.8–5.6)	0.8366
Satisfaction	4.8 (3.4–6.0)	4.4 (2.8–5.3)	0.3937
Pain	4.8 (2.5–6.0)	3.8 (1.6–5.7)	0.2234
Full-scale score	27.7 (17.5–30.7)	23.8 (15.4–29.3)	0.3395

Values are expressed in median (interquartile range)

**Table 3** Postoperative FSFI scores: controls vs. endometriosis patients

Variable	Controls (n = 38)	Endometriosis (n=64)	<i>p</i> -value
Desire	3.6 (3.0–4.7)	3.6 (2.4–4.8)	0.5157
Arousal	4.5 (3.1–5.3)	4.5 (3.0–5.4)	0.9889
Lubrication	5.6 (3.7–6.0)	5.4 (3.5–6.0)	0.5860
Orgasm	4.8 (3.3–6.0)	4.6 (2.4–5.6)	0.3810
Satisfaction	5.2 (4.0–6.0)	5.2 (4.0-6.0)	0.8374
Pain	5.6 (2.8–6.0)	5.2 (0.0-6.0)	0.5542
Full-scale score	28.6 (22.0–31.4)	27.9 (18.0–32.4)	0.8059

Values are expressed in median (interquartile range)

**Table 4**FSFI scores of patients with endometriosis: preoperativevs. postoperative values

Variable	Preoperative (n=64)	Postoperative (n=64)	<i>p</i> -value
Desire	3.0 (2.4–3.8)	3.6 (2.4–4.8)	0.0008
Arousal	3.9 (2.0–4.8)	4.5 (3.0-5.4)	0.0542
Lubrication	4.4 (3.3–5.7)	5.4 (3.5–6)	0.2641
Orgasm	4.4 (2.8–5.6)	4.6 (2.4–5.6)	0.7904
Satisfaction	4.4 (2.8–5.3)	5.2 (4.0-6.0)	0.0024
Pain	3.8 (1.6–5.7)	5.2 (0.0-6.0)	0.2880
Full-scale score	23.8 (15.4–29.3)	27.9 (18.0–32.4)	0.0499

Values are expressed in median (interquartile range), Wilcoxon signed rank test with continuity correction. Significant p-values are set in italics

While no difference was found here between the two groups, sexual desire was the only parameter that significantly improved post surgically in both the endometriosis and the control group. However, it has to be underlined that sexual desire can be influenced by a multitude of psychological factors (e.g., performance anxiety, depression) [43], which leads us to question whether this improvement might be due to a mere positive psychological reaction caused by the beneficial effects of surgery in women affected by gynecological diseases, underlining

Variable	Preoperative (n=38)	Postoperative (n=38)	<i>p</i> -value
Desire	3.0 (2.4–3.6)	3.6 (3.0–4.7)	0.0254
Arousal	3.9 (2.5–5.0)	4.5 (3.1–5.3)	0.1990
Lubrication	5.1 (3.25–6.0)	5.6 (3.7–6.0)	0.2271
Orgasm	4.8 (2.1–5.9)	4.8 (3.3–6.0)	0.2635
Satisfaction	4.8 (3.4–6.0)	5.2 (4.0-6.0)	0.2693
Pain	4.8 (2.5–6.0)	5.6 (2.8–6.0)	0.3359
Full-scale score	27.7 (17.5–30.7)	28.6 (22.0-31.4)	0.1944

 Table 5
 FSFI scores of controls: preoperative vs. postoperative

Values are expressed in median (interquartile range), Wilcoxon signed rank test with continuity correction. Significant *p*-values are set in italics

the importance of further studies in this collective with a longer follow-up period.

#### Strength and limitations

values

A strength of our study is the prospective cohort design of well-characterized study groups [39, 44]. While laparoscopy has been replaced as the gold-standard for the diagnosis of DIE, adenomyosis, and endometriomas, it allows the confirmation or exclusion of peritoneal lesions [45] and a more precise classification of the disease. In all endometriosis patients, the disease was intraoperatively scored by a member of our certified Endometriosis Centre according to rASRM and ENZIAN [20-22] classifications and all visible endometriotic lesions were resected. In contrast to other studies [28, 46], our collective comprised all endometriosis subtypes (peritoneal, ovarian, DIE). While this allowed the analysis of the changes of the FSFI in all endometriosis subtypes, representing the whole entity of this disease, the lack of stratification according to endometriosis subtype, in particular to the occurrence of DIE lesions, can be seen as a limitation of our study, as the results do therefore not reflect the possible influence of certain lesions on particular aspects of sexual function, such as dyspareunia. Furthermore, all patients who had used hormonal contraceptives within 3 months prior to the study [28, 41] were not included in our study, hereby eliminating the potential bias of hormonal contraceptives regarding sexual function [14].

