

Review

The Diagnosis and Treatment of Caesarean Scar Defect: Progress and Controversies

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Abstract

Objective: A caesarean scar defect (CSD), a niche in the uterus, is an emerging long-term complication after caesarean section (CS). With rising rates of CS, the risk of post-caesarean iatrogenic complications has increased, although many women may be diagnosed incidentally while asymptomatic. We present a review of the diagnosis and treatment of CSD. Mechanism: We describe the current state of treatment of CSD in terms of diagnosis and the different treatment options. Findings in Brief: The diagnosis can be confirmed by transvaginal sonography, hysteroscopy and other imaging examinations. At present, therapy is dependent on the presence of symptoms, the size of the CSD, and secondary infertility. Non-invasive management includes oral contraceptives and the levonorgestrel-releasing intrauterine system (LNG-IUS), which can improve the clinical symptoms of prolonged menstrual periods to some certain extent. Currently, the principal therapy is surgery, and micro-invasive surgical management includes transvaginal, hysteroscopic and laparoscopic procedures. However, there is no unified surgical selection standard, and the selection of surgical methods and the effect of surgical management are also controversial and need to be further studied. Conclusions: The aim of this review is to summarize and compare the selection of diagnostic and therapeutic methods and the differences in the curative effects of these therapeutic methods for CSD.

Keywords: caesarean scar defect (CSD); non-invasive treatment; laparoscopy; transvaginal sonography; hysteroscopy

1. Introduction

Caesarean section (CS) is one of the most frequently carried out surgical operations in modern obstetrics worldwide [1]. In recent decades, the caesarean delivery (CD) rate has increased significantly [2]. This has resulted in a globally increasing discussion of the impact and complications of the procedure [3]. The rates of some of these complications, such as uterine rupture, ectopic pregnancy, placenta previa and stillbirth, are constantly increasing [4]. However, there has been paid more attention to the main long-term complications of CS [1] which is caesarean scar defect (CSD), and obstetrics complication of CSD which are caesarean scar pregnancy and Placenta accrete spectrum disorders.

CSD refers to a myometrial defect resembling a pouch on the uterine incision after CS. More specifically, it is characterized by the separation of the myometrium at the incision of a previous caesarean scar on the lower uterine segment [5]. This defect contributes to the occurrence of symptoms such as abnormal uterine bleeding [6], chronic pelvic pain, caesarean scar pregnancy, dysmenorrhea and secondary infertility [7]. The guidelines for the diagnostic and therapeutic criteria of CSD are still unclear. At present, the treatment choices for CSD include conservative and surgical treatment, such as oral contraceptives, the levonorgestrel intrauterine system (LNG-IUS) and transvaginal, laparoscopic, or hysteroscopic repair.

The aim of this review is to provide a comprehensive and wide overview of the current literature by analysing the diagnosis and medical and surgical management of CSD to provide a reference for the selection of a clinical therapy project.

2. Diagnosis

At present, there are no definitive diagnostic criteria for CSD [1]. The diagnosis is based mainly on the medical history, clinical symptoms and auxiliary examination of patients. Patients with a history of CS have different clinical symptoms, such as chronic pelvic pain, postmenstrual spotting [8], infertility, and scar pregnancy, and some patients may be diagnosed incidentally while asymptomatic. Therefore, auxiliary examination should be further performed. There are currently multiple imaging methods that can be used for the diagnosis of CSD, such as ultrasonography, hysteroscopy, sonohysterography, and magnetic resonance imaging (MRI), which can be applied to evaluate the lower anterior myometrium and make a diagnosis of CSD. A prospective cohort study mentioned that the incidence of CSD was 44.4% at 6 months after CS. In addition, lower abdominal pain and scar pain were more pronounced when CSD was observed in transvaginal sonography (TVS) [9].

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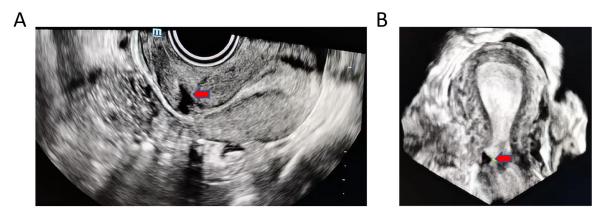


Fig. 1. Ultrasound images of CSD. (A) Transvaginal two-dimensional ultrasound. (B) Transvaginal three-dimensional ultrasound. The red arrow indicates the location of the CSD. CSD, caesarean scar defect.

