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[Intervention Review]

# Microwave therapy for cervical ectropion

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## ABSTRACT

### Background

In western medicine, cervical ectropion is considered as a normal finding that does not require treatment despite the red and inflamed appearance of the cervix. However, cervical ectropion is considered as one of the most common types of chronic cervicitis in China. Topical treatments for cervical ectropion, including microwave tissue coagulation, are widely used in many hospitals in China they are considered likely to reduce the chances of development of abnormal metaplasia and infection.

### Objectives

To compare the efficacy and potential side effects of microwave tissue coagulation with other interventions or no intervention in the treatment of cervical ectropion.

### Search methods

We searched The Cochrane Library, MEDLINE, EMBASE, Chinese Biomedical Literature Database (CBM), Chinese Medical Current Contents (CMCC), CAJ Full-text Database and the Chinese Scientific Journals Database. We also searched for related literature on the Internet with search engines such as Google and searched the reference lists of articles and handsearched relevant Chinese journals. The search of these databases was conducted from inception to May 2009.

### Selection criteria

Only authentic randomised controlled trials (RCTs) were considered for inclusion.

### Data collection and analysis

Two review authors independently interviewed the original authors of claimed RCTs published in China and then assessed the risk of bias of three RCTs and extracted data.

### Main results

No studies were found that met the participant inclusion criteria of the protocol. One hundred and thirty-three studies were reported as RCTs, but only only three studies were identified as authentic RCTs. However, it was not possible to confirm that the participants were

symptomatic prior to treatment. and the methodological quality of the three RCTs was generally low. No trial compared treatment to no intervention and most trials didn't assess relief of symptoms or quality of life and satisfaction, which are very important to women.

### **Authors' conclusions**

There were no RCTs in symptomatic women with cervical ectropion comparing microwave therapy with another treatment or no treatment. Although microwave therapy improved the appearance of the cervix compared with both laser and interferon-alpha suppository therapy, any other benefit for the women was not clear.

## **PLAIN LANGUAGE SUMMARY**

### **Microwave therapy for cervical ectropion**

In western countries, cervical ectropion or erosion caused by the movement of columnar epithelium onto the vaginal portion of the cervix is considered a normal physiological process not requiring intervention. In China, however, cervical ectropion is considered as one of the most common types of chronic cervicitis and is often treated. One of the topical treatments widely used is microwave tissue coagulation (MTC). From Chinese databases, the authors identified 131 potential randomised controlled clinical trials (RCTs) that compared microwave therapy to no treatment or other treatments. Only three of these studies involving a total of 540 participants were identified as RCTs. Two studies (420 participants) compared MTC versus laser, and one study (120 participants) compared MTC with an interferon-alpha suppository. The follow-up period of the RCTs was adequate. None of the studies assessed relief of symptoms or quality of life and satisfaction, which are very important to women. The authors of the review considered that they could not answer the review question about the appropriate use of microwave therapy as only these three low quality RCTs were found. Although the trials showed improved appearance of the cervix with microwave treatment compared with the control therapy, it was not possible to ascertain whether the women were symptomatic before treatment. The review protocol also required participants to be symptomatic with mucopurulent discharge or to have contact bleeding before treatment. At present, intervention for inflammatory cervical ectropion is still controversial.

## BACKGROUND

### Description of the condition

The term cervical ectropion (CE) has replaced that of cervical erosion. It is also sometimes referred to as the transformation zone of the cervix. A cervical ectropion occurs when, due to hormonal changes, the columnar epithelium moves onto the vaginal portion of the cervix. A cervical ectropion appears red and may have an inflamed appearance due to the columnar epithelium being thinner than squamous epithelium. This makes the underlying blood vessels more apparent. Along with the red appearance, the columnar epithelium may also secrete mucus which, if abundant, causes a vaginal discharge (Chang 1991). In western countries, cervical ectropion is considered as a physiological process not requiring treatment. In China, however, cervical ectropion is considered as one of the most common types of chronic cervicitis and it is thought that severe cervical ectropion may cause infertility (Daljic 2001). As a result, treatment for cervical ectropion is widely practiced in hospitals in China. It is reported that 20% to 50% of Chinese women of reproductive age have cervical ectropion (Zhang 1995; Cai 1997; Peng 2002; Xiao 2003; Zhang 2003).

In the Chinese medical literature, the occurrence of cervical ectropion is not considered to be caused solely by estrogenic hormonal changes; inflammation and trauma have also been implicated in its etiology (Yue 2000). Inflammatory ectropion presents as a defect or degeneration and necrosis of cervical squamous epithelium combined with infiltration of large numbers of inflammatory cells as evidenced using histopathology. On a smear, superficial, intermediate and basement cells can be found with degenerated and necrotic cells, mild dyskaryotic cells and large numbers of neutrophils; small numbers of monocytes, lymphocytes, plasmocytes and macrophages can also be seen (Tang 1995).

