

Original Article

Short-term advantages of laparoscopic uterine vessel occlusion in the management of women with symptomatic myoma

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Accepted 12 July 2012

Abstract

Objective: The aim of this study was to evaluate the short-term therapeutic outcome of women with symptomatic uterine myomas treated with laparoscopic uterine vessel occlusion (LUVO) or laparoscopic myomectomy (LM).

Methods: Ninety-five patients with symptomatic, uncomplicated myomas warranting surgical treatment who expressed a strong desire to retain their uterus were included in this study. Fifty-two patients underwent LUVO and 43 underwent LM. The outcome was measured by comparing blood loss, surgical time, postoperative recovery, postoperative pain (visual analog scale), complications, and success rate in both groups.

Results: The general characteristics of the patients were similar in both groups. There were no statistical differences in febrile morbidity, complications, success rate, therapeutic efficacy (symptom relief), and satisfaction rate between the two groups. LUVO had advantages over LM, including less surgical time, minimal blood loss, lower visual analog scale score, and rapid postoperative recovery.

Conclusions: Both LUVO and LM might be effective in the management of symptomatic myomas in selected cases, but LUVO seemed to be more acceptable and less invasive in this 1-year short-term follow-up.

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Keywords: laparoscopic myomectomy; laparoscopic uterine vessel occlusion; myoma

Introduction

Approximately 20–40% of women of reproductive age are known to have uterine myomas [1], suggesting that uterine

myomas are remarkably common [2]. Most of these myomas are asymptomatic [3], although some may cause symptoms that require a definite treatment [4]. There are many therapeutic strategies, but the management of women with symptomatic uterine myomas depends on the patient's age, the reasons for treatment, the issue of fertility preservation, and the patient's preference [5]. Although hysterectomy has long been considered a good choice if women have completed childbearing [6], there are many other therapeutic approaches available for preservation of the uterus [7–11], because psychologically, the uterus has been regarded as the regulator

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and controller of important physiologic functions, a sexual organ, a source of energy and vitality, and a maintainer of youth and attractiveness [12].

Laparoscopic surgery has seen significant improvement in its techniques and instruments and has been developed and used in the management of various kinds of benign diseases [13–16]. Although some concerns about the role of laparoscopic myomectomy (LM) as a treatment option for symptomatic uterine myomas are still present [17], it has become more acceptable. Many comparison studies have evaluated the safety and feasibility of abdominal myomectomy and LM [18–22]. These studies showed that LM is clearly associated with shorter hospitalization, faster recovery, less expense, less pain, less blood loss, less fever, and fewer surgical complications compared with abdominal myomectomy. Pregnancy rates and recurrence rates appear to be comparable between LM and abdominal myomectomy [22]. A recent study comparing LM and minilaparotomic (MLT) myomectomy also showed the significant benefits of LM compared with MLT myomectomy, if these treatments were applied in a highly selected population (women with uterine fibroids with a size less than 8 cm and the number less than 5) [21,22]. Therefore, the use of LM can be considered as one of the choices in the management of women with symptomatic uterine myomas.

Although the uterine-sparing treatment of choice for symptomatic uterine myomas has been myomectomy [23–26], uterine artery embolization (UAE) was introduced in 1995 as an alternative technique for treating fibroid tumors [27]. Since then it has become increasingly accepted as a minimally invasive, uterine-sparing procedure, and studies have reported relief of excessive menstrual bleeding or pressure in 80–90% of patients [28–32]. These studies have also shown a reduction in leiomyoma and uterus size 3–12 months after the procedure, as measured by ultrasonography or magnetic resonance imaging [30]. A similar concept, known as laparoscopic uterine vessel occlusion (LUVU) or laparoscopic uterine artery occlusion (LUAO), was first reported as a treatment for myomas in 1999 or earlier [33,34]. The difference between the LUAO and LUVU methods is that LUAO is uterine artery occlusion alone without simultaneous blockage of the uterine vessels and the anastomotic sites between the uterine vessels and the ovarian vessels, whereas LUVU is uterine artery occlusion with simultaneous blockage of the uterine vessels and the anastomotic sites between the uterine vessels and the ovarian vessels. Similar relief of symptoms and reduction of the uterus and leiomyoma size were reported in 2001 in a 7–12-month follow-up of 87 patients after LUVU [35]. Since that time, there has been rapid growth in the use of this treatment with various modifications, such as simultaneous myomectomy, and there has been considerable research into its outcome [36–42]. For example, two studies with follow-ups as long as 36 months after surgery have confirmed the results of LUVU [43,44].