2.1 Transvaginal Ultrasound (TVUS)

TVUS is a simple, low-cost, most common usage and non-invasive examination to evaluate the integrity of the lower myometrium on the anterior wall [10,11]. During routine scans, the use of a low-resolution ultrasound machine could neglect the pouch. Togas Tulandi *et al.* [1] reviewed the literature and discovered that the detection rate of CSD can reach 42%–88% when used by TVUS. Cecilia Fabres reported that TVUS was highly 100% correlated with hysteroscopy in the diagnosis of CSD, and the two diagnostic methods have similar positive and negative predictive values [12]. TVUS and sonohysterography showed no significant difference in evaluating and measuring the thicknesses of the residual myometrium, and TVUS examination readily identified this anatomic defect during the period of postmenstrual bleeding [13].

Compared with transabdominal ultrasound, TVUS decreased the limitation of conditions, such as intestinal gas accumulation, degree of bladder filling and fat layer thickness of the lower abdominal wall, and TVUS can more clearly show the shape of the CSD and its adjacency. Three-dimensional TVUS can clearly display the coronal plane, which makes up for the limitation of two-dimensional TVUS (Fig. 1A) and more intuitively and comprehensively shows the lesion morphology (Fig. 1B). Consequently, it is considered to be the preferred method for screening CSD. However, the level of the sonographers and the timing of the examination can easily influence the result of the CSD, especially the missed diagnosis of the smaller CSD.

2.2 Sonohysterography (SHG)

Ultrasound contrast agent is injected into the uterine cavity, and TVUS shows typical wedge-shaped or cystic dark liquid in the lower segment of the uterus. The CSD is more common in SHG than in TVUS, which shows that SHG has higher sensitivity [14–18]. Interestingly, the scar defect in SHG seems deeper and larger [14,19]. SHG is classified into saline infusion sonohysterography (SIS) and gel instillation sonography (GIS) according to which ultra-

sound contrast agent is used. As the contrast agent increases the contrast between the defect and uterine wall, SIS has a higher specificity and sensitivity than TVUS for identifying CSD [20]. One study demonstrated that 56.0% of women with a history of CS had a niche detected by GIS [17]. Van der Voet *et al.* [19] reported that the niches were visible in all women using GIS at 6–12 weeks after CS, and the prevalence of defects detected by GIS (64.5%) was higher than that detected by TVUS (49.6%). However, some researchers believe that SHG can enlarge defects and thus the size of the CSD [21].

2.3 Hysterography (HSG)

The contrast agent is perfused into the uterine cavity, and the defect is observed on X-ray. The CSD appears as cystic, banded or linear defects on X-ray imaging. HSG can be used to evaluate the CSD, but the width and length of the defect are increased compared with those on ultrasound, and this is associated with the outside pressure during the examination [22]. HSG has some limitations, such as the lack of measurement of myometrial thickness and the accumulation of mucus or blood in the CSD, which can lead to a missed diagnosis of CSD [23]. Consequently, HSG has been gradually replaced by SHG.

2.4 Hysteroscopy

Through hysteroscopy, the structure of the CSD can be directly seen in the depression of the uterine incision, such as the "valve" on the incision margin, highly vascularized on the surface, accumulation of old blood or mucus and sometimes the growth of endometrial tissue in the depression. Hysteroscopy is currently considered to be the best method for the diagnosis of CSD due to its direct and clear visualization and potential therapy [12]. Dominguez *et al.* [24] mentioned that there is gradual agreement to recommend hysteroscopic repair of CSD as the preferred method as long as the residual myometrium is at least 2.5–3 mm. However, it cannot be used to assess the residual myometrial thickness (TRM).