Cervical ectropion has been associated with both the combined oral contraceptive pill and chlamydia trachomatis (Critchlow 1995). Changing the method of contraception may be considered for those women who are unable to accept the increase in physiological discharge associated with cervical ectropion. Screening for chlamydia and other sexually transmitted infections should be considered for those women who are at risk. Ectropion has also been found in 5% to 25% of women presenting with postcoital bleeding (Selo-Ojeme 2004). These women need appropriate follow up with a cervical smear and colposcopy.

Clinical macroscopic examination describes three grades (I, II and III) based on size, the severity of the lesion and the rate of growth of columnar epithelium. These are described as simple pattern, granular pattern and mammilla pattern (Yue 2005). Macroscopically it may be difficult to distinguish cervical ectropion from cervical intraepithelial neoplasia (CIN), cancer in situ. Thus a cervical smear for cytological examination should be routinely performed to exclude dysplasia. In addition, colposcopy and a biopsy may also be needed for histopathologic examination.

### Description of the intervention

Microwave therapy uses electromagnetic waves at a frequency of up to 2.45 billion Hz. The biophysical effect at the target site is through both an internal heating effect and a non-heating effect. Unlike the external heating of other physical treatments,

microwave tissue coagulation (MTC) involves internal heating using human tissue as the heat source (Guan 2004).

Treatments are only performed after excluding malignancy and specific or non-specific cervical or vaginal infections. The purpose of treatment is to induce necrosis and exfoliation of the columnar epithelium at the affected site and allow repair by neofornative squamous epithelium. This, in turn, is considered likely to reduce the chances of development of abnormal metaplasia and infection.

### How the intervention might work

When the microwave electrode touches the topical lesion, a microwave magnetic field forms in the tissue and the small-scope high temperature produced instantly, which induces degeneration, coagulation and necrosis of the topical tissue protein. New squamous cells then migrate from peripheral tissue and repair the wound. The general non-heating effect of microwave induces dilation of superficial blood vessels, acceleration of blood flow and enhancement of tissue metabolism, which in turn promote the elimination of inflammatory products. There is, therefore, a synergistic action within the therapy (Guan 2004).

### Why it is important to do this review

Presently, the rationale for use of microwave therapy for asymptomatic cervical ectropion is unclear.

Although this systematic review was first published in the Cochrane Database of Systematic Review in 2007, many methods, including microwave treatment, are still used to treat cervical ectropion in China. It is necessary to evaluate these interventions.

## OBJECTIVES

To compare the efficacy and the potential side effects of microwave tissue coagulation with other interventions or no intervention in the treatment of cervical ectropion.

## METHODS

### Criteria for considering studies for this review

#### Types of studies

Only authentic randomised controlled trials (RCTs) were included in the review.

#### Types of participants

Women aged 18 to 60 years with cervical ectropion who were symptomatic with mucopurulent vaginal discharge or contact bleeding, or both. The women were diagnosed by vaginal examination and graded by ectropion size and tissue pattern.

#### Diagnostic criteria

Macroscopic examination to discriminate between acute and chronic cervicitis.

Three grades were based on ectropion size: grade I, covering one third of the cervix; grade II, from one third to two thirds; grade III, over two thirds of the cervix.

Three patterns were defined based on the severity of the lesion and the growth of columnar epithelium: simple pattern, granular pattern and mammilla pattern. A simple pattern was

where the surface of the ectropion was planus and covered by simple columnar epithelium; the granular pattern was where the surface was not smooth and the glandular epithelium was hyperplasmic, accompanied by hyperplasmic mesenchyma; the mammilla pattern was where the surface was more severely uneven.

### Exclusion criteria

- Asymptomatic ectropion.
- Women who were pregnant or post-menopausal
- Women with cervical intraepithelial neoplasia (CIN1, 2, 3), cancer in situ, or early cervical cancer diagnosed through a Papanicolaou test, histological examination or colposcopy.
- Women with vaginal or cervical infection (vulvovaginal candidiasis, bacterial vaginosis, trichomonas vaginitis, chlamydial or gonococcal infection).

### Types of interventions

Microwave tissue coagulation (MTC) versus no treatment or another treatment such as laser coagulation, infrared light, an interferon-alpha suppository (Aoping suppository), the combination of MTC with an interferon-alpha suppository, electrocautery or cryotherapy.

### Types of outcome measures

#### Primary outcomes

- 1) Relief of symptoms - vaginal discharge, contact bleeding as assessed by the woman or health professional, or both.
- 2) Quality of life or satisfaction.
- 3) Appearance of the cervix.