The aim of this study was to assess the short-term outcome of LUVU and LM in the management of symptomatic uterine myomas.

Materials and methods

Patients

The population of this study was derived from the Myoma Registry in the Department of Obstetrics and Gynecology (initiated from 1995). Women booked for myoma treatment who had symptomatic uterine myomas were invited to participate in the current study. All the myoma treatment included uterine-sparing and uterine-sacrificed treatments. Uterine-sparing treatment included medical treatment, radiologic approach, and surgical procedures; for example, myomectomy, which was performed through traditional exploratory laparotomy, MLT, ultraminilaparotomy, laparoscopy, vagina, and hysteroscopy approaches; and blockage of the uterine vessels, such as LUVU, uterine vessel occlusion (UVO), LUAO, uterine artery occlusion (UAO); or any combination of the above-mentioned therapy. These women were informed that they could choose to be treated with any one of the previously mentioned procedures. All women were treated based on their willingness and preference. Patients fulfilled the following criteria of the Taiwan National Health Insurance Bureau (NHIB) to comply with the definition of uncomplicated myomas: symptomatic myoma; a wish to retain the uterus; absence of previous abdominal or pelvic surgery; a number of visible uterine masses (myomas) less than or equal to five intramural or subserous myomas (without peduncle); a maximum diameter of no more than 8 cm; and an absence of prominent or significant pelvic adhesion on clinical evaluation. The phrase “symptomatic” included either menstrual problems such as menorrhagia and pain, or compression syndrome, including a bulgelike sensation and frequency. To make this study even more uniform and consistent, the general characteristics of the patients in the two groups were evaluated continuously during the study period using a limited number of evaluated items, such as age, body mass index (BMI), and biochemical blood data. This study was designed to compare the short-term therapeutic outcome of women with uncomplicated uterine myomas treated with LUVU and LM. Therefore, the following criteria should be met: a single operator; uncomplicated myoma; complete 1-year follow-up; LUVU alone or LM alone; and adequate sample size. There were 95 patients who were analyzed, including 52 women (54.7%) undergoing LUVU (LUVU group) and the remaining 43 (45.3%) undergoing LM (LM group).

Surgical procedures

Both procedures were performed under general anesthesia with the patient in the dorsolithotomy position and the bladder catheterized.

The detailed procedure for the LUVU group has been described previously [34–36]. In brief, after establishing a laparoscopic surgical field, the anterior leaf of the broad ligament was opened with scissors, keeping the peritoneum at proper tension by shifting the uterus to the opposite side. A vertical 2- to 3-cm incision was made on the triangle enclosed by

the round ligament, external iliac artery, and infundibulopelvic ligament with careful dissection of the ureter and the internal iliac artery, and adequate bleeding avoidance. The bilateral uterine artery was then thoroughly occluded. The anatomic sites between the uterine vessels and the ovarian vessels were also occluded.

The detailed procedure used in the LM group has been reported previously [26], with some modification. Myomas were extracted through the 12-mm suprapubic port by morcellation with the help of an electromechanical morcellator (Ethicon, San Angelo, TX, USA). The myometrial edges were closed in one or two layers, according to the depth of the uterine wound by means of polyglactin 0 sutures.

Evaluation parameters

The parameters we considered for comparing the two groups were: surgical time (minutes); blood loss; interval between completing surgery and tolerance of food intake; maximal fever; and complications such as blood transfusion, wound infection, or hematoma. A visual analog scale (VAS) applicable to the wound of each group was used to evaluate postoperative pain for 24 hours after surgery. The VAS consisted of a nongraduated 10-cm line ranging from “no pain” to “pain as bad as it could be”.

Leiomyoma-related symptoms, for example menstrual problems such as menorrhagia and pain, or compression syndrome, including a bulgelike sensation and frequency, were assessed using a yes/no improvement questionnaire at 3 months and 12 months after surgery. Recurrence was defined as any presence of “no” in a yes/no improvement questionnaire at 3 months and 12 months after surgery. A yes/no questionnaire was used at the discharge date and at the 3-month and 12-month follow-ups to evaluate satisfaction.