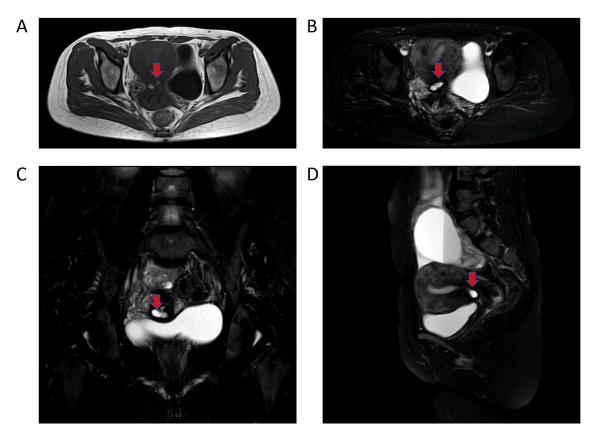


Fig. 2. Nuclear magnetic resonance (MRI) images of CSD. (A) Transverse section of the pelvis (T1W1). (B) Transverse section of the pelvis (T2W1). (C) Coronal section of the pelvis (T2W1). (D) Sagittal section of the pelvis (T2W1). The red arrow indicates the location of the CSD. CSD, caesarean scar defect.

2.5 Magnetic Resonance Imaging (MRI)

MRI of CSD shows the occurrence of an iso-signal or high-signal on T1-weighted imaging (W1) (Fig. 2A) and a high-signal on T2W1 (Fig. 2B–D). MRI has advantages in the display of soft tissue and clearly images of the site of the CSD and the absence of the myometrium. Additionally, it has the advantages of its non-invasiveness and lack of radiation. Taru Gupta *et al.* [25] reported that MRI was better than TVUS, but the difference in the rate of CSD identification was not statistically significant. However, the disadvantages of MRI include its high cost and longer examination time, so it is not the best first-choice option to screen for CSD.

Consequently, TVUS is more affordable, is non-invasive and yields reliable results, and it is the best first-choice option to screen for CSD [23]. For some cases, several studies have suggested SIS as a diagnostic tool for planning surgical treatment [6,26].

3. Treatment

The therapeutic options for CSD comprise various regimens depending on the plans for pregnancy, the presence of clinical symptoms, secondary infertility and the willingness to undergo surgery [27,28]. However, the efficacy of therapy has yet to be ascertained. For the diagnosis

of CSD with no plan for pregnancy, routine surgical repair is not recommended [23]. Therapy options for CSD are as follows.

3.1 Conservative Treatment

3.1.1 Oral Contraceptives

When there is no plan for pregnancy or the patient declines surgery, short-acting oral contraceptives are the typical choices for abnormal uterine bleeding (AUB) as the clinical manifestation. This is a short-term treatment. It can effectively improve the symptoms of AUB but does not heal the defect, so the symptom recurrence and the complications of hormones are still important issues [29].

3.1.2 Levonorgestrel Intrauterine System (LNG-IUS)

There are quite a few studies on the application of the LNG-IUS to CSD. Xuyun Zhang *et al.* [30] performed a prospective evaluation of five methods applied to treat CSD, and the duration of menstruation was similar before and after treatment in the LNG-IUS group. Thus, given this lack of difference, the therapeutic application of the LNG-IUS is inadequate, but the long-term clinical efficacy of the LNG-IUS remains to be determined.



A B

Fig. 3. CSD in repeated CS. The blue arrow points to the CSD. (A) The lower segment of the uterus in late pregnancy after opening the abdominal cavity. (B) Lower segment of the uterus after removal of the fetus following incision of the uterus. CSD, caesarean scar defect. CS, caesarean section.

3.2 Surgical Treatment

The indication for surgical treatment is a diagnosis of CSD with corresponding clinical symptoms and the experience of therapeutic effects by the patient. Surgical treatment for CSD can prevent recurrence and restore the normal anatomy of uterus after drug cessation. Therefore, there are many studies suggesting different surgical approaches to repair the CSD [7,27,31,32], including laparotomy, hysteroscopy (HP), laparoscopy (LP), transvaginal repair (TVR), and combined HP with LP (LH). Currently, minimally invasive surgery is the most common procedure for CSD. The timing of surgery is generally 2–3 days after the completion of menstruation if menstruation is normal and generally between 7–9 days if bleeding during menstruation is abnormal.