#### Secondary outcomes

##### Assessment criterion of curative effect

- 1) Healing well: cervix had become smooth and glossy, ectropion had disappeared or lesion on cervical surface had become squamous metaplasia with the new squamous epithelium coloured on iodine testing.
- 2) Not healing well: reduction in the grade of cervix ectropion or no change after treatment.

##### Additional outcome measure

- 1) The rate of bleeding during operation.

##### Follow-up period

For measurement of cervical ectropion, a follow-up period of 8 to 12 weeks was considered adequate.

For symptoms of vaginal discharge and contact bleeding (and quality of life, satisfaction), a follow-up period of more than three months was considered adequate.

### Search methods for identification of studies

#### Electronic searches

We searched the Cochrane Menstrual Disorders and Subfertility Group Trials Register, Cochrane Central Register of Controlled Trials (CENTRAL) (*The Cochrane Library* 2009, Issue 2), MEDLINE (January 1966 to May 2009), EMBASE (January 1974 to May 2009), Chinese Biomedical Literature Database (CBM) (January 1978 to May 2009), The Chinese Medical Current Contents (CMCC) (January 1994 to May

2009), CAJ Full-text Database (CNKI) (January 1994 to May 2009), and Chinese Scientific Journals Database (VIP) (January 1989 to May 2009). No language restrictions were made. We used the search strategies detailed in [Appendix 1](#); [Appendix 2](#); and [Appendix 3](#).

#### Searching other resources

We also searched for related literature on the Internet with search engines such as Google (30 May 2009), the reference lists of articles and handsearched relevant Chinese journals that were most likely to publish trials on cervical ectropion. We made attempts to contact some experts and manufacturers but did not identify any related studies in this way.

### Data collection and analysis

#### Selection of studies

We developed a standardized data extraction form and performed a pilot to ensure consistency and completeness. Two review authors (LYL, ZP) independently selected the potential studies for inclusion. Two review authors telephone interviewed the authors of claimed RCTs and considered trials for inclusion using a specially designed questionnaire (see Additional [Table 1](#)).

#### Data extraction and management

It was planned for two review authors (TJ, MB) to independently enter data into a data extraction form. Discrepancies would be resolved by a third review author (LYL). We obtained missing data from authors wherever possible. Data were to be checked and entered into RevMan by one review author (LYL). However, in the event there were no eligible studies.

#### Assessment of risk of bias in included studies

We assessed the risk of bias of each trial in terms of adequate sequence generation, allocation concealment, blinding, that incomplete outcome data were addressed, the trial was free of selective reporting and free of other biases. These were classified as 'Yes' for low risk of bias; 'No' for high risk of bias; or 'Unclear' for moderate risk of bias, according to the guidelines of the Cochrane Handbook of Systematic Reviews of Interventions 5.0.1 and described by [Wu 2007](#). Sensitivity analyses were to be undertaken on the basis of whether the quality factors were 'Yes', 'No' or 'Unclear'. Differences were to be resolved by discussion among the review authors.

The following characteristics were assessed.

#### Sequence generation

(1) Yes, adequate sequence generation was reported using one of the following approaches: random number tables, computer-generated random numbers.

(2) No, other methods of allocation used that appeared to be biased.

(3) Unclear, did not specify one of the adequate methods outlined in (1) but mentioned 'random'.

#### Allocation sequence concealment

(1) Yes, adequate measures to conceal allocations with the key being that the person who generated the allocation sequence did not recruit or allocate the participants, such as central randomisation; serially-numbered, opaque, sealed envelopes;

or another description that contained convincing elements of concealment.

(2) No, inadequately concealed allocation where the description reported an approach that did not fall into one of the categories in (1). Did not conceal allocation.

(3) Unclear, unclearly concealed trials in which the author did not report an allocation concealment approach at all.

### **Blinding**

(1) Blinding of participants (Yes, No or Unclear).

(2) Blinding of caregivers (Yes, No or Unclear).

(3) Blinding of outcome assessment (Yes, No or Unclear).

### **Incomplete outcome data addressed**

(1) Trials where an intention-to-treat analysis was possible and few participants were lost to follow up.

(2) Trials which reported exclusions, as listed in (1), but exclusions were less than 10%.

(3) No reporting on exclusions, exclusions of at least 10%, or wide differences in exclusions between groups.

### **Free of selective reporting**

(1) Yes, if all the important outcomes (the primary outcomes stated in the review) were reported or if the trial protocol was available and all of the trial's pre-specified (primary and secondary) outcomes that were of interest to this review were reported.

(2) No, if not all the pre-specified outcomes were reported, the primary outcomes were changed, or if some of the important outcomes were incompletely reported.

(3) Unclear, if there is insufficient information to assess whether the risk of selective outcome reporting was present.

### **Free of other bias**

Source of funding:

(1) Yes, if the trial was unfunded;

(2) No, if the trial was funded;

(3) Unclear, if the source of funding was not clear.

