

Research on central sensitization of endometriosis-associated pain: a systematic review of the literature

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Abstract: Endometriosis-associated pain afflicts an enormous number of women who suffer from endometriosis. There is an urgent need to explore the pathogenesis of endometriosis-associated pain to identify targets for treatment of hyperalgesia. A search was conducted in PubMed, Web of Science, Embase, and the Cochrane Library using the search terms “endometriosis” AND (“pain” OR “hyperalgesia” OR “nociception” OR “allodynia”) AND “central sensitization”. The search was limited to articles published in English from 01/01/2008 to the present. Among the search results, 15 articles were eligible for systematic review, including 6 reviews, 6 human studies (one in the form of a conference abstract only), and 3 animal studies. The articles were classified into 4 lists to describe the mechanism of endometriosis-associated pain and synthesize different aspects of research on it. In conclusion, there is a need to explore the mechanism of endometriosis-associated pain in terms of innervation, vascularization, local inflammation, cross-correlated visceral sensitization, and central sensitization to identify the target molecules and signaling pathways of key genes and relevant biomarkers through new techniques, all with the goal of developing a more comprehensive treatment strategy for endometriosis than is currently available.

Keywords: endometriosis-associated pain, neurogenic inflammation, mechanism, central sensitization

Introduction

Endometriosis, a condition in which lesions made of endometrium form ectopically outside the uterus, is a common gynecological disease in reproductive-age women. One in ten women has been diagnosed with this type of lesion. Recently, several common effects of endometriosis, including pain (found in 80% of patients), infertility (60%) and/or pelvic mass (40%),¹ have been classified as a syndrome. Studies have indicated that women with endometriosis-associated pain manifest several symptoms: chronic pelvic pain (CPP), dysmenorrhea, dyspareunia, and dyschezia. Additionally, substantial burdens can arise from associated nociceptive conditions such as viscerovisceral hyperalgesia syndrome, painful bladder syndrome (formerly called interstitial cystitis) and irritable bowel syndrome.²

The diagnosis of endometriosis depends on laparoscopy, in which the cyst or nodule can be seen directly in the peritoneal cavity. Laparoscopy is also the most effective way to eliminate the ectopic lesion. However, most patients still suffer from persistent pain after surgery. Some women with this disease can relieve pain

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via oral acyeterion, gonadotropin-releasing hormone agonist (GnRH-a), or nonsteroidal anti-inflammatory drugs.³ However, there remains a potential problem because clinically significant results require prolonged and repeated administration, which causes side effects in a large number of patients. Therefore, an imperative need exists for alternative, more mechanism-based treatments to ease the extreme hyperalgesic symptoms of endometriosis.

A growing body of evidence attests that patients with endometriosis endure pain associated with abnormal angiogenesis and the growth of novel nerve fibers in close proximity to ectopic lesions. Endometriotic lesions create an inflammatory environment and change the quality or quantity of inflammatory mediators or neurotransmitters, thereby stimulating peripheral nerve sensitization by remodeling the structure of peripheral synapses and accelerating conduction along nerve fibers. Berkely and McAllister⁴⁻⁶ discovered that ectopic cysts harvested from rat models with established endometriosis and from human patients develop their own C-fiber (sensory afferent) and sympathetic (autonomic efferent) nerve supply. The supply is rooted in nerve fibers innervating sites near the lesions; these fibers sprout branches into the growths. In 2003, Bajaj⁷ and his team proposed that central sensitization may be involved mechanistically in the development and maintenance of endometriosis-related pain. Those researchers hypothesized that persistent nociceptive input from endometriotic tissues might result in increased responsiveness among dorsal horn neurons processing input from the affected viscera and somatic tissues. Their subsequent study found a reduced pain threshold but improvement in the reaction to pricking and in mechanical hyperalgesia in 10 patients with laparoscopically confirmed endometriosis who suffered from pelvic pain. Spisak,⁸ by evoking blood-oxygen-level-dependent (BOLD) responses in a block-design functional magnetic resonance imaging (fMRI) experiment, identified that central sensitization to chronic pain can be observed as altered connectivity in key regions of the nociceptive network. An increasing number of studies focus on the relationship between differences in gene expression and the central sensitization mechanism of endometriosis-associated pain.