Statistical analysis

A sample size of 44 in each group will have 80% power to detect the differences in means of the selected outcome variables using a two-group Student *t* test with a 0.05 two-sided significance level.

SPSS Software Version 15.0 for Windows (SPSS Inc., Chicago, IL, USA) was used for statistical analysis. Descriptive statistics were presented as means \pm standard deviation or percentages. Means were compared by unpaired Student *t* test, and proportions were compared by Chi-square or Fisher exact tests, as appropriate. All calculated *p* values were two-tailed, and a *p* value less than 0.05 was considered statistically significant.

Results

Mean age, BMI, symptoms resulting from myoma, preoperative hemoglobin level, number of myomas, and maximal myoma diameter were similar in the two groups (Table 1). Significantly, surgical time in the LUVU group was shorter than in the LM group (36.9 ± 4.9 minutes vs. 86.7 ± 17.0 minutes); in addition, the amount of blood loss during the

Table 1
Baseline characteristics of the enrolled women.

	LUVU (<i>n</i> = 52)	LM (<i>n</i> = 43)	<i>p</i>
Age (y)	41.8 \pm 3.5	42.3 \pm 3.3	0.408
Body mass index (kg/m ²)	22.5 \pm 1.3	22.4 \pm 1.5	0.998
Preoperative HgB (g/dL)	9.4 \pm 1.8	9.7 \pm 1.7	0.391
Symptom			
Pain	19.2% (<i>n</i> = 10)	16.3% (<i>n</i> = 7)	0.781
Menorrhagia	69.2% (<i>n</i> = 36)	69.8% (<i>n</i> = 30)	0.936
Bulge sensation	46.2% (<i>n</i> = 24)	39.5% (<i>n</i> = 17)	0.375
Frequency	40.4% (<i>n</i> = 21)	34.9% (<i>n</i> = 15)	0.415
Myoma			
Number (<i>n</i>)	2.0 \pm 0.7	1.5 \pm 0.6	0.062
Max diameter (cm)	5.9 \pm 0.8	5.8 \pm 0.8	0.081

HgB = hemoglobin level; LM = laparoscopic myomectomy; LUVU = laparoscopic uterine vessel occlusion.

procedure in the LUVU group was also less than that in the LM group (24.7 ± 5.8 mL vs. 143.7 ± 81.7 mL). Postoperative recovery was significantly better in the LUVU group compared with the LM group because the mean of the maximal postoperative temperature of the LUVU group was significantly lower than that of the LM group, the VAS score in the LUVU group was significantly lower than that in the LM group, and food intake occurred sooner after surgery in the LUVU group compared with the LM group.

The complication rate seemed to be higher in the LM group, although it did not reach statistical significance (Table 2). Five patients in the LM group were considered as having complications and some of them had complications comprising one or more events, two with abdominal conversion due to technique difficulty, three with blood transfusion because of a more than 500-mL blood loss during surgery, two with ileus, and one with spiking fever ($\geq 39.0^\circ\text{C}$). There was only one patient in the LUVU group classified as having a complication. This patient had widespread subcutaneous ecchymosis extending from the right flank to the right thigh.

Most patients in both groups had symptom relief and elevated hemoglobin levels (ranging from 85.7% to 94.4% in the LUVU group and 83.3% to 95.3% in the LM group, based on different kinds of symptoms) at the first follow-up, which was evaluated at the end of 3 months after surgery (Table 3). These symptom-improvement effects continued in the 1-year follow-up in both groups. The cumulative recurrence rate in the LUVU group was 9.6% and 15.4% at the end of 3 months and 12 months after surgery, respectively, compared with 11.6% and 18.6% at the end of 3 months and 12 months after surgery, respectively, in the LM group. Neither was statistically significant.

Most patients in both groups still had symptom relief and had maintained the hemoglobin level (ranging from 80.0% to 90.4% in the LUVU group and 76.7% to 93.0% in the LM group) at the end of 12 months after surgery (Table 3). Patients in both groups showed a high immediate satisfaction rate (100% in the LUVU group vs. 97.7% in the LM group), and this satisfaction continued to the 3-month follow-up and the 1-year follow-up, as seen in Table 3, without statistical difference.

Table 2
Surgical and postoperative parameters following LUVU and LM.