In addition, we have not found previous studies that describe the effect of surgery on FSFI changes in patients with endometriosis compared to controls.

We find a key factor of conducting endometriosis studies to be the selection of the control group [17, 18]. In contrast to some other studies that simply chose asymptomatic women, we strived to define a control group in which the presence of endometriosis had been surgically excluded. All our patients however suffered from benign

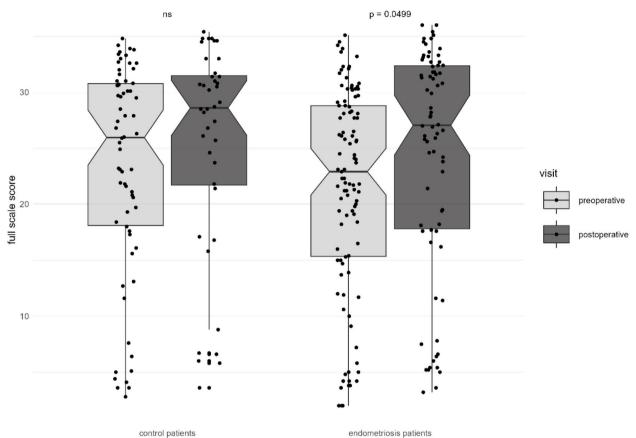


Fig. 1 Changes of the FSFI full-scale score pre- and postoperatively, control patients vs. endometriosis patients

gynecological diseases that affected their pelvic organs and potentially by extension their sexuality. Finding an "ideal control group" is in our opinion a difficulty that needs to be solved through further studies.

Another limitation of our study is the short time of follow-up. Our follow-up time was eight to 18 weeks postoperatively. In order to truly evaluate a long-term effect of surgical excision of endometriotic lesions, further data with a longer follow-up period should follow.

#### Conclusions

This study demonstrates that the surgical removal of endometriotic lesions can lead to significant improvements in sexual function, as measured by the FSFI within 8 to 18 weeks' post-surgery. Unlike surgeries performed for other benign gynecological conditions, the benefits observed in women with endometriosis suggest that impaired sexual function could become an increasingly important factor in the decision-making process for surgical intervention in these patients. This is particularly relevant for those suffering from deep dyspareunia and other forms of sexual dysfunction, which are prevalent in endometriosis.

In a clinical context, these results underscore the importance of addressing sexual health as a key component of patient care in endometriosis management. By highlighting the positive outcomes of surgical treatment on sexual function, this research supports the consideration of surgery not only for alleviating pain but also for improving the overall quality of life for women with endometriosis. This evidence may guide clinicians in recommending surgery earlier in the treatment process for patients experiencing significant sexual dysfunction, thereby potentially altering the current therapeutic landscape. Future studies with larger sample sizes and longer follow-up periods are necessary to further validate these findings and solidify the role of surgical intervention in the management of sexual dysfunction associated with endometriosis.

#### Abbreviations

BMI Body mass index

- DIE Deep-infiltrating endometriosis
- FSFI Female Sexual Function Index
- GnRH Gonadotropin-releasing hormone
- IUD Intrauterine device
- n Number
- rASRM Revised American Society of Reproductive Medicine

VAS Visual analogue scale

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#### Authors' contributions

A.P. and M.G. contributed equally to this study and share first authorship. All authors contributed equally to conception and design of study as well as manuscript finalization. AP, MG, SI, FH, LS, JH, HH, LK, CB, and RW were involved in the conception, design, and interpretation of the results. AP, MG, SI, LS, JH, HH, LK, and CB acquired the data. AP, MG, and RW conducted the analyses and wrote the first draft of the manuscript. HH, LK, and RW were the responsible surgeons. AP, MG, SI, and FH did statistical analysis. All authors read and approved the final manuscript.

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#### Data availability

The data that support the findings of this study are not openly available due to reasons of sensitivity and are available from the corresponding author upon reasonable request.

#### Declarations

#### Ethics approval and consent to participate

The study protocol was approved by the Ethics Committee of the Medical University of Vienna (EC no. 1907/2019). All patients gave their verbal and written informed consent at the time of study inclusion.

#### **Consent for publication**

Not applicable.

#### **Competing interests**

The authors declare no competing interests.

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