This review was organized with the aim of systematically synthesizing the literature published to date regarding the central sensitization mechanism of endometriosis-associated pain. The goal of this undertaking is to unearth the underlying pathogenesis and provide reliable

evidence to aid the search for novel methods of nonhormone target therapy for endometriosis-associated pain.

Methods

Literature search

Articles and review papers retrieved from the databases PubMed, Web of Science, Embase, and the Cochrane Library. Gray literature was excluded. The following search terms were used: “endometriosis” AND (“pain” OR “hyperalgesia” OR “nociception” OR “allodynia”) AND “central sensitization”. The search was limited to articles published in English from 01/01/2008 to the present.

Data selection

All studies identified by the searches were screened for inclusion. If our inclusion criteria were not all addressed in the abstract, then the methods section of the paper was screened. The inclusion criteria were as follows: (1) women or animals as the study subjects; and (2) women with endometriosis accompanied by pain, or animals with induced endometriosis-like lesions and hyperalgesia. The types of study designs included for review were randomized controlled trials (RCTs), cohort studies, cross-sectional studies, observational studies and reviews. Articles in languages other than English were excluded. Some potentially useful papers were excluded from the present meta-analysis because their data were too heterogeneous. The abstracts were double-checked by at least two authors to determine whether the reports fit the inclusion criteria for this study (Figure 1).

Data extraction

The articles found in the search were classified into the categories of reviews, research articles and conference presentations; furthermore, the research articles were subdivided into human research and animal research. For data collection and analysis, two authors independently extracted key data from the selected studies into several data tables according to the above classification. The tables contained general information such as author name, year of publication, locations and other characteristics (study characteristics, eligibility criteria, interventions, outcome measurements, etc.). All included articles were stored in EndNote software to assist the reviewers in managing data and to enable a third author to eliminate

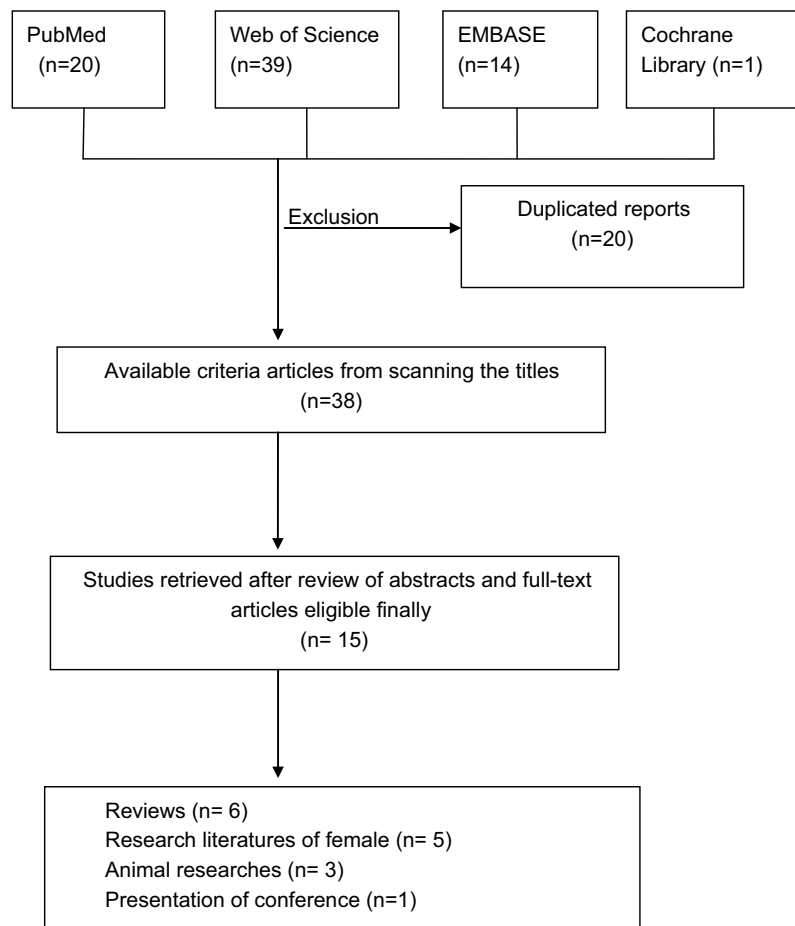


Figure 1 Flow chart of selection processes for eligible studies.

duplicate publications. All disagreements regarding study inclusion or data extraction were resolved by another author.

