

# Uterine artery embolization using gelatin sponge particles for symptomatic focal and diffuse adenomyosis

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**Objective:** To evaluate the effectiveness of uterine artery embolization (UAE) with gelatin sponge particles for symptomatic focal and diffuse adenomyosis. **Materials and Methods:** This was a retrospective study conducted between January 2014 and December 2019. All women underwent UAE for symptomatic adenomyosis without leiomyomas. Gelatin sponge particles were used in all cases. Patients were assessed for effectiveness of symptom control, changes in uterine volume, and degree of tumor necrosis on magnetic resonance (MR) imaging. **Results:** One hundred and sixty-three patients underwent UAE. There were statistically significant differences in bleeding score, pain score, and uterine volume at three months after UAE ( $P < 0.001$ ,  $P < 0.001$ , and  $P < 0.001$ , respectively). Complete tumor necrosis occurred in 66.9% of cases after UAE. Proportion of complete necrosis were significantly different in cases with focal adenomyosis compared with diffuse adenomyosis ( $P = 0.023$ ). Symptom recurrence occurred in 20.6% of cases at 12-month follow-up. Focal adenomyosis patients with complete necrosis had no recurrent menorrhagia and dysmenorrhea. No major complications were observed. **Conclusions:** UAE using gelatin sponge particles is an effective and safe treatment for symptomatic adenomyosis. Patients with focal adenomyosis with complete necrosis after UAE had no symptom recurrence at midterm follow-up.

## Keywords

Uterine artery embolization (UAE); Adenomyosis; Gelatin sponge particles

## 1. Introduction

Adenomyosis is described as the ectopic growth of endometrial tissues into the myometrium with adjacent smooth muscle hyperplasia [1]. The prevalence of adenomyosis is 20.9% [2], and the two most common presenting symptoms are menorrhagia and dysmenorrhea [3]. Surgical hysterectomy is considered to be the only definitive treatment for patients with symptomatic adenomyosis [4]. Although conservative therapies such as hormone therapy (oral contraceptive pills, selective estrogen/progesterone receptor modulators, a levonorgestrel releasing intrauterine device, danazol, aromatase inhibitors) and minimally invasive therapy are also widely used in treating adenomyosis, hormone therapies are unable to achieve mass resorption, and symptom recurrence

will occur when therapy is discontinued [5]. Other methods such as high intensity focused ultrasound (HIFU) have limited availability, relative high cost, unknown fertility outcomes, and strict indications [6].

Uterine artery embolization (UAE) on the other hand is minimally invasive, relatively low cost, allows preservation of uterus, and has fewer complications a low complication rate [7]. Therefore, UAE is more favourable compared with many hormone therapies and HIFU. Therefore, UAE is considered an alternative therapeutic method compared to many hormone therapies and HIFU.

UAE has emerged as a possible treatment option for patients with symptomatic adenomyosis [8–10]. Traditionally, UAE studies have used non spherical polyvinyl alcohol particles, with subsequent introduction of other embolic agents. These include tris-acryl gelatin microspheres (EmbosphereR), acrylamido polyvinyl alcohol microsphere, Bead Block, and spherical PVA microspheres (PVA; Contour SE) for symptomatic adenomyosis [8–12]. The disadvantages of non-spherical PVA particles were variation in the size of the particles and the tendency for them to aggregate resulting in obstruction of microcatheter and proximal vessel occlusion or unpredictable level of occlusion [13, 14]. Gelatin sponge particles have been used since 2001 as the primary embolic material in intervention radiology, and blood vessels occluded by gelatin sponge can be recanalized within several weeks [15]. Therefore, gelatin sponge particles are advantageous in that they are low-cost, temporary embolic agents that enable reintervention [15, 16].

There are few reports of UAE using gelatin sponge particles in the treatment of symptomatic adenomyosis [17].

Therefore, we aimed to investigate the effectiveness and safety of UAE with gelatin sponge particles for patients with symptomatic adenomyosis.



**Fig. 1.** MR images from a 30-year-old woman with focal adenomyosis. (1A) T2-weighted sagittal image shows asymmetrical thickening of uterine posterior wall (white arrow) with multifocal high signal spots. (1B) Gadolinium-enhanced T1-weighted sagittal image corresponding to (1A). (2A) T2-weighted sagittal image 3 months after embolization shows area of very low SI (white arrow). (2B) Gadolinium-enhanced T1-weighted sagittal image 3 months after embolization show no enhancement at the adenomyotic region (white arrow) suggesting complete necrosis.

