Review Article

Endometriosis as a risk factor for colorectal cancer

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Abstract

Endometriosis is a common benign disease in women of reproductive age, it has been associated with an increased risk of various malignancies that is defined by certain histological criteria mainly 80% in ovary and 20% in extragonadal sites such as intestine, rectovaginal septum, abdominal wall, pleura and others; the greatest risk for colorectal cancer is women with adenomyosis or endometriosis; Several genetic alterations have been found in the risk of endometriosis associated with cancer; The symptomatology, imaging and endoscopic characteristics simulate other inflammatory and malignant lesions that make the preoperative diagnosis of extragonadal endometriosis difficult. This is a review of the knowledge about endometriosis and its potential risk of malignancy, particularly with colorectal cancer.

Background

Endometriosis is a proliferative disease that is defined as the presence of endometrial glands and stroma outside the uterine cavity; or in extraterine sites, it is a common chronic gynecological disease; the incidence in women of reproductive age is 5% to 17%, its cause is unknown; But, the accepted hypothesis is the implantation of endometrial tissue in the peritoneal cavity due to retrograde menstruation, or when endometrial tissues and cells adhere to the surfaces of the peritoneum, annexes and other pelvic organs [1-4]. The main symptoms are dysmenorrhea, pelvic pain and infertility. Although endometriosis is considered a benign condition, it shares some characteristics of cancer proliferation, such as invasion, tissue damage, neoangiogenesis and spread to distant organs [3].

Despite epidemiological evidence, the association between endometriosis and cancer has not been elucidated so far.

Endometriosis is hormonally dependent on estrogens, it is associated with oxidative stress, inflammatory pathways that are activated in its microenvironment [12,13], but, molecular events involved in malignant transformation are under investigation, and early molecular alterations have been identified such as the alteration of a tumor suppressor gene, the mutation of the ARID1A and loss of its encoded BAF250a [5]. The association between endometriosis and other hormone-dependent cancers particularly endometrial cancer (EC) and breast cancer (CM), share common risk factors (FR),
such as hyperestrogenism, some reproductive characteristics, obesity, the administration of menopausal hormone therapy (THM) and Type 2 diabetes mellitus (DM2) [13-15]. When clinical-pathological characteristics are compared between primary CE and synchronous epithelial ovarian cancer (EOC) [15], the incidence of endometriosis is higher in patients with EC than in EOC (100% vs. 35%), other malignant neoplasms such as colorectal cancer which is one of the most frequent intra-abdominal cancers in women that could exist in association with endometriosis. Uterine adenomyosis or internal endometriosis is when the ectopic endometrial glands and stroma are impregnated to the myometrium that can also be associated with an increased risk of cancer [3,14,15].

The clinical presentation of intestinal endometriosis is usually asymptomatic, or with gastrointestinal bleeding, nausea, vomiting, abdominal pain, defecation pain; diarrhea, constipation, rectum-vaginal colonic mass, intussusception, intestinal obstructions and intestinal perforation are observed; Symptomatology worsens during menstruation by 40%. Imaging and endoscopy of the intestinal tract simulates other inflammatory and malignant lesions; the definitive diagnosis before surgery is difficult, only clinical suspicion can prevent it [2], consider patients with endometriosis, especially postmenopausal patients with recurrence of symptoms [16-19] (Table 1).

**Endometriosis and cancer risk**

Women with endometriosis are at risk of developing cancer of 87.2 per 10,000 patients/year, or risk ratio of (OR) 1.8 with a 95% CI of 1.4 to 2.4; the time to develop endometriosis cancer was 34.3 months (18.7–46.8 months) and that is when they have adenomyosis; and increases with age, with OR 2.3 for those aged 31 to 40, 2.9; for 41 to 50 years and 4.2 for those 50 years of age or older [15-17].

**Imaging and pathology studies of intestinal endometriosis**

The diagnostic suspicion of intestinal endometriosis is mainly clinical, based on the symptomatology; but, the lack of pathognomonic signs makes diagnosis difficult; Even during surgery, intestinal endometriosis is confused with neoplasms, despite the use of computed tomography (CT) [1,18], Figure 2 or magnetic resonance imaging (MRI), the detection of wall nodules within the attached masses, when the characteristics Atypical sequences in MRI in T2 suggest a possible malignancy (Figure 3). In general it is diagnosed by histological findings after surgical resection [19] (Figures 4-6).

Pathological and immunohistochemical staining (IHQ) is essential to make a diagnosis, IHQ stains, which include CK 7, CK 20, vimentin and estrogen receptors (RE) [6] are useful to distinguish between adenocarcinoma arising from endometriosis and adenocarcinoma Primary intestinal [1,6]. The endometrioid glands are usually immunoreactive for CK7, RE, and stromal cells are positive for CD10 and RE.
intestinal glands express CDX2 and CK20, while showing a negative expression of CK7, RE or CD10. PAX8 was shown to be expressed in gynecological cancers [1].