Results

Seventy-four publications were identified from PubMed, Web of Science, Embase and the Cochrane Library using the specified search terms. Twenty duplicates were removed from the list of search results, and the remaining studies were screened for inclusion and exclusion criteria by reading the abstracts and full texts. Ultimately, 15 articles were eligible for systematic review, including 6 reviews, 6 human studies (one of which was in the form of a conference abstract only), and 3 animal studies .

The 15 articles were classified into 4 categories in order to describe and synthesize information from similar study types. The human studies are summarized in Table 1; the animal studies are presented Table 2; the reviews are analyzed in Table 3, and the conference presentations are explained individually in Table 4.

Discussion

In recent years, studies on endometriosis-related pain have focused on CPP in women of reproductive age, but most patients with endometriosis have also shown some specific allodynia, such as abdominal myofascial pain syndrome,²⁴ visceral cross-organ nociception and muscle fascial pain;^{25,26} therefore, in recent years, these ambiguous pain locations have become a research hotspot for endometriosis-related pain. Most of the researchers have reviewed some original research articles accumulated over years the years, involving case-control, prospective, retrospective, randomized controlled and nonrandomized controlled study designs. Generally, pain patients who were diagnosed laparoscopically with endometriosis were assigned to an experimental group, while pain patients without endometriosis were classified in a control group, and pain assessment was performed with a visual analogue scale (VAS) scoring system. The pain patients received an intervention, such as medication (oral or intravenous) or acupuncture, sometimes accompanied by adjuvant

Table 1 The human studies for selected studies

Author	Rocha et al ⁹	Gurian et al ¹⁰	Deitos et al ¹¹	He et al ¹²	Chen et al ¹³
Year	2015	2015	2015	2010	2018
Locations	Brasil	Brasil	Brasil	China	USA
Participants	CPP=40 Group 1: endometriosis, n=24 (age: 32.9±9.0) ; Group 2: abdominal myofascial pain syndrome, n=16 (age: 32.7±7.8)	CPP=58 (age: 43.3) G1: endometriosis=12, G2: myofascial syndrome=18, G3: others=28	CTTH =30, MPS =29, FM =22, somatic/visceral nociception=27, endometriosis=32	Patients with endometriosis=100 (age: 34.4±7.4)	CPP (SFPN+) =25
N=?	Group3:a healthy, n=25 □age: 35.4 ±6.7)	Not given	pain-free controls n=37	Pain without endometriosis= 70 (age:33.4±7.1)	CPP (SFPN-) =14
Control N=?	prospective	prospective	prospective	prospective	retrospective
Typical study	prospective	prospective	prospective	prospective	retrospective
Pain measurement	VAS: G1=79.5±15.4, G2=85.9±11.0; threshold pain (kg/cm ²): G1=1.0±0.1 G2=1.9±0.2 G3=2.6±0.2	VAS=62; Basal pain threshold=14.2 electrical pain thresholds: G1=12.1; G2= 16.1	VAS>40mm (ie, moderate or severe pain). Pain associated with functional disability lasted >3 months.	VAS: G1=6.20 (1.74) G2=4.72 (1.71) MPQ: G1=11.48 (3.86) G2=9.19 (3.90) ST: G1=2.94 (0.83) G2=2.81 (0.85) PT: G1=11.25 (3.99) G2=13.60 (5.15)	LUTS, Vaginal or ovarian symptoms, Dyspareunia, Cystocele or vaginal mesh, Neurological symptoms, Depression, Substance dependence.
Intervention	Injections of 2 mL 0.5% lidocaine in G1 and G2	Oral or physical therapy for 6 months	Not intervention.	Surgery	Skin punch biopsies of the right upper thigh, using the PGP 9.5 and CD3 marker the tissues.
Parameter or Outcome	G1: VAS & NO:r=0.67 (95% CI: 0.35 to 0.85), P<0.0001; threshold & NO: r=-0.53 (-0.78 to -0.14), P<0.0001 G2: VAS & NO:r=-0.64 (-0.89 to 0.10), P=0.20; threshold & NO:r=-0.12 (-0.65 to 0.49), P=0.88	VAS=39; electrical pain thresholds: G1=16.5; G2= 19.3	Groups of patients with CSS presented higher expression: TNF- α =28.61±12.74pg/mL, BDNF=49.87±31.86ng/mL. Controls group: TNF- α =21.41±5.74pg/mL, BDNF=14.09±11.80ng/mL. BDNF: screen CSS from controls: AUC=0.86, cutoff=13.3 I ng/mL, sensitivity=95.06%, specificity=56.76%, moderate-severe depressive symptoms: AUC=0.81 cutoff=42.83ng/mL, sensitivity=56.80%, specificity=100%. TNF- α : moderate-severe depressive symptoms: AUC=0.97; cutoff=22.1 I pg/mL, sensitivity=90%, specificity=91.3%	VAS: G1=5.07 (1.35) G2=4.60 (1.67) MPQ: G1=8.93 (2.64) G2=8.90 (3.58) ST: G1=2.97 (0.67) G2:NG PT: G1=12.36 (3.69) G2:NG The score of VAS and MPQ is higher in patient group; The pain threshold is progressively improved in patient group.	A decrease in epidermal small fiber nerve density or small fiber loss in punch biopsy tissues of CCP patient with more comorbid conditions (SFPN+ group).