## 2. Materials and methods

### 2.1 Patients

This retrospective study was approved by our Institutional Review Board (approval number 2020-SCMC-02-012) and informed consent was waived. We reviewed the medical records of patients who underwent UAE for symptomatic uterine adenomyosis between January 2014 and December 2019. Their ages ranged from 30 to 53 years (median age, 42 years). Their body mass index ranged from 19 to 32.8 kg/m<sup>2</sup> (mean, 23.7 kg/m<sup>2</sup>) and their parity ranged from 0 to 3 (mean, 1). All women were premenopausal and had clinical symptoms including menorrhagia, pelvic pain, and bulk-related symptoms associated with uterine adenomyosis.

Exclusion criteria were acute pelvic infection, pregnancy, gynecologic malignancy, endometriosis and contraindications or allergy to iodinated contrast medium. There were no cases of concomitant leiomyomas. Patients desiring future pregnancy were not excluded, and they were provided with information regarding potential benefits and risks associated with UAE. The follow-up period ranged from eight to 15 months (mean, 12.3 months).

### 2.2 Imaging

All patients underwent unenhanced and enhanced pelvic MR imaging with a 3.0-T Ingenia unit (Philips Healthcare, Eindhoven, the Netherlands). Axial and sagittal MR imaging were obtained before the procedure and at the 3-month follow-up. The T1-weighted images were obtained using the following parameters: repetition time (TR) msec/echo time (TE) msec, 570/10; field of view, 23 cm; matrix, 400 × 240; thickness, 4 mm; gap, 0.4 mm. Breath-hold T2-weighted images were obtained using the following parameters: TR msec/TE msec, 4200/80; echo train, 18; field of view, 23 cm; matrix, 430 × 275; thickness, 4 mm; gap, 0.4 mm. For this study, MR imaging images were evaluated by two radiologists with 15 years and four years of experience, who were blinded to patient data and each other's readings. Disagreements in interpretation were resolved by consensus.

Adenomyosis was considered when diffuse (> 12 mm) or focal thickening of the junctional zone was present. The pres-

ence of myometrial punctate high-signal foci was considered ancillary evidence of adenomyosis [18]. Adenomyosis was subdivided into two types by MR imaging: focal and diffuse. Focal adenomyosis was defined as an actual circumscribed mass within the myometrium. Diffuse adenomyosis was defined as diffuse ectopic growth of the endometrium with widening of junctional zone. Uterine volumes were calculated according to the equation of a prolate ellipse (length × depth × width × 0.5233), and the length of the uterus means only the uterus corpus, not including the cervix [19]. Necrosis was defined as the absence of contrast enhancement on T1-weighted imaging and was classified into three categories: complete, partial, and none. Complete necrosis was defined as well-defined necrosis that almost completely replaced adenomyosis high signal intensity on T1-weighted image, low signal intensity on T2-weighted image, and no contrast enhancement on enhanced T1-weighted images and represented over 90% of the non-perfusion area of adenomyosis (Fig. 1). Incomplete necrosis was defined as ill-defined necrosis that did not totally cover the adenomyosis considered to have occurred when some adenomyotic tissue remained though necrosis was clearly present in other areas and represented less than 90% of the non-perfusion area of adenomyosis (Fig. 2). No necrosis was defined as absence of necrosis in adenomyosis and represented 0% of the non-perfusion area of adenomyosis [20]. Complete necrosis between focal adenomyosis and diffuse adenomyosis was assessed by the procedure and at the 3-month follow-up MR imaging.

### 2.3 Angiography and embolization procedure

A unilateral femoral artery approach was performed in all cases under local anesthesia. A 5.0-F RUC catheter (Cook, Bloomington, IN, USA) was placed in the internal iliac artery and a coaxial 3-F microcatheter (Stride Hi-flow; Asahi Intecc, Osaka, Japan) was advanced distally into the uterine artery. Because of the possible need for reintervention, embolization was performed only using gelatin sponge particles (SPONGOSTAN; Johnson & Johnson, Skipton, UK) for symptomatic adenomyosis. Before the embolization proce-