3.2.1 Laparotomy

Laparotomy is the traditional surgery for the CSD repair. Because of the large surgical field and ease of operation, it is suitable for CSD with serious adhesions in the pelvic cavity. However, due to its disadvantages regarding the large scale of operative trauma and slow postoperative recovery, it is not the first choice to treat CSD at present.

3.2.2 Transvaginal Repair (TVR)

TVR is performed by dissecting the bladder from the cervix and uterus, exposing the cervix and lower uterine segment to clarify the position of the CSD, resecting the defect and suturing the two layers. TVR is a minimally invasive and effective surgical approach [33]. Zhou et al. [34] suggested that TVR relieved the symptoms of postmenstrual spotting and anatomically corrected the scars. Moreover, the optional timing for TVR is an interval ≤ 2.5 years between caesarean sections, and the standard healing

criterion for CSD is a thickness of the remaining muscular layer of ≥ 8.5 mm at 6 months. Zhang *et al.* [35] proposed that TVR has the advantages of lower cost and shorter operation time than LP for correcting the CSD. A novel therapeutic approach for repairing a symptomatic CSD by using vaginal natural cavity transluminal endoscopic surgery [36]. TVR is helpful for anatomic reconstruction of CSD. However, TVR requires the surgeon to be experienced in vaginal procedures, and the bladder peritoneal fold and the anterior wall of the lower uterine segment often develop adhesions after caesarean section, making the surgical field hard to expose. Furthermore, the size of the defect is usually a few millimetres, and it is difficult to locate the position accurately.

3.2.3 Hysteroscopy (HP)

Electric resection is performed by placing the hysteroscope and an electric resection ring (or an electric rolling ball) into the uterine cavity, and hysteroscopic electric resection is then performed. HP can remove the upper valve of the CSD by blocking the outflow of menstrual blood [37]. HP contributes to alleviating the clinical symptoms of AUB, but it cannot change the anatomic abnormalities of the CSD [31]. In a retrospective review, Xie et al. [38] reported that the operative efficacy of HP (64.5%) was significantly inferior to that of vaginal surgery (93.5%), whereas patients who underwent HP had a shorter operative time (25 vs. 55 min) and less blood loss (10 vs. 50 mL). Bingqing Lv et al. [39] compared HP combined with LP to HP alone, and the results showed similar pregnancy rates between the two methods (61.5% vs. 73.3%; p = 0.505); however, the interval between the operation and pregnancy was much shorter in the HP group than in the HP combined with LP group (10 vs. 27 months). One study showed that the effective



rate of menstrual improvement was 59%–100% [40]. Additionally, Florio *et al.* [26] stated that bipolar electrocision should be performed when the TRM muscles are not too thin (≥ 2 mm), and electrical coagulation should be performed if there is no active bleeding in the CSD. Another study also suggested that HP may be considered for patients who do not plan to become pregnant and with TRM > 3 mm or clinical symptoms [41]. A meta-analysis reported that hysteroscopic remodelling appears to be a treatment choice in the management of symptomatic CSD [42]. Channeled (360°) hysterectomy for symptomatic CSD patients may result in increased TRM, reduced CSD, improved symptoms and high patient satisfaction after the procedure [43]. Currently, whether TRM is a limitation of HP is controversial.

3.2.4 Laparoscopy (LP)

LP has the advantages of a full field of vision, less trauma, shorter operation time and faster recovery. It has been advocated for defects (TRM < 3 mm) and patients with symptoms who plan to become pregnant [44]. Based on common practice and expert opinion, Roy Mashiach recommended 2.5 mm as the cut-off value [27]. Conventional LP or robotic-assisted LP can be used by a skilled laparoscopic surgeon to repair the CSD [45]. The reported operation time is 65 ± 26 min [46], and robotic excision requires 240 minutes [47]. Vervoort et al. [48] reported a prospective cohort study that evaluated the effectiveness of an LP resection for CSD-related symptoms, fertility-related problems, and ultrasound findings in 101 women. In this study, the results showed that postmenstrual spotting and dysmenorrhoea were reduced by the laparoscopic approach, and the TRM was enlarged 6 months after the surgery. Olivier Donnez et al. [44] reported that eight women (44%) became pregnant and performed CS at 38-39 weeks of gestation among the 18 women with infertility. A two-centre observational study reported that CSD repair via robot-assisted laparoscopy is a feasible potential therapy, but further research into this technique is still needed [49].