(Continued)

Table 1 (Continued).

Author	Rocha et al ⁹	Gurian et al ¹⁰	Deitos et al ¹¹	He et al ¹²	Chen et al ¹³
Conclusion	The plasma NO level may be directly involved in the pathophysiologic pain of central sensitization in women with endometriosis	Increasing the electrical pain threshold may provide an additional evidence for reducing the pain intensity between central sensitization and CPP.	BDNF, TNF- α as neuroplasticity mediators could play a vital role as screening tools for central sensitivity pain patients.	Central sensitization may be a possible mechanism underlying various forms of pain associated with endometriosis.	SFPN may supports a more dynamic relationship between the peripheral and central sensitization in complex CPP.

Abbreviations: G1, Group1, G2, Group2, G3, Group3; CPP, chronic pelvic pain; VAS, visual analogue scale; NO, nitric oxide; CSS, Central sensitivity syndrome; BDNF, brain derived neurotrophic factor; MPS, myofascial pain syndrome; CTTT, chronic tension-type headache; FM, fibromyalgia; MPQ, McGill Pain Questionnaire; ST, Sensory threshold; SFPN, small fiber polyneuropathy; LUTS, lower urinary tract symptoms.

therapy; subsequently, the effectiveness of the treatment was evaluated using a pain scale, such as the VAS, Iowa Pain Thermometer (IPT) or McGill Pain Questionnaire (MPQ),⁹ or a protein biomarker, such as BDNF, TNF- α , or the PEGylated form of a specific protein.^{11,27} Some studies have found that although the pain was relieved after the experimental treatment measures, it recurred after a few months; this phenomenon is related to the central sensitization mechanism of endometriosis.^{23,28–30} Nagabukuro³¹ has confirmed that endometriotic pelvic lesions contain abnormal neurovascular proliferation. These phenomena may be caused by abnormalities in conduction after peripheral sensitization, leading to central neurological abnormalities in the phenomenon known as central sensitization. The five articles cited above are the most comprehensive studies on the central sensitization mechanism of pain in endometriosis. Their subject populations were mainly from Brazil, China and the United States. This geographical scope is relatively limited and not representative of the entire world. Consequently, the results of the analysis are applicable only to the region, and the quality of evidence needs to be improved.