**Fig. 2. MR images from a 44-year-old woman with diffuse adenomyosis.** (1A) T2-weighted sagittal image shows diffuse enlargement of uterus with multifocal bright spots. (1B) Gadolinium-enhanced T1-weighted sagittal image shows diffuse enhancement of adenomyosis. (2A) T2-weighted sagittal image 3 months after embolization shows area of very low SI (white arrow) and marked decrease in uterine size. (2B) On gadolinium-enhanced T1-weighted image 3 months after embolization, non-enhancing portions (white arrow) are seen in uterus, suggesting partial necrosis.

ture, the gelatin sponge particles were mixed with 40 mL of 1 : 1 saline solution-contrast agent mixture (Iomeron {Iomeperol}; Bracco, Milano, Italy). A 40-mL mixture with 150-350  $\mu\text{m}$  gelatin sponge particles was injected at the beginning of embolization into each uterine artery. If the flow rate of the mixture decreased in the intramural uterine artery, a mixture with 350-560  $\mu\text{m}$  particles was injected. When near obstruction of the intra-uterine vessel was achieved, a mixture with 560-710  $\mu\text{m}$  particles was injected into each uterine artery. The embolization continued until complete

cessation of blood flow was achieved in the proximal ascending uterine artery during ten cardiac beats. Preparation of gelatin sponge particles and embolization procedures were the same as previously described [21].

#### 2.4 Pain control

Pain within 24 hours of the procedure was primarily managed through an intravenous patient-controlled analgesia pump containing 1,500 mg fentanyl sulfate and 150 mg ketorolac tromethamine. Use of patient-controlled analge-



**Table 1. Baseline characteristics and outcomes.**

	Pre procedure (n = 163)	3-month follow-up (n = 163)	<i>P</i> value		12-month follow-up (n = 102)	<i>P</i> value
Age (years)	42 (30-53)					
Body mass index (kg/m <sup>2</sup> )	23.71 (18.97-32.77)					
Parity (no.)	1 (0-3)					
Hb (g/dL)	8.25 (3.80-10.00)	12.7 (9.20-14.70)	< 0.001		12.1 (10.7-14.2)	
WBC (10 <sup>3</sup> /uL)	5.37 (2.12-9.75)	5.48 (2.93-11.52) <sup>a</sup>	0.425			
CA125 (U/mL)	74.56 (6.37-653.00)	45.87 (6.21-450.10)	< 0.001		30.25 (6.01-78.39)	< 0.001
Bleeding score						
Focal type (n = 46)	7 (6-9)	2 (2-6)	0.004		2 (2-5)	0.004
Diffuse type (n = 56)	8 (6-9)	2 (1-6)	0.007		2 (1-6)	0.007
Overall (102)	8 (6-9)	2 (1-6)	< 0.001		2 (1-6)	< 0.001
Pain score						
Focal type (n = 45)	7 (5-9)	2 (1-6)	0.007		2 (1-5)	0.006
Diffuse type (n = 50)	7 (5-9)	2 (1-6)	0.026		2 (1-6)	0.02
Overall (n = 95)	7 (5-9)	2 (1-6)	< 0.001		2 (1-6)	< 0.001
Uterine volume (cm <sup>3</sup> )						
Focal type (n = 78)	242.8 (100.3-610)	130.6 (58.9-460)	0.001			
Diffuse type (n = 85)	310.4 (180.8-688.8)	173.3 (100-652.1)	0.012			
Overall (n = 163)	274.9 (100.3-688.8)	138.05 (58.9-652.1)	< 0.001			

<sup>a</sup> Blood data at 1-week follow-up.

sia pump was started after the procedure in the angiography suite. Nonsteroidal anti-inflammatory drugs were additionally administered by intravenous injection if needed and oral analgesic agents were prescribed simultaneously.

### 2.5 Clinical follow-up

Clinical response was assessed by questionnaire including symptomatic changes in menorrhagia and dysmenorrhea before UAE, and at the 3-month and 12-month follow-up visits. Number of used pads was used to quantify bleeding. Points were assigned for bleeding based on the number of pads used and were categorized into: 0 points (no pad was used), one point (1-3 pads), two points (4-6 pads), three points (7-9 pads), four points (10-12 pads), five points (13-15 pads), six points (16-18 pads), seven points (19-21 pads), eight points (22-24 pads), nine points (25-27 pads), 10 points ( $\geq 28$  pads). A visual analog scale (VAS) was used to describe pain, ranging from 0 (no pain) to 10 (extreme pain). Patients were asked to report improvement in symptoms on a scale of 0 (little bleeding, no pain) to 10 (severe menorrhagia, severe pain) according to the number of pads used, analgesics, and individual experiences.