Baseline imbalance:

(1) Yes, if there was no baseline imbalance in important characteristics;

(2) No, if there was a baseline imbalance due to chance or due to imbalanced exclusion after randomisation;

(3) Unclear, if the baseline characteristics were not reported.

### **Measures of treatment effect**

For dichotomous data, two-by-two tables were generated for each trial and the data expressed as a risk ratio (RR) with 95% confidence intervals (CI), for example cure rate and rate of bleeding during operation.

### **Unit of analysis issues**

The included trials were to have a simple parallel group design where participants were individually randomised to treatment and control groups.

### **Dealing with missing data**

We contacted the original authors to request missing data. Then we would discuss why data may be missing.

### **Assessment of heterogeneity**

We were to assess statistical heterogeneity (variation) between the results of different studies included in the meta-analysis by inspecting the scatter in the data points on the graph and the overlap in their confidence intervals. The Chi<sup>2</sup> test was used with significance set at a P value less than 0.1 and the I<sup>2</sup> statistic to estimate the total variation across studies due to heterogeneity, where I<sup>2</sup> less than 25% was considered as low-level heterogeneity, 25% to 50% as moderate level and higher than 50% as high-level heterogeneity (Higgins 2003).

### **Assessment of reporting biases**

Potential publication bias was not tested by a funnel plot because there were not sufficient trials.

### **Data synthesis**

It was planned to analyse abstracted data following the guidelines of the Cochrane Handbook for Systematic Reviews of Interventions (Higgins 2009) and to use RevMan software to complete the data analysis. The results of individual studies would be pooled only if they were clinically similar with respect to the participants, interventions and clinical criteria for defining outcomes.

Published graphs were to display the results using the fixed-effect approach unless there was statistical heterogeneity, in which case the random-effects model was to be used. However, unless the results are robust to both models, they would need to be treated with caution. The results may be converted to absolute measures, such as the number needed to treat (NNT). Continuous data were to be expressed as mean differences (MD) and 95% confidence intervals. Meta-analytic methods for continuous data assume that the underlying distribution of the measurements is normal. Results were also given as reported in the publication, for example as median and range with non-parametric tests of significance.

### **Subgroup analysis and investigation of heterogeneity**

It was planned to perform subgroup analysis according to different interventions and the grades of cervical ectropion.

### **Sensitivity analysis**

We planned to conduct a sensitivity analysis and to explore the effect of including only trials with adequate methodology.

## **RESULTS**

### **Description of studies**

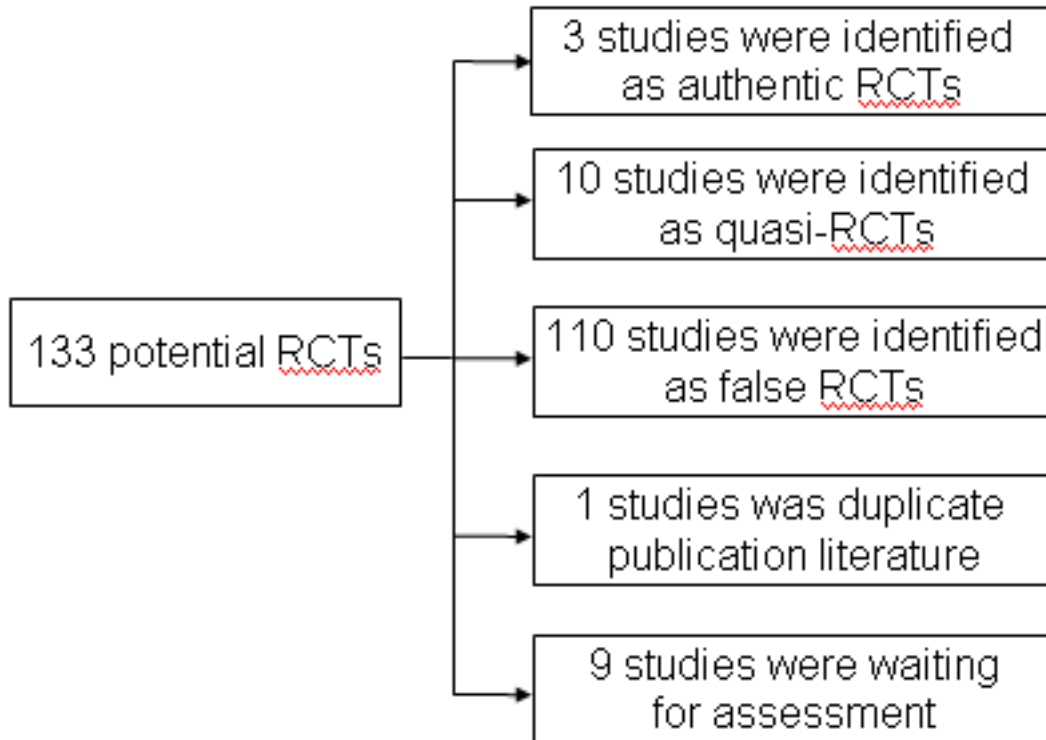
#### **Results of the search**

No English language studies were identified in *The Cochrane Library*, MEDLINE or EMBASE. One hundred and thirty-three

potentially eligible Chinese trials were identified using the described search strategy and inclusion criteria. We tried our best to contact the authors of claimed RCTs to get missing information

about randomisation and other data. No unpublished studies, grey literature or ongoing trials were found (see [Figure 1](#)).