In three studies of animal models of the central sensitization mechanism of endometriosis-associated pain, pain intensity was evaluated in mice by behavioral experiments, for instance, the hot plate test, von Frey filaments, and the open field test, with or without drug intervention. Subsequently, the brains of the mice were dissected in search of target proteins or genes in candidate regions via staining and sequencing. Li and Greaves^{14,15} found changes in functional regions of the genome through sequencing of candidate regions, and they also detected abnormal expression of some genes as a result of central sensitization. Vicuna et al³² identified that the expression of Serpina3n and Lct were downregulated in the insula of endometriotic mice, which may play a vital role in central sensitization to endometriosis-associated pain of central sensitization. Hence, they assumed that an intervention against central sensitization would relieve the symptoms of pain by altering these abnormally regulated genes. This hypothesis, however, is based on animal experiments and is not mature enough for clinical application. There is insufficient evidence to conclude that patients with endometriosis-related pain have genetic changes similar to those in animal models; therefore, this conclusion cannot be extended to humans at present. However, preclinical animal exploration provides an experimental basis for future clinical research.

Table 2 Animal studies for selected studies

Author	Li et al ¹⁴	Greaves et al ¹⁵	Dodds ¹⁶
Year	2018	2017	2018
Location	USA	UK	Australia
Species	Female C57BL/6 mice	Female C57BL/6 mice	Female C57BL/6 mice
ENDO, N=?	12	18	5 (endometrial fragments from 5 donor mice were injected into abdomen cavity)
Control, N=?	Sham =12	OVX + E2=6 OVX + E2+PBS=6 Sham =6	6 (sterile saline were injected into abdomen cavity)
Pain measurement	Hot plate test/Open field test/Tail suspension test	Open field test/abdominally directed licking/Von frey test	No measurement
Intervention	-	Injection the inhibitor of TRPV1, JNJ 17203212, EP4 antagonist L-161982, the EP2 antagonist TG6-10-1 and EP2 antagonist (PF-04418948) in all groups.	ENDO: endometrial fragments from 5 donor mice were injected into abdomen cavity. Control: sterile saline were injected into abdomen cavity.
Parameter or Outcome	<ul style="list-style-type: none"> Altered CNS electrophysiology: pain, anxiety, and depression result from impairment in GABAergic and glutamatergic transmission onto neurons in the amygdale. Differentially expressed genes in the brains: upregulated(Gpr88, Glra3 in insula; Chrb4, Npas4 in the hippocampus; Lcn2 in the amygdala); downregulated(Lct, Serpina3n in insula; Nptx2 in amygdala). 	<ul style="list-style-type: none"> EP2, EP4, COX-1, COX-2, PGE2, increased in endometriosis lesions. EP2, Cox-1, Scn11a and Trpv1 mRNA concentrations were increased in DRG. The EP2 antagonism could reverse both peripheral and secondary hyperalgesia. 	<ul style="list-style-type: none"> Astrocytic GFAP and microglial CD11b were highly expressed in immunoreactions of Spinal cords (T13-S1) as endometriosis-like lesions.
Conclusion	Gene abnormal expression in brain result from the mechanism of central sensitization, which provide evidence for molecular targets to cure pain.	EP2 receptor antagonism could be a key target for the potential therapies of endometriosis-associated pain.	Endometriosis-like lesions resulted in the adaptations in nonneuronal, immune-like cells of the central nervous system to modulate central sensitization and pain.

Abbreviations: EMS, induced- endometriosis by surgery; Sham, Sham surgeries for controls; OVX, ovariectomised; COX-1, *cyclooxygenase-1*; COX-2, *cyclooxygenase-2*; DRG, dorsal root ganglia; GFAP, glial fibrillary acidic protein.

Endometriosis-related pain is currently considered a form of neuropathic or neuroinflammatory pain. A large number of experimental studies, both human and animal, have demonstrated that abnormal microscopic neurogenesis and angiogenesis occur in ectopic lesions, supporting the development, density, infiltration and even metastasis of ectopic endometrium.^{4,17,20,33} The perception of pain is caused by this abnormal proliferation of nerve fibers and vessels. Inflammatory factors are released into the sensory afferent nerve at the distal end of the lesion by noxious stimuli, and the nociceptive signal is transferred to the nerve root of the dorsal horn of the spinal cord. After simple handling, the pain signal is transferred to the thalamus, the brain stem, and finally the cerebral cortex.³⁴ If the lesion sites are stimulated persistently by those abnormal factors, the transmission of

