Caesarean section scar endometriosis: quo vadis?

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Endometriosis is characterized by the presence of endometrial tissue implants outside the uterine cavity that responds to hormonal stimulation. These implants can be detected in all areas surrounding the uterus, ovaries, posterior cul-de-sac, ligaments of the uterus, pelvic peritoneum, and rectovaginal septum. Endometriosis may be infrequently found in the thorax, gastrointestinal tract, appendix, urinary tract, central nervous system, nose, umbilicus, lower limbs, and cutaneous cellular tissues. Caesarean scar endometriosis, id est, the cutaneous endometriosis, is the most common extrapelvic form, vulgo, and is located in scars following obstetric and/or gynecologic surgical procedures, such as caesarean delivery, hysterotomy, hysterectomy, episiotomy, ectopic pregnancy, salpingostomy, and tubal ligations, but scarcely in scars following appendicectomy, in the laparoscopic trocar and amniocentesis needle tracts. Diagnosis of surgical scar endometriosis following caesarean section, possessing an incidence of 0.03-0.4%, is not an easy process due to being often mistaken for a suture granuloma, lipoma, abscess, cyst, desmoid tumors, malignancies, incisional hernia, or a strange body¹⁻⁵. Cellular transport theory, coelomic metaplasia theory, and the endometrial tissue reaching the surgical scar through the lymphatic or vascular pathways in order to develop into scar endometriosis afterward are accused and argued in the pathophysiology of the disease, to date⁵. Although mass in a caesarean section scar with symptoms of cyclic pain associated with menstruation is nearly pathognomonic, imaging modalities assist in identifying the condition. In spite of all odds, histopathologic evaluation is the major tool for confirmation¹⁻⁴. Surgical scar endometrioses are known as possessing a potential for the progression of transformation, which rarely transpires for the malignant degeneration, accounting for 0.3–1%^{1,6}. Herein, the interval of time from the onset of the benign lesion to the development of malignant form in caesarean section scar endometriosis has been defined as a broad variation, ranging from 3 to 39 years with a mean of 17 years^{1,7}. In the upfront surgery setting,

in particular, wide surgical excision with a safety margin with or without reconstruction has been recommended for the surgical procedures of endometriosis, per se, the gold standard treatment of choice^{1,6}. As well as to avoid the possible transformation, some authors recommend a wide excision with at least 10 mm margins in order to prevent the recurrence⁶. Some authors recommended surgical resection with margins at least 5 mm in diameter and depth^{8,9}. Although the pathogenesis of endometriosis is not precisely known, immunologic factors, metaplasia, and confounding factors such as diagnosis of endometriosis before the first delivery, breastfeeding, previous surgery, and hormonal contraception are important. Theoretically, pregnancy, per se, a state of the altered immune response, and caesarean section could augment the risk of developing endometriosis. Some authors emphasized that the cases with two caesarean sections did not augment the risk of being diagnosed with endometriosis, compared with one caesarean section. In addition, it was stated that those who diagnosed with caesarean scar endometriosis after the first caesarean section are no longer at risk of developing endometriosis for the first time. *Inter alia*, some authors proclaimed that caesarean scar endometriomas are more common after unlabored caesarean sections^{10,11}. In fine, bene diagnosticur, bene curatur. Reddite ergo quae sunt Caesaris, Caesari.

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REFERENCES

- Sengul I, Sengul D, Soares Junior JM. Interpretations on a rare localization of endometriosis: labium minus. Rev Assoc Med Bras (1992). 2021;67(1):1-2. https://doi.org/10.1590/1806-9282.67.01.001
- Sengul I, Sengul D, Kahyaoglu S, Kahyaoglu I. Incisional endometriosis: a report of 3 cases. Can J Surg. 2009;52(5):444-5. PMID: 19865584
- 3. Sengul D, Sengul I. Scar endometriosis: a review. Remedy Open Access. 2016;1(1): 1038.
- Sengul I, Sengul D. Clinical presentation of incisional (cesarean section scar) endometriosis vary with the size of the lesion. Erciyes Med J. 2011;33(2):171-2.
- Nominato NS, Prates LFVS, Lauar I, Morais J, Maia L, Geber S. Caesarean section greatly increases risk of scar endometriosis. Eur J Obstet Gynecol Reprod Biol. 2010;152(1):83-5. https://doi. org/10.1016/j.ejogrb.2010.05.001
- Kocher M, Hardie A, Schaefer A, McLaren T, Kovacs M. Cesareansection scar endometrioma: a case report and review of the literature.

- J Radiol Case Rep. 2017;11(12):16-26. https://doi.org/10.3941/jrcr.v11i12.3178
- 7. Matter M, Schneider N, McKee T. Cystadenocarcinoma of the abdominal wall following caesarean section: case report and review of the literature. Gynecol Oncol. 2003;91(2):438-43. https://doi.org/10.1016/j.ygyno.2003.07.003
- 8. Boccara D, Runz AD, Marco O, Chaouat M, Mimoun M. Primary skin umbilical endometriosis: about one case. J Clin Case Rep. 2016;6(3):100742. https://doi.org/10.4172/2165-7920.1000742
- 9. Taniguchi F, Hirakawa E, Azuma Y, Uejima C, Ashida K, Harada T. Primary umbilical endometriosis: unusual and rare clinical presentation. Case Rep Obstet Gynecol. 2016;2016: 9302376. https://doi.org/10.1155/2016/9302376
- Andolf E, Thorsell M, Källén K. Caesarean section and risk for endometriosis: a prospective cohort study of Swedish registries. BJOG. 2013;120(9):1061-5. https://doi.org/10.1111/1471-0528.12236
- 11. Sengul I, Sengul D, Kahyaoglu S, Kahyaoglu I. Insizyonel endometriozis: üc olguluk bir sunu. Ulusal Cerrahi Kongresi 2008 (UCK 2008), Antalya, Turkey, May 28-31, 2008. Abstract no. GEN-15, p.123.