3.2.5 Hysteroscopy Combined with Laparoscopy

When the CSD does not protrude from the serosa layer, LP does not directly observe the CSD. Thus, LP alone may not completely resect the scar. LH can solve the problems of the poor location ability of LP and incomplete repair of HP for CSD. During HP, the scope of CSD can be identified by the light transmission of the hysteroscope, and suturing of the CSD can be confirmed after laparoscopic repair [23]. During LP, the bladder can be fully pushed down, and the risk of bladder injury can be minimized. In addition, surgical treatment of other pelvic diseases, such as endometriosis, can be immediately performed [50]. LH is better than LP or HP alone, but the disadvantages of LH are its longer operation time and higher cost.

3.2.6 Detect and Repair CSD during Caesarean Section

In addition to some of the specialised procedures for repairing CSD mentioned above, there is another type of repairment of CSD when found in a repeated CS. In repeat lower segment caesarean section (LSCS), we found some women with a significant scar defect in the anterior uterine wall (Fig. 3). One prospective study noted that a modified surgical technique concentrated on identification, mobilisation and location of the retracted myometrial fringes by a unique experienced obstetrician [51]. The purpose of this is to make the recovery of anatomical results consistent. The result showed no diverticula in all cases at three months postpartum on pelvic ultrasound. The mean residual scar thickness was 8.4 mm) [51]. In summary, it makes sense to discover and reconstruct the structure of the lower myometrium while repeating the cesarean section.

4. Controversies

At present, there is still no uniform standard regarding the principles of treatment of CSD worldwide. The following aspects remain controversial: (1) the basis for the diagnosis of CSD on symptoms, imaging or a combination of both; (2) the necessity and duration of pharmacological treatment; and (3) the criteria for selecting different surgical procedures for CSD.

5. Conclusions

Post-caesarean CSD has become a common clinical condition. The most direct way to reduce the CSD is to reduce the rate of caesarean delivery. The associations between CSD and pelvic pain, dysmenorrhea and infertility requires more study. The most popular therapy appears to be hysteroscopic isthmoplasty. However, the efficacy of laparoscopic or vaginal surgery remains to be seen. These treatments should be considered for specific patients until an optimal treatment for symptomatic women with CSD is identified. At this point, we still need randomized controlled trials to obtain concrete evidence to support effective treatment for CSD.

Author Contributions

ZW and YK designed and performed the research study. SL and FH analyzed the data. All authors contributed to editorial changes in the manuscript. All authors read and approved the final manuscript.

Ethics Approval and Consent to Participate

Not applicable.

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Conflict of Interest

The authors declare no conflict of interest.