### 2.6 Statistical analysis

Statistical analysis was performed with SPSS version 21.0 software (IBM Institute, Chicago, IL, USA). Continuous variables were expressed as mean and SD. Analyses were performed with Wilcoxon signed rank test for continuous variables and Pearson's chi-square tests or Fisher's exact test for categorical variables. A *P*-value < 0.05 was considered sta-

tistically significant.

## 3. Results

UAE was successfully performed in all 163 patients. The symptom outcomes and changes in uterine volume are shown in Table 1. The median scores of menorrhagia and pelvic pain at 12 months were reduced from 8 (6-9) to 2 (1-6) and from 7 (5-9) to 2 (1-6), respectively (*P* < 0.01). Mean uterine volume reduction from  $297.68 \pm 159.77$  to  $178.38 \pm 130.71$  (*P* < 0.01) was observed in 41% of patients at three months.

Of the 163 patients, 78 patients had focal adenomyosis and 85 had diffuse adenomyosis. Complete necrosis of adenomyosis after UAE was seen in 109 patients (66.9%), partial necrosis was seen in 45 patients (27.6%), and no necrosis was seen in nine patients (5.5%) (Table 2). Statistically significant differences were found in the occurrence of complete necrosis between focal adenomyosis and diffuse adenomyosis (*P* = 0.023).

Symptom recurrence occurred in 20.6% of patients at 12 months' follow-up after UAE (Table 2). Three patients had a hysterectomy. Patients who had focal adenomyosis with complete necrosis had no recurrent menorrhagia and dysmenorrhea. Of the nine patients with no necrosis after UAE, three patients exhibited no improvement in clinical symptoms after UAE and four patients had recurrent symptoms.

Ten patients with complete and partial necrosis of adenomyosis experienced transient amenorrhea. However, most patients resumed menstruation within seven months. Two patients aged 47 years and 50 years did not resume

**Table 2. Symptom recurrence after 12-month according to adenomyosis type and degree of necrosis.**

	n	Necrosis		
		Complete	Partial	None
Focal adenomyosis	74			
Recurrent dysmenorrhea	4	0/74 (0%)	2/74 (2.7%)	2/74 (2.7%)
Recurrent menorrhagia	2	0/74 (0%)	1/74 (1.4%)	1/74 (1.4%)
Diffuse adenomyosis	81			
Recurrent dysmenorrhea	8	2/81 (2.5%)	3/81 (3.7%)	3/81 (3.7%)
Recurrent menorrhagia	7	0/81 (0%)	3/81 (3.7%)	4/81 (4.9%)

menstruation until 12 months after UAE. There were 98 women of childbearing age before the procedure, and twelve of them. Twelve women attempted natural pregnancy and three women attempted artificial reproduction after procedure. Three women succeeded in becoming pregnant, one of whom underwent a natural pregnancy and the other two underwent artificial reproduction. One woman who conceived naturally, delivered a full-term baby. Two women who became pregnant with artificial reproduction had one miscarriage and the other had follow-up loss. Three women succeeded in becoming pregnant. One woman conceived naturally and delivered a full-term baby. Two women became pregnant after artificial reproduction, one had a miscarriage at 6 weeks, the other was lost to follow up.

There was no procedure related complications. Four patients complained of allergic reaction or rash, a minor complication, and they recovered with appropriate medications. Three other patients who reported symptoms of urinary tract infection after discharge were also recovered with medication. Two patients complained of urinary retention, and they recovered with conservative managements. There were no major complications that required readmissions.

#### 4. Discussion

Gelatin sponge is inexpensive and has been used as a safe and effective embolic agent for more than 30 years in UAE for some uterine disorders [22, 23]. Despite their merits, gelatin sponge particles have recently become available as an approved embolic agent in Korea.

It has been reported that UAE offers significant short- and long-term improvements in symptoms for both pure or combined adenomyosis [11]. In our study, there were significant symptom improvements or uterine volume reduction, ( $P < 0.01$ ), at three months after UAE. These results are similar to those reported by Kim *et al.* [24] in UAE using nonspherical polyvinyl alcohol particles.