**Figure 1. Flow chart of the study excluded process**



**Included studies**

No studies were available for review.

**Excluded studies**

One hundred and twenty-four studies were excluded ([Characteristics of excluded studies](#)). Ten studies were identified as quasi-randomised controlled trials from the description of the authors; 105 were identified as non-RCTs and excluded. The others were excluded as there was no documentation on whether the women who had cervical ectropion were symptomatic. One was a duplicate publication. Nine studies were not assessed because we were unable to access the full-text versions or failed to contact the authors. Finally, only three studies were identified as true randomised controlled trials. The three trials were single centre and had a parallel group design. However, it was not possible to ascertain whether any of these three randomised studies met the participant inclusion or exclusion criteria as there was no documentation as to whether the women who had cervical ectropion were symptomatic. We have presented the findings of these three RCTs in Additional tables (see Additional [Table 2](#); [Table 3](#); [Table 4](#)).

**Risk of bias in included studies**

No studies were available for review. We assessed the risk of bias of the excluded RCTs.

**Allocation**

None

**Blinding**

None

**Incomplete outcome data**

None

**Selective reporting**

None

**Other potential sources of bias**

None

**Effects of interventions**

No studies were found that met the participant inclusion criteria of the protocol. Although three studies were identified as authentic RCTs, from 131 potential RCTs, it was not possible to confirm that the participants were symptomatic prior to treatment.



## DISCUSSION

### Summary of main results

In China, cervical ectropion is considered as one of the most common types of chronic cervicitis and requires treatment. Three randomised studies involving 540 participants were found. Two studies (Song 1994; Xu 2004) compared MTC versus laser (420 participants) and one study (Wang 2003) compared MTC versus interferon-alpha suppository. Relief of symptoms, quality of life and satisfaction were not reported in any of the three studies. The study results suggested that microwave therapy gave similar or greater improvements in the appearance of the cervix over laser or interferon suppository use. One study reported the rate of bleeding during operation (Song 1994). The difference was statistically significant (RR 0.15, 95% CI 0.06 to 0.37; 300 participants, one trial) between the MTC and laser groups. Other adverse effects such as bleeding and topical infection after operation were also reported in Song 1994. No differences were found between the two groups. The other study (Xu 2004) reported a statistically significant difference for the duration of vaginal discharge post-treatment (mean difference -3.30, 95% CI -5.27 to -1.33; 120 participants, one trial).

Wang 2003 reported that bleeding after operation occurred in a few participants with no difference between MTC and an interferon-alpha suppository (120 participants, one trial).

However, none of these trials employed assessor blinding, which raises the possibility of assessor bias. In addition, there were no data to clarify whether these women were symptomatic before treatment or received any benefit from the treatment in addition to improved cervical appearance. No study compared microwave therapy with no treatment. This is important as the rationale for use of this therapy for cervical ectropion is not clear.

### Overall completeness and applicability of evidence

Up to now, no clearly applicable evidence has been found to show any benefit from MTC in the treatment of cervical ectropion.

### Quality of the evidence

The information on quality issues for the three RCTs was poorly reported. Although we were finally able to access this information, from the original authors, the risk of recall bias is high. There were also varying standards of classification for the grade and subtype of cervical ectropion. Consideration of sample sizes was not reported, which may lead to a low-test power. We found that most Chinese authors misunderstood the concept of randomisation. One hundred and thirty-three studies were reported as RCTs, however only three were authentic RCTs (2.3%, 3/131). Some of studies were experience collections or case collections and most of them were written retrospectively.

The quality of the three RCTs was poor (see Additional Table 5).

There was not enough information in the trial reports to determine whether the allocation sequence was adequately concealed. We contacted the authors by telephone. Two trials used random number tables (Wang 2003; Xu 2004). One trial used adequate concealment (envelopes) prior to allocation (Song 1994). Concealment was not used in the other two trials (Wang 2003; Xu 2004).

No trial mentioned blinding. By interviewing the authors it was identified that only one trial was single blind, for the participants (Song 1994). The assessors were not blinded in any trial.

As follow up and withdrawal rates were not reported in all trials, and no participant 'Flow diagram' was available, it is unclear whether all the participants were followed up. As a result, we judged the losses to follow up and withdrawal from the trials by the data presented.