pain signals will change, magnifying future pain and forming more intense memories of pain in the cerebral cortex. During the process of signal transduction, the function of the corresponding immune cells and cytokines changes accordingly, promoting the enhancement and amplification of the pain signals to some extent.³⁵ As-Sanie³⁶ and his colleagues determined that women with endometriosis-associated CPP shown increased levels of combined glutamine and glutamate (Glx) within the anterior insula and increased anterior insula connectivity to the medial prefrontal cortex (mPFC), which may play a role in the pathophysiology of CPP independent of the presence of endometriosis. Of course, the degree of pain is positively correlated with the density of nerves at the lesion. The higher the density of nerve fibers, the more pronounced the pain. Reviews by Liu³⁷ and

Table 3 Reviews for selected studies

Author	Year	Location	Personality	Common
Asante et al ¹⁷	2011	USA	<ul style="list-style-type: none"> • Circulating Markers of Endometriosis: CA-125, ICAM-1. • Central sensitization was induced in dorsal horn neurons by an increasing in excitatory synaptic transmission, mediated via the NMDA and AMPA receptors, or by a loss of inhibitory synaptic transmission, mediated via GABA and glycine receptors . 	<ul style="list-style-type: none"> • Endometriosis-related pain is a type of neuropathic pain or a neurogenic inflammation pain. • unique vascular and neural supplies via neuroangiogenesis were identified in location lesions. • Autonomic Nervous System Changes and defective immunosurveillance and inflammatory hyperresponsiveness. • Patients with DIE and bowel endometriosis show more several pain symptoms cause of a higher nerve fibres density compared to other sites.
Brawn et al ¹⁸	2014	UK	<ul style="list-style-type: none"> • Changes in brain structure: reduction in brain volume by neuroimaging. • Activity of the HPA axis. • Predisposition to other chronic conditions. 	
Laux-Biehlmann et al ¹⁹	2015	Germany	<ul style="list-style-type: none"> • Noxious and innocuous stimuli in endometriosis lesions cause inflammatory pain. • Escherichia coli found in menstrual blood and peritoneal fluid validated that the DAMPs and PAMPs play a vital role in endometriosis-associated inflammation. • TRPV1-positive nerve fibers is relevant to CPP. • Nervous system response further increases peritoneal inflammation result from higher concentration of SP and CGRP in lesions, whilst NK1R gene polymorphism rs881 may be a pathogenic role in endometriosis. 	
McKinnon et al ²⁰	2015	USA	<ul style="list-style-type: none"> • Analyze the implication between endometriosis and the molecules in peritoneal fluid or neurogenic inflammation environment including: ENA-78, IL-1β, IL-6, IP-10, IL-33, Leptin, MCP1, MK, NGF, OPG, PAEP, PAAPP-A, RANTES and TNFα. 	
Morotti et al ²¹	2017	UK	<ul style="list-style-type: none"> • Offer the precise terminology on assessment of pain. • TRPV1, CCL2, BDNF, VEGF, and NT4/5 in cysts are highly expressed. • Women with endometriosis-associated CPP were detected several volumetric modifications in specific brain areas, such as thalamus, insula, putamen, etc. • HPA axis, the endocrine pathway, present a dysfunction by central change, demonstrated a positive correlation between cortisol reductions and both infertility and dyspareunia. 	
Aredo et al ²²	2017	USA	<ul style="list-style-type: none"> • A framework for evaluating such sensitization and myofascial trigger points in a clinical setting is presented. • Painful MTrPs may serve as an additional source of nociceptive input. • Botulinum Toxin Type A may obtain satisfactory therapy effect in alleviating the myofascial pelvic pain associated with endometriosis via blocking the transmission of signals that stimulate muscle fibers. 	

Abbreviations: CA-125, cancer antigen-125 or carbohydrate antigen-125; ICAM-1, serum-soluble intercellular adhesion molecule-1; CPP, chronic pelvic pain; HPA, Hypothalamic–Pituitary–Adrenal; NMDA, glutamate N-methyl-D-aspartate; AMPA, α -amino-3-hydroxy-5-methyl-4-isoxazole-propionate; GABA, γ -aminobutyric acid; DAMPs, damage-associated molecular patterns; PAMPs, pathogen-associated molecular patterns; TRPV1, transient receptor potential vanilloid 1; SP, neuropeptide substance P; CGRP, calcitonin gene-related peptide; NK1R, neurokinin 1 receptor, is one of the SP receptor; DIE, deep infiltrating endometriosis.