**Discussion**

Endometriosis is defined histologically as the presence of endometrial glands and stroma outside the uterine cavity; common benign disease in women of reproductive age, the frequency of intestinal endometriosis varies from 3% to 34% [16], it affects the intestinal tract in 15% to 37% of patients with pelvic endometriosis [17]. Many theories about its pathogenesis have been proposed, the most accepted proposes retrograde menstruation and subsequent implantation of endometrial cells implanted in the peritoneum and pelvic viscera, which is facilitated by immune alterations; It is located in various anatomical sites, peritoneum, ovary, fallopian tubas, cervix, vagina, vulva, rectovaginal septum, uterus-sacral ligaments, intestine, rectosigmoid, bladder, uterus and skin [17].

Cancer develops in 5.5% of patients with endometriosis; 21.3% of cases originate in extragonadal pelvic sites, intestinal tumors associated with endometriosis are even more rare; in patients with pelvic endometriosis, it mainly affects the rectosigmoid colon, followed by the proximal colon, small intestine, blind and appendix. Malignant transformation of endometriosis without pelvic involvement is rare and its actual incidence is unknown; but, it simulates a neoplasm of the gastrointestinal tract [1,6], in a review of endometrioid adenocarcinomas that arise in colorectal endometriosis, of 50 cases only in 22 neoplastic transformation, it was an adenocarcinoma.

The others included sarcomas and mixed Müllerian tumors; progression to cancer has been linked to hyperestrogenism and to classify cancer as a result of endometriosis, it requires histopathological criteria; proposed by Sampson for the first time in 1925 and are: 1) presence of malignant and benign endometrial tissue in the same organ; 2) that the cancer arises from the tissue and does not invade it from another place; and 3) the finding of tissue similar to the endometrial stroma surrounding the characteristic glands [6].

The symptoms of intestinal endometriosis include abdominal pain, abdominal distension, signs and symptoms of gastrointestinal obstruction, rectorrhagia, etc., depending on the segment of the affected intestine, they can be cyclic in 40% that are usually aggravated during menstruation. Preoperative diagnosis of intestinal endometriosis through imaging is difficult and rare due to other common intestinal pathologies [20]. Treatments of intestinal endometriosis depend on presentation symptoms and operative findings. Medical hormone therapy with surgery provides less recurrence of endometriosis [20,21], it is rarely successful in severe symptomatic disease or intestinal obstruction, surgery with intestinal resection and anastomosis is necessary, Superficial lesions can be removed and definitive treatment is Excision is hysterectomy with bilateral salpingo-oporectomy and total excision of endometrial foci by laparoscopy or by laparotomy, with multidisciplinary team; it is associated with improvements in the quality of life, it represents a safe approach to the treatment of intestinal endometriosis or colorectal cancer; the risk of subsequent colorectal cancer was elevated in patients with coexisting adenomyosis with other extragonadal endometriosis, OR, 13.04, not including carcinoma in situ; of cancers related to endometriosis, colorectal is the second most common extragonadal site for malignant transformation of endometriosis [23,24] this malignant transformation is associated with hyperestrogenism [25], the theory of malignant endometrial transformation only explains a small proportion of patients with colorectal cancer; In women with coexisting adenomyosis, there are shared etiological factors for these 2 sequential events [21]. 80% of all neoplasms associated with extragonadal endometriosis occurred in the rectum and sigmoid colon, mostly are adenocarcinomas. Adenocarcinoma that arises from endometriosis often mimics primary intestinal adenocarcinoma [1].

The possible association between endometriosis and cancer by current molecular studies through routes related to inflammation, oxidative stress and hyperestrogenism [3,12] and alterations mutations of the tumor suppressor gene PTEN or ARID1A for malignant transformation [12].

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and the microenvironment of endometriosis and associated cancer share similar cytokines and mediators [23-25]. Epidemiological evidence supports molecular carcinogenesis with a link between adenomyosis and gynecological cancers, including colorectal cancer. Personalized treatment of women with endometriosis/adenomyosis is required through advice on early detection of cancer. There is no consensus on the therapeutic approach to treat malignant neoplasms associated with endometriosis, it is recommended that patients diagnosed with carcinomas associated with extragonadal endometriosis, which are limited to the lower pelvic cavity, can benefit from adjuvant pelvic radiotherapy, hormone therapy, may have a similar efficacy in malignant neoplasms associated with endometriosis with progesterone receptors [1].

Recently a new concept has been developed in the pathogenesis of endometriosis: the “neurological hypothesis”. showed an absolute correlation with the anatomic distribution of the pelvic sympathetic nervous system showed that there is a close histological relationship between endometriotic lesions of the large intestine and nerves in this area. Endometrioid lesions appear to infiltrate the wall of the large intestine preferably along the nerves, even at a distance from the palpated lesion. Endometriosis and its possible malignant changes should be taken into account in the differential diagnosis of intestinal masses in women. In addition, the clinical suspicion of malignancy should be aroused in patients with abdominal pain or rectal bleeding and a history of quiescent endometriosis [6].

Globally, cancer-associated endometriosis is very rare and it is really appropriate to consider it a premalignant condition; but, there is considerable controversy in the literature about the relationship between endometriosis and cancer [6] Due to the malignant potential, patients with endometriosis should have adequate hormonal management even after surgery and estrogens without opposition should generally be avoided in these patients [16,25].

Conclusion

Endometriosis is associated with an increased risk of cancer, including colorectal cancer mainly when coexisting with adenomyosis. Studies on the association of adenomyosis and the risk of colorectal cancer are needed to clarify whether the malignant transformation of colorectal endometrial implants.

References


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