The three trials did not report on 'relief of symptoms' or 'quality of life or satisfaction', which were primary outcomes for this review. We were unable to obtain any information about the protocol for these trials. They were judged 'Unclear' due to insufficient information.

No trial mentioned other potential sources of bias, and the source of funding was not clear.

### Potential biases in the review process

We failed to find any unpublished or ongoing trials. This may have led to potential bias.

### Agreements and disagreements with other studies or reviews

This review was originally published two years ago and there has not been any other systematic review about this intervention. It is not clear whether the methods, including microwave treatment, have any benefit for women. However, women with cervical ectropion continue to have treatment with various topical treatments. Many of these studies are published in Chinese journals and suggest effectiveness but none of them meet the requirements of a randomised controlled clinical trial.

## AUTHORS' CONCLUSIONS

### Implications for practice

Although the appearance of the cervix can be improved with microwave and other therapies it is not clear whether this has any benefit for the women. Women presenting with increased vaginal discharge from an ectropion can be advised that this is physiological, following swabs to rule out sexually transmitted infections, bacterial vaginosis and candidiasis. Postcoital bleeding, with an associated ectropion, requires follow up with cervical cytology and colposcopy to exclude cervical cancer or its precursors.

### Implications for research

Further studies should be undertaken and they need to consider the rationale for treating cervical ectropion, including primary outcomes that are important for women. Only symptomatic women with cervical ectropion should be included and their characteristics clearly reported. Treatment needs to be compared with no treatment. Correct randomisation is required for allocating the participants along with allocation concealment, long-term follow up, blind assessment of the results, presentation of complete outcome data, clarification of other potential sources of bias and more careful and detailed monitoring of adverse effects. Reporting on the quality issues should be improved according to Consolidated Standards of Reporting Trials (CONSORT) Statement (Schulz 2010).

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**CHARACTERISTICS OF STUDIES**
**Characteristics of excluded studies** [ordered by study ID]

Study	Reason for exclusion
Ban 2004	Non-RCT (through call to the author, we found it was a pseudo-randomised controlled trial)
Cao 2006	Non-RCT (according to looking up study in detail)

Study	Reason for exclusion
Chen 1999	Non-RCT (according to looking up study in detail)
Chen 2002a	Non-RCT (according to looking up study in detail)
Chen 2003	Non-RCT (according to looking up study in detail)
Chen 2006a	Non-RCT (according to looking up study in detail)
Chen 2006b	Non-RCT (through call to the author, we found it was a pseudo-randomised controlled trial)
Chen 2007	Non-RCT (according to looking up study in detail)
Chen 2008	There was no documentation as to whether the women who had cervical ectropion were symptomatic
Cheng 2005	Non-RCT (through call to the author, we found it was a pseudo-randomised controlled trial)
Cheng 2006a	Non-RCT (according to looking up study in detail)
Dai 2002	Non-RCT (through call to the author, we found it was a pseudo randomised controlled trial)
Dai 2004	Non-RCT (according to looking up study in detail)
Dai 2008	Non-RCT (according to looking up study in detail)
Dang 2003	Non-RCT (through call to the author, we found it was a pseudo randomised controlled trial)
Fan 2003	Non-RCT (according to looking up study in detail)
Fan 2005	Non-RCT (according to looking up study in detail)
Feng 2008	Non-RCT (through call to the author, we found it was a pseudo-randomised controlled trial)
Gan 2006	Non-RCT (according to looking up study in detail)
Gao 2005	Non-RCT (according to looking up study in detail)
Gu 2005a	Non-RCT (through called for the author, we found it was a pseudo randomised controlled trial)
Gu 2005b	Non-RCT (according to looking up study in detail)
Guan 2001	Non-RCT (through call to the author, we found it was a pseudo randomised controlled trial)
Guo 1997	Retrospective clinical trial (according to looking up study in detail)
Guo 2003	Non-RCT (through call to the author, we found it was a pseudo randomised controlled trial)
Guo 2008	There was no documentation as to whether the women who had cervical ectropion were symptomatic
Han 2007	Non-RCT (through call to the author, we found it was a pseudo-randomised controlled trial)
Hao 2006	Quasi-randomised controlled trial (through call to the author)
He 2000	Non-RCT (through call to the author, we found it was a pseudo- randomised controlled trial)