	LUVU (n = 52)	LM (n = 43)	p
Surgical time (min) (95% CI)	36.9 ± 4.9 (35.0–38.7)	86.7 ± 17.0 (79.2–94.2)	0.0001
Blood loss (mL) (95% CI)	24.7 ± 5.8 (22.9–26.5)	143.7 ± 81.7 (102.8–184.7)	0.0001
Complications (95% CI)	1.9% (n = 1) (0–5.6)	11.6% (n = 5) (2.0–21.2)	0.069
Max fever (°C) (95% CI)	37.4 ± 0.2 (37.3–37.4)	37.7 ± 0.5 (37.5–37.9)	0.0001
VAS (95% CI)	3.1 ± 0.5 (2.9–3.2)	3.6 ± 0.3 (3.5–3.7)	0.0001
Tolerance to food intake (h) (95% CI)	4.7 ± 3.1 (3.7–5.6)	16.6 ± 7.3 (13.9–19.2)	0.0001
Satisfaction rate (95% CI)	100% (52/52)	97.7% (42/43) (93.2–100)	0.323

CI = confidence interval; complications = detailed information in the text; LM = laparoscopic myomectomy; LUVU = laparoscopic uterine vessel occlusion; VAS = visual analog scale.

Discussion

In gynecologic and reproductive surgeries, patients have been treated with different strategies and instruments depending on what is currently in vogue [45–47]. There is a growing consensus that patients undergoing surgery for benign gynecologic diseases would benefit from a minimally invasive approach [48]. The surgical accuracy and early postoperative advantages of laparoscopy have been documented recently in the treatment of benign and malignant gynecologic diseases [49,50]. In treating symptomatic uterine myoma, both LUVU [6,9,10,30,31,33–42] and LM [17–23,51] have been reported to provide satisfactory results.

We further confirmed in this study the concept that currently available instruments make LM feasible, although the wide application of this approach is limited by the size and number of myomas reasonably removed and the technical difficulty of the procedure and the laparoscopic suturing [17,32,50]. The criteria for our study population were limited to the definition of uncomplicated uterine myomas. However, the technical difficulties might have been overestimated with careful selection of a given population, such as those fulfilling the previously mentioned criteria. In this study, almost all LM operations were

performed without difficulty. LM surgical time ranged from 56 minutes to 164 minutes, with a mean of 87 minutes. These data were neither superior nor inferior to that of reports from all available studies, because almost all spent more than 1 hour, ranging from 62 minutes to 223 minutes, to complete the surgery, except for one study reporting a range of 30–140 minutes [52]. Furthermore, compared with abdominal myomectomy either through conventional exploratory laparotomy (mean = 99 minutes) or ultraminilaparotomy (mean = 98 minutes) [10], LM was not inferior. With regard to the surgical time, it was not surprising that LUVU was significantly superior to LM because the mean surgical time in the LUVU group was only 37 minutes, which was less than half of the mean time spent in the LM group (87 minutes).

The amount of estimated blood loss during LM in this study was 144 mL; this seemed to be a little more than in our previous study [23], which enrolled patients with the same criteria. The blood loss in the LUVU group was minimal. Furthermore, considering the complication rate in both groups, LUVU was also superior to LM, although this difference did not reach statistical significance. Based on the previously discussed findings, LUVU was a more feasible procedure in the management of uncomplicated uterine myoma, compared with LM.

In terms of the postoperative recovery in both groups, LUVU also provided significant advantages, including less postoperative pain (lower VAS score), fewer patients suffering from postoperative fever with resultant lower mean temperatures postoperatively, and a shorter interval from surgery to tolerance to food intake.

LUVU had significant advantages, not only in shorter surgical time and less blood loss, but also in better postoperative recovery with lower VAS score, less fever, and a shorter interval to food intake after surgery. In terms of 1-year durable symptom control, they were evaluated at 3 months and 12 months after surgery, respectively.

All myoma-related symptoms in the LM group, either menstrual problems such as menorrhagia and pain, or compression syndrome, including a bulgelike sensation and frequency, showed significant improvement, although they varied from 83.3% to 94.1%, at the end of 3 months after surgery. The hemoglobin level increased from 9.7 g/dL to 11.8 g/dL. The efficacy of symptom relief was maintained for 1 year, with 76.7% to 88.2% of patients showing a symptom relief status. In addition, the hemoglobin level was still maintained at 11.5 g/dL.