Table 4 the presentation at conference

Author	Year	Location	Hypothesis	Methods	Results	Conclusions
Guo et al ²³	2009	China	women with endometriosis have increased pain perception as compared with women without.	VDS, VAS and IPT (VAS and MPQ scores) assess the their severity of dysmenorrheal between endometriosis and without endometriosis.	Women with endometriosis had a significantly higher VAS and MPQ scores than without endometriosis.	Central sensitization may well be a possible mechanism for various types of pains associated with endometriosis, may underlie both pathological and adaptive functions in the affected visceral areas.

Abbreviations: VDS, verbal descriptor scale; VAS, visual analogue scale; IPT, ischemic pain test; MPQ, McGill Pain Questionnaire.

Serrano³⁸ hold that deep infiltrating endometriosis (DIE) and ectopic growths on the intestinal wall have greater neurological density than ectopic lesions in other parts, meaning that this type of patient will have more severe pain.

The exploration of biomarkers has played an important part in research efforts to characterize the mechanism of endometriosis pain. CA125 has been widely used in the detection of endometriosis, but no study, to our knowledge, demonstrates clearly that the increase in CA125 is positively correlated with the degree of pain. Existing research merely indicates that the amount of CA125 in patients with endometriosis is elevated, which can serve as supplemental diagnostic sign.^{17,39} Some studies also suggest that ICAM-1 is associated with the degree of pain in endometriosis.¹⁷ A review by Brawn¹⁸ states that chronic, repeated local pain stimuli affect the normal activity of the hypothalamic–pituitary–adrenal (HPA) axis and may then exacerbate pain through a reduction in cortisol levels. Meanwhile, the structure of specific brain areas is changed, such as the periaqueductal gray (PAG), a vital region in the descending pain modulatory pathways;⁴⁰ the volume of the PAG is increased in women with endometriosis pain compared to those without pain, as observed by seed-based resting functional connectivity magnetic resonance imaging (fcMRI). Laux-Biehlmann¹⁹ summarized the important role that damage-associated molecular patterns (DAMPs) and pathogen-associated molecular patterns (PAMPs) play in pain signaling from ectopic lesions. At the same time, it is also recognized that TRPV1, substance P (SP) and calcitonin gene-related peptide (CGRP) show high expression in lesions and nearby sites that are subject to pelvic pain. On a related note, the NK1R gene polymorphism rs881 may play an important role in this process, providing a new possibility for pain treatment targets, but further

verification is needed. From the review of McKinnon,²⁰ it is not difficult to see that many other inflammatory factors such as Leptin, MCP1, MK, NGF, OPG, PAEP, PAAPP-A, RANTES and TNF- α are relevant to the pain associated with endometriosis, which merits further attention in the field of inflammatory pain. Morotti's view is similar to that of Brawn and further proposes that volumetric changes in the thalamus, insula and putamen result from long-term endometriosis-related pain.²¹ Aredo²² took another perspective to explain endometriosis-related pain at sites such as myofascial trigger points (MTrPs); he noted that botulinum toxin type A can alleviate myofascial pain by blocking signal transmission.

A study presented by Guo et al²³ also used questionnaires to evaluate the symptoms of CPP in patients. The investigators conducted a brief analysis of the underlying cause of intractable endometriosis-associated pain driven by central sensitization.

In summary, we can conclude that endometriosis-associated pain is closely related to central sensitization, as validated in both animal experiments and human case-control studies. Attention should be focused on the molecular pathways of pain signaling, with the intention of identifying the target-molecule signaling pathways of key genes and relevant biomarkers through new techniques, blocking the transmission of amplified signals by effective methods, and blocking or reversing the results of central sensitization from neurotoxic stimulation in order to relieve pain and improve quality of life in patients suffering from endometriosis-associated pain.

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Disclosure

The authors report no conflicts of interest in this work.

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