References

- [1] Tulandi T, Cohen A. Emerging Manifestations of Cesarean Scar Defect in Reproductive-aged Women. Journal of Minimally Invasive Gynecology. 2016; 23: 893–902.
- [2] Li HT, Hellerstein S, Zhou YB, Liu JM, Blustein J. Trends in Cesarean Delivery Rates in China, 2008-2018. The Journal of the American Medical Association. 2020; 323: 89–91.
- [3] Bij de Vaate AJM, van der Voet LF, Naji O, Witmer M, Veersema S, Brölmann HAM, et al. Prevalence, potential risk factors for development and symptoms related to the presence of uterine niches following Cesarean section: systematic review. Ultrasound in Obstetrics & Gynecology. 2014; 43: 372–382.
- [4] Sandall J, Tribe RM, Avery L, Mola G, Visser GH, Homer CS, et al. Short-term and long-term effects of caesarean section on the health of women and children. Lancet. 2018; 392: 1349–1357.
- [5] Vervoort AJMW, Uittenbogaard LB, Hehenkamp WJK, Brölmann HAM, Mol BWJ, Huirne JAF. Why do niches develop in Caesarean uterine scars? Hypotheses on the aetiology of niche development. Human Reproduction. 2015; 30: 2695–2702.
- [6] Tower AM, Frishman GN. Cesarean scar defects: an underrecognized cause of abnormal uterine bleeding and other gynecologic complications. Journal of Minimally Invasive Gynecology. 2013; 20: 562–572.
- [7] Abacjew-Chmylko A, Wydra DG, Olszewska H. Hysteroscopy in the treatment of uterine cesarean section scar diverticulum: A systematic review. Advances in Medical Sciences. 2017; 62: 230–239.
- [8] Borges LM, Scapinelli A, de Baptista Depes D, Lippi UG, Coelho Lopes RG. Findings in patients with postmenstrual spotting with prior cesarean section. Journal of Minimally Invasive Gynecology. 2010; 17: 361–364.
- [9] Gozzi P, Hees KA, Berg C, David M, Wernecke KD, Hellmeyer L, et al. Frequency and associated symptoms of isthmoceles in women 6 months after caesarean section: a prospective cohort study. Archives of Gynecology and Obstetrics. 2023; 307: 841– 848.
- [10] Marotta ML, Donnez J, Squifflet J, Jadoul P, Darii N, Donnez O. Laparoscopic repair of post-cesarean section uterine scar defects diagnosed in nonpregnant women. Journal of Minimally Invasive Gynecology. 2013; 20: 386–391.
- [11] Brölmann HAM, BijdeVaate M, van der Voet L, Naji O, Hehenkamp W, Bourne T, *et al.* Letter to the Editor JMIG on the Article "Cesarean scar defects: an underrecognized cause of abnormal uterine bleeding and other gynecologic complications" (2013;20; 562-572). Journal of Minimally Invasive Gynecology. 2014; 21: 498–499.
- [12] Fabres C, Aviles G, De La Jara C, Escalona J, Muñoz JF, Mackenna A, *et al.* The cesarean delivery scar pouch: clinical implications and diagnostic correlation between transvaginal sonography and hysteroscopy. Journal of Ultrasound in Medicine. 2003; 22: 695–700; quiz 701–702.
- [13] Chang Y, Tsai EM, Long CY, Lee CL, Kay N. Resectoscopic treatment combined with sonohysterographic evaluation of women with postmenstrual bleeding as a result of previous cesarean delivery scar defects. American Journal of Obstetrics and Gynecology. 2009; 200: 370.e1–370.e4.
- [14] Osser OV, Jokubkiene L, Valentin L. Cesarean section scar