Table 3
Three-month and 12-month postoperative follow-up of the two groups of women.

	F-U (months)	LUVU (n = 52)	LM (n = 43)	p
Hemoglobin level	3	12.0 ± 0.5	11.8 ± 0.7	0.191
	12	11.7 ± 0.7	11.5 ± 0.9	0.951
Relief of symptoms (yes)				
Pain	3	90.0% (9/10)	85.7% (6/7)	1.00
	12	80.0% (8/10)	85.7% (6/7)	1.00
Menorrhagia	3	94.4% (34/36)	83.3% (28/30)	1.00
	12	88.9% (32/36)	76.7% (23/30)	0.729
Bulge sensation	3	91.7% (22/24)	94.1% (16/17)	1.00
	12	83.3% (20/24)	88.2% (15/17)	0.691
Frequency	3	85.7% (18/21)	93.3% (14/15)	0.627
	12	85.7% (18/21)	80.0% (12/15)	1.00
Recurrence rate	3	9.6% (5/52)	11.6% (5/43)	0.750
	12	15.4% (8/52)	18.6% (8/43)	0.676
Satisfaction rate	3	94.2% (45/52)	95.3% (41/43)	1.00
	12	90.4% (47/52)	93.0% (40/43)	1.00

F-U = follow-up; LM = laparoscopic myomectomy; LUVU = laparoscopic uterine vessel occlusion.

All of these factors contributed to the high satisfaction rate in the LM group. A similar efficacy was shown in the LUVU group, because symptom relief was reported as 85.7% to 94.4% at the end of 3 months after surgery, and 80.0% to 88.9% at the end of 12 months after surgery. This also contributed to the more than 90% satisfaction rate up to the 1-year follow-up.

Although LUVU seems to have its advantages in this 1-year study, there are many unanswered questions regarding LUVU and LM. For example, the long-term outcome and future reproductive outcome are unknown, and this study was not a randomized study, which resulted in some selection bias. In addition, we could not compare the myoma changes after each treatment because, although the evaluable items in the LUVU group were the complete disappearance of the myomas, the decreased number of myomas (tumor shrinkage to invisible status), and decreased size of the myomas, the myomas in the LM group were theoretically removed during surgery; thus, the reappearance of myomas in the LM group during follow-up meant recurrence.

A publication evaluating the midterm clinical and first reproductive results of the comparison between UAE and myomectomy hinted that although UAE is less invasive and as symptomatically effective and safe as myomectomy, myomectomy appears to have superior reproductive outcomes in the first 2 years after treatment [53]. Therefore, some studies suggested that myomectomy should be recommended as the treatment of choice over UAE in most patients desiring future fertility [54], even though most pregnancies following UAE have good outcomes [55]. In comparing reproductive outcomes between UAE and LUAO in women with symptomatic uterine myomas, a recent study showed that the pregnancies of women who were treated with uterine embolization were at significantly increased risk for spontaneous abortion when compared with the pregnancies of women treated with LUAO [56]. In our previous studies, we also demonstrated that immediate follicle-stimulating hormone changes were not significant after LUAO procedure, but women treated with LUVU were associated with a greater risk of a significant increase in follicle-stimulating hormone level at the first month after operation, which may be a reflection of diminished ovarian function, although the therapeutic effects seemed to be better in those patients who underwent LUVU [57,58].

In conclusion, this study showed the acceptability of both LUVU and LM in the management of symptomatic women with uncomplicated uterine myomas for the purpose of short-term symptom control. The effectiveness of both procedures is similar, but the LUVU group experienced less surgical and postoperative suffering, so LUVU seemed to be a better choice in this short-term 1-year follow-up. Based on these short-term advantages of LUVU, the strategy to use this procedure in the management of a certain group of patients might be worthy of further study.

Acknowledgments

The authors (Dr Wang and Dr Chang) contributed equally to this article. This work was supported in part by grants from

Veterans General Hospitals University System of Taiwan Joint Research Program (VGHUST99-G4), Taipei Veterans General Hospital (V99C1-085, V100C-054, V101C1-128, V101E4-004, and V101E5-006), and the National Science Council (NSC 99-2314-B-010-009-MY3), Taiwan.

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