Study	Reason for exclusion
He 2009	There was no documentation as to whether the women who had cervical ectropion were symptomatic
Hu 1996	Non-RCT (through call to the author, we found it was a pseudo- randomised controlled trial)
Hu 2002	Retrospective clinical trial (through call to the author)
Hu 2007	Non-RCT (according to looking up study in detail)
Hu 2008	Non-RCT (according to looking up study in detail)
Huang 1997	Non-RCT (through call to the author, we found it was a pseudo- randomised controlled trial)
Huang 2000	Retrospective clinical trial (according to looking up study in detail)
Ji 2003	Non-RCT (through call to the author, we found it was a pseudo- randomised controlled trial)
Jiang 2001	Non-RCT (through call to the author, we found it was a pseudo- randomised controlled trial)
Lei 2007	Non-RCT (according to looking up study in detail)
Li 2002	Non-RCT (through call to the author, we found it was a pseudo- randomised controlled trial)
Li 2003	Non-RCT (according to looking up study in detail)
Lin 2004	Non-RCT (through call to the author, we found it was a pseudo- randomised controlled trial)
Liu 2001a	Quasi-randomised controlled trial (through call to the author)
Liu 2001b	Quasi-randomised controlled trial (through call to the author)
Liu 2002	Quasi-randomised controlled trial (through call to the author)
Liu 2003a	Non-RCT (through call to the author, we found it was a pseudo randomised controlled trial)
Liu 2003b	Non-RCT (through call to the author, we found it was a pseudo randomised controlled trial)
Liu 2003c	Non-RCT (through call to the author, we found it was a pseudo-randomised controlled trial)
Liu 2004	Non-RCT (according to looking up study in detail)
Liu 2005	Retrospective clinical trial (according to looking up study in detail)
Liu 2006	Non-RCT (according to looking up study in detail)
Long 2003	Quasi-randomised controlled trial (Although called for the author)
Lu 2004	Non-RCT (Through called for the author, we found it was a pseudo randomised controlled trial)
Luo 2004	Non-RCT (Through called for the author, we found it was a pseudo randomised controlled trial)
Luo 2007	Non-RCT (according to looking up study in detail)
Lv 2004a	Non-RCT (through call to the author, we found it was a pseudo-randomised controlled trial)

Study	Reason for exclusion
<a href="#">Lv 2004b</a>	Non-RCT (through call to the author, we found it was a pseudo-randomised controlled trial)
<a href="#">Ma 2001</a>	Non-RCT (through call to the author, we found it was a pseudo-randomised controlled trial)
<a href="#">Mo 2004</a>	Non-RCT (through call to the author, we found it was a pseudo-randomised controlled trial)
<a href="#">Pan 1999</a>	Non-RCT (according to looking up study in detail)
<a href="#">Qin 2001</a>	Non-RCT (through call to the author, we found it was a pseudo-randomised controlled trial)
<a href="#">Ran 2004</a>	Non-RCT (through call to the author, we found it was a pseudo-randomised controlled trial)
<a href="#">Ruan 2007</a>	Non-RCT (through call to the author, we found it was a pseudo-randomised controlled trial)
<a href="#">Shan 2002</a>	Quasi-randomised controlled trial (through call to the author)
<a href="#">Shao 2003</a>	Non-RCT (through call to the author, we found it was a pseudo-randomised controlled trial)
<a href="#">Shu 2006</a>	Non-RCT (according to looking up study in detail)
<a href="#">Song 1994</a>	RCT ( There was no documentation as to whether the women who had cervical ectropion were symptomatic)
<a href="#">Sun 2006a</a>	Non-RCT (according to looking up study in detail)
<a href="#">Sun 2006b</a>	Non-RCT (according to looking up study in detail)
<a href="#">Wang 1996</a>	Non-RCT (Through called for the author, we found it was a pseudo randomised controlled trial)
<a href="#">Wang 1997</a>	Retrospective clinical trial (Through called for the author)
<a href="#">Wang 2001</a>	Non-RCT (through call to the author, we found it was a pseudo-randomised controlled trial)
<a href="#">Wang 2002</a>	Non-RCT (through call to the author, we found it was a pseudo-randomised controlled trial)
<a href="#">Wang 2002c</a>	Non-RCT (we found it was a pseudo-randomised controlled trial)
<a href="#">Wang 2003</a>	RCT ( There was no documentation as to whether the women who had cervical ectropion were symptomatic.)
<a href="#">Wang 2006a</a>	Non-RCT (according to looking up study in detail)
<a href="#">Wang 2006b</a>	Non-RCT (according to looking up study in detail)
<a href="#">Wang 2008</a>	Non-RCT (according to looking up study in detail)
<a href="#">Wei 2008</a>	There was no documentation as to whether the women who had cervical ectropion were symptomatic
<a href="#">Wen 2002</a>	Non-RCT (through call to the author, we found it was a pseudo-randomised controlled trial)
<a href="#">Wen 2003</a>	Non-RCT (through call to the author, we found it was a pseudo-randomised controlled trial)
<a href="#">Wu 1996</a>	Non-RCT (through call to the author, we found it was a pseudo-randomised controlled trial)