- defects: agreement between transvaginal sonographic findings with and without saline contrast enhancement. Ultrasound in Obstetrics & Gynecology. 2010; 35: 75–83.
- [15] Antila-Långsjö R, Mäenpää JU, Huhtala H, Tomás E, Staff S. Comparison of transvaginal ultrasound and saline contrast sonohysterography in evaluation of cesarean scar defect: a prospective cohort study. Acta Obstetricia et Gynecologica Scandinavica. 2018; 97: 1130–1136.
- [16] Osser OV, Jokubkiene L, Valentin L. High prevalence of defects in Cesarean section scars at transvaginal ultrasound examination. Ultrasound in Obstetrics & Gynecology. 2009; 34: 90–97.
- [17] Bij de Vaate AJM, Brölmann HAM, van der Voet LF, van der Slikke JW, Veersema S, Huirne JAF. Ultrasound evaluation of the Cesarean scar: relation between a niche and postmenstrual spotting. Ultrasound in Obstetrics & Gynecology. 2011; 37: 93– 99.
- [18] Jastrow N, Chaillet N, Roberge S, Morency AM, Lacasse Y, Bujold E. Sonographic lower uterine segment thickness and risk of uterine scar defect: a systematic review. Journal of Obstetrics and Gynaecology Canada. 2010; 32: 321–327.
- [19] van der Voet LF, Bij de Vaate AM, Veersema S, Brölmann HAM, Huirne JAF. Long-term complications of caesarean section. The niche in the scar: a prospective cohort study on niche prevalence and its relation to abnormal uterine bleeding. BJOG: an International Journal of Obstetrics and Gynaecology. 2014; 121: 236– 244.
- [20] Thurmond AS, Harvey WJ, Smith SA. Cesarean section scar as a cause of abnormal vaginal bleeding: diagnosis by sonohysterography. Journal of Ultrasound in Medicine: Official Journal of the American Institute of Ultrasound in Medicine. 1999; 18: 13–16; quiz 17–18.
- [21] Pešková V, Kacerovský M. Cesarean scar defect manifestation, diagnostics, treatment. Ceska Gynekologie. 2020; 85: 282–287.
- [22] Dosedla E, Gál P, Calda P. Association between deficient cesarean delivery scar and cesarean scar syndrome. Journal of Clinical Ultrasound. 2020; 48: 538–543.
- [23] Sipahi S, Sasaki K, Miller CE. The minimally invasive approach to the symptomatic isthmocele - what does the literature say? A step-by-step primer on laparoscopic isthmocele - excision and repair. Current Opinion in Obstetrics & Gynecology. 2017; 29: 257–265.
- [24] Dominguez JA, Pacheco LA, Moratalla E, Carugno JA, Carrera M, Perez-Milan F, et al. Diagnosis and management of isthmocele: a SWOT analysis. Ultrasound in Obstetrics & Gynecology. 2023. (online ahead of print)
- [25] Gupta T, Singal K, Gupta N, Kohli S, Kanyal M. Comparative Study of USG and MRI in Evaluation of Isthmocele. Journal of Obstetrics and Gynaecology of India. 2021; 71: 292–296.
- [26] Florio P, Filippeschi M, Moncini I, Marra E, Franchini M, Gubbini G. Hysteroscopic treatment of the cesarean-induced isthmocele in restoring infertility. Current Opinion in Obstetrics & Gynecology. 2012; 24: 180–186.
- [27] Mashiach R, Burke YZ. Optimal Isthmocele Management: Hysteroscopic, Laparoscopic, or Combination. Journal of Minimally Invasive Gynecology. 2021; 28: 565–574.
- [28] Kulshrestha V, Agarwal N, Kachhawa G. Post-caesarean Niche (Isthmocele) in Uterine Scar: An Update. Journal of Obstetrics and Gynaecology of India. 2020; 70: 440–446.
- [29] Tahara M, Shimizu T, Shimoura H. Preliminary report of treatment with oral contraceptive pills for intermenstrual vaginal bleeding secondary to a cesarean section scar. Fertility and Sterility. 2006; 86: 477–479.
- [30] Zhang X, Yang M, Wang Q, Chen J, Ding J, Hua K. Prospective evaluation of five methods used to treat cesarean scar defects. International Journal of Gynaecology and Obstetrics. 2016; 134: 336–339.



- [31] He Y, Zhong J, Zhou W, Zeng S, Li H, Yang H, et al. Four Surgical Strategies for the Treatment of Cesarean Scar Defect: A Systematic Review and Network Meta-analysis. Journal of Minimally Invasive Gynecology. 2020; 27: 593–602.
- [32] Setúbal A, Alves J, Osório F, Sidiropoulou Z. Demonstration of Isthmocele Surgical Repair. Journal of Minimally Invasive Gynecology. 2021; 28: 389–390.
- [33] Luo L, Niu G, Wang Q, Xie HZ, Yao SZ. Vaginal repair of cesarean section scar diverticula. Journal of Minimally Invasive Gynecology. 2012; 19: 454–458.
- [34] Zhou J, Yao M, Wang H, Tan W, Chen P, Wang X. Vaginal Repair of Cesarean Section Scar Diverticula that Resulted in Improved Postoperative Menstruation. Journal of Minimally Invasive Gynecology. 2016; 23: 969–978.
- [35] Zhang Y. A Comparative Study of Transvaginal Repair and Laparoscopic Repair in the Management of Patients With Previous Cesarean Scar Defect. Journal of Minimally Invasive Gynecology. 2016; 23: 535–541.
- [36] Baekelandt JF, Kapurubandara S. A novel approach using vaginal natural orifice transluminal endoscopic surgery to repair a symptomatic uterine isthmocele. Fertility and Sterility. 2023; 119: 328–330.
- [37] Wang CJ, Huang HJ, Chao A, Lin YP, Pan YJ, Horng SG. Challenges in the transvaginal management of abnormal uterine bleeding secondary to cesarean section scar defect. European Journal of Obstetrics, Gynecology, and Reproductive Biology. 2011; 154: 218–222.
- [38] Xie H, Wu Y, Yu F, He M, Cao M, Yao S. A comparison of vaginal surgery and operative hysteroscopy for the treatment of cesarean-induced isthmocele: a retrospective review. Gynecologic and Obstetric Investigation. 2014; 77: 78–83.
- [39] Lv B, Xie X, Liu C, Lin Y. Laparoscopic combined with hysteroscopic repair or operative hysteroscopy in the treatment of symptomatic cesarean-induced diverticulum. Medecine Sciences. 2018; 34: 47–51.
- [40] Pan Y, Liu MB. The value of hysteroscopic management of cesarean scar pregnancy: a report of 44 cases. Taiwanese Journal of Obstetrics & Gynecology. 2017; 56: 139–142.
- [41] Nezhat C, Falik R, Li A. Surgical management of niche, isth-mocele, uteroperitoneal fistula, or cesarean scar defect: a critical rebirth in the medical literature. Fertility and Sterility. 2017;