While we found complete necrosis of adenomyosis in 66.9% of patients in the current study, the corresponding estimates reported by Kim *et al.* [24] with nonspherical polyvinyl alcohol particles and Lohle *et al.* [25] with calibrated tris-acryl gelatin microspheres were 82.5% and 44.1%, respectively. Our findings of statistically significant differences in complete necrosis between focal adenomyosis and diffuse adenomyosis ( $P = 0.023$ ) is inconsistent with estimates reported by

Kim *et al.* [24] who found no significant differences in the occurrence of complete necrosis between focal adenomyosis and diffuse adenomyosis. We believe that these differences may be related to differences in embolic material characteristics, sample size, and utero-ovarian anastomoses [26]. Unlike leiomyomas, previous studies with pathologic examination of specimens showed that embolic materials are randomly distributed in the myometrium in adenomyosis, and blood vessels feeding adenomyosis are not clearly defined [27, 28].

Pelage *et al.* [29] performed embolization with PVA particles or tris-acryl gelatin microspheres and reported that eight (53%) of 15 women had complete resolution at 1-year follow-up. In our study, 20.6% of patients had symptom recurrence at 12 months' follow-up. We assume that the discrepancy between these studies may be related to differences in embolic material used. Pelage *et al.* [29] and Kim *et al.* [24] used permanent embolic materials, such as PVA particles or tris-acryl gelatin microspheres, with variable diameters of 150-250  $\mu\text{m}$  to 500-900  $\mu\text{m}$ . However, in the present study, we used 150-250  $\mu\text{m}$  gelatin sponge particles at the beginning of embolization and later used predominantly 250-350  $\mu\text{m}$  particles.

Occurrence of complete necrosis of adenomyosis is thought to be positively associated with embolization [30, 31]. When adenomyosis had complete necrosis after UAE, there was no recurrence of menorrhagia. Only two out of 38 patients (5.7%) with diffuse type adenomyosis had recurrent dysmenorrhea in our study. This result was consistent with those reported by Kim *et al.* [24]. Therefore, we believe that achievement of complete necrosis of adenomyosis is the target in UAE for symptomatic adenomyosis.

A previous study by Kim *et al.* [32] using gelatin sponge particles, which assessed ovarian reserve using AMH after UAE for symptomatic uterine fibroids, also reported that UAE influenced ovarian function, but younger ovaries had greater capacity for recovery after ovarian damage. Therefore, we speculate that a variety of factors, including utero-ovarian anastomosis, embolic agent size, embolic agent type, extent of embolization, and the endpoint of embolization, may be associated with ovarian damage [12, 33].

In our study, one patient with complete necrosis of focal adenomyosis, a 30-year-old woman who desired future pregnancy, succeeded in becoming pregnant. Twenty-four months after UAE, this woman delivered a full-term baby. However, Liang *et al.* [34] reported pregnancy following UAE for adenomyosis is likely to be complicated and cannot be recommended at this time. So, larger studies should be performed for more conclusive results for patients wishing to preserve fertility.

In the present study, eight patients developed transient amenorrhea. Two patients over 47 years of age did not resume menstruation until 12 months after UAE. The frequency of minor complications was comparable with that of other studies. There were no major complications after UAE with gelatin sponge particles.

Our study has some limitations. Firstly, a significant number of patients were lost to follow up. This may have induced a potential bias. Secondly, we only included patients with adenomyosis without myoma. However, adenomyosis and myoma often coexist. Therefore, the results are not necessarily valid for all women with adenomyosis. Thirdly, the current study was conducted at a single institution with a relatively short follow-up period. So, multicenter study should be conducted to confirm these findings. Finally, this study was not a comparison study, therefore, further randomized controlled trials to compare the effects of UAE with non-spherical polyvinyl alcohol particles on imaging and clinical outcomes with those of UAE with gelatin sponge particles among patients with symptomatic adenomyosis are needed.

## 5. Conclusions

Despite its limitations, UAE using gelatin sponge particles is an effective and safe treatment for symptomatic adenomyosis without leiomyomas. Patients with focal adenomyosis with complete necrosis after UAE had no symptom recurrence at midterm follow-up.

## Author contributions

YG and MO designed the research study. JY and C-W performed the research. YG and JY analyzed the data. YG, MO and JY wrote the manuscript. All authors contributed to editorial changes in the manuscript. All authors read and approved the final manuscript.

## Ethics approval and consent to participate

This retrospective study was approved by Samsung Changwon hospital Institutional Review Board (SCMC 2020-02-012) and informed consent was waived.

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## Conflict of interest

We certify that there is no conflict of interest with any financial or other potential conflict of interest.

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