Study	Reason for exclusion
Wu 2003	Retrospective clinical trial (through call to the author)
Wu 2004	Non-RCT (through call to the author, we found it was a pseudo-randomised controlled trial)
Wu 2007	There was no documentation as to whether the women who had cervical ectropion were symptomatic
Xie 1999	Quasi-randomised controlled trial (Through called for the author)
Xie 2003	Non-RCT (through call to the author, we found it was a pseudo-randomised controlled trial)
Xie 2006	Non-RCT (according to looking up study in detail)
Xie 2007	Non-RCT (according to looking up study in detail)
Xu 2000	Non-RCT (through call to the author, we found it was a pseudo-randomised controlled trial)
Xu 2004	RCT ( there was no documentation as to whether the women who had cervical ectropion were symptomatic.)
Xu 2005	Non-RCT (through call to the author, we found it was a pseudo-randomised controlled trial)
Xu 2007a	Non-RCT (through call to the author, we found it was a pseudo-randomised controlled trial)
Xu 2007b	Non-RCT (according to looking up study in detail)
Xue 2002	Non-RCT (we found it was a pseudo-randomised controlled trial)
Xue 2004	Non-RCT (according to looking up study in detail)
Yang 1997	Retrospective clinical trial (according to looking up study in detail)
Yang 2001	Non-RCT ( According to looking up study in detail)
Yang 2007	Non-RCT (according to looking up study in detail)
Yang 2008a	Non-RCT (according to looking up study in detail)
Yang 2008b	Non-RCT (according to looking up study in detail)
Ye 2005	Non-RCT (Through called for the author, we found it was a pseudo randomised controlled trial)
You 2004	Quasi-randomised controlled trial (Through called for the author)
Yu 2003	Non-RCT (according to looking up study in detail)
Yu 2004	Retrospective clinical trial (through call to the author)
Yu 2008	Non-RCT (according to looking up study in detail)
Zhan 2001	Non-RCT (through call to the author, we found it was a pseudo-randomised controlled trial)
Zhang 1999	Non-RCT (through call to the author, we found it was a pseudo-randomised controlled trial)
Zhang 2001	Retrospective clinical trial (according to looking up study in detail)

Study	Reason for exclusion
Zhang 2002	Retrospective clinical trial (through call to the author)
Zhang 2005a	Non-RCT (Through called for the author, we found it was a pseudo randomised controlled trial)
Zhang 2005b	Non-RCT (through call to the author, we found it was a pseudo-randomised controlled trial)
Zhang 2005c	Non-RCT (through call to the author, we found it was a pseudo-randomised controlled trial)
Zhang 2006a	Duplicate publication literature (It was the same study with "Zhang 2006b")
Zhang 2006b	Quasi-randomised controlled trial (Although called for the author)
Zhang 2007	Non-RCT (according to looking up study in detail)
Zhao 1997	Non-RCT (through call to the author, we found it was a pseudo-randomised controlled trial)
Zhao 2003	Quasi-randomised controlled trial (through call to the author)
Zhao 2007	Non-RCT (according to looking up study in detail)
Zhao 2008	Non-RCT (according to looking up study in detail)
Zhong 2005	Non-RCT (according to looking up study in detail)
Zhou 2002	Non-RCT (Through called for the author, we found it was a pseudo randomised controlled trial)
Zhu 1997	Non-RCT (through call to the author, we found it was a pseudo-randomised controlled trial)
Zhu 2007	Non-RCT (according to looking up study in detail)

### Characteristics of studies awaiting assessment [ordered by study ID]

#### Chen 2001

Methods	RCT duration: not yet known Randomisation method not stated Blinding: not yet known Number of women randomised: n = 204 Number of women analysed: n = 204 Unit of comparison: individuals.
Participants	Inclusion criteria: women with cervical erosion without cervical cancer and infusorium, neisseria gonorrhoeae vaginitis and mould infection Microwave group (n = 106): Grade I: 22; Grade II: 58; Grade III: 26 Age: 32y; No.of pregnancy: 2.2; No. of births: 1.2 Electrocautery group (n = 98):

**Chen 2001** (Continued)

Grade I: 28; Grade II: 50; Grade III: 20

Age: 30y; No. of pregnancy: 3.1; No. of births: 1.6

Source of participants: medical outpatients.

**Interventions**

Treatment: microwave tissue coagulation (MTC)  
Control: electrocautery.

Time of treatments: during 3 to 7 days after menstrual period.

**Outcomes**

Efficacy: cure rate (according to appearance of cervix)

Adverse reaction: bleeding during operation, bellyache pain during operation, more bleeding during decrustation than menstruation

The follow-up period: 8 to 12 weeks

**Notes**

Location: China

Setting: Department of Gynaecology and Obstetrics of Nanyou Hospital, Zhanjiang City and No. 422 Liberation Army Hospital

Funding: unclear

Failed to contact the author, and it is unclear whether the trial is an authentic RCT