- 107: 69-71.
- [42] Vrijdaghs V, Dewilde K, Froyman W, Van den Bosch T. Hysteroscopic management of caesarean scar defects. Journal of Obstetrics and Gynaecology. 2022; 42: 816–822.
- [43] Casadio P, Raffone A, Alletto A, Filipponi F, Raimondo D, Arena A, *et al.* Postoperative morphologic changes of the isthmocele and clinical impact in patients treated by channel-like (360°) hysteroscopic technique. International Journal of Gynaecology and Obstetrics. 2023; 160: 326–333.
- [44] Donnez O, Donnez J, Orellana R, Dolmans MM. Gynecological and obstetrical outcomes after laparoscopic repair of a cesarean scar defect in a series of 38 women. Fertility and Sterility. 2017; 107: 289–296.e2.
- [45] Setubal A, Alves J, Osório F, Guerra A, Fernandes R, Albornoz J, et al. Treatment for Uterine Isthmocele, A Pouchlike Defect at the Site of a Cesarean Section Scar. Journal of Minimally Invasive Gynecology. 2018; 25: 38–46.
- [46] Li C, Guo Y, Liu Y, Cheng J, Zhang W. Hysteroscopic and laparoscopic management of uterine defects on previous cesarean delivery scars. Journal of Perinatal Medicine. 2014; 42: 363– 370
- [47] Yalcinkaya TM, Akar ME, Kammire LD, Johnston-MacAnanny EB, Mertz HL. Robotic-assisted laparoscopic repair of symptomatic cesarean scar defect: a report of two cases. The Journal of Reproductive Medicine. 2011; 56: 265–270.
- [48] Vervoort A, Vissers J, Hehenkamp W, Brölmann H, Huirne J. The effect of laparoscopic resection of large niches in the uterine caesarean scar on symptoms, ultrasound findings and quality of life: a prospective cohort study. BJOG: An International Journal of Obstetrics and Gynaecology. 2018; 125: 317–325.
- [49] Cardaillac C, Salmon C, Vaucel E, Gueudry P, Lavoue V, Nyangoh Timoh K, et al. Robot-assisted laparoscopy repair of uterine isthmocele: A two-center observational study. International Journal of Gynaecology and Obstetrics. 2023; 160: 244–248.
- [50] Li C, Tang S, Gao X, Lin W, Han D, Zhai J, et al. Efficacy of Combined Laparoscopic and Hysteroscopic Repair of Post-Cesarean Section Uterine Diverticulum: A Retrospective Analysis. BioMed Research International. 2016; 2016: 1765624.
- [51] Siraj SHM, Lional KM, Tan KH, Wright A. Repair of the myometrial scar defect at repeat caesarean section: a modified surgical technique. BMC Pregnancy and Childbirth. 2021; 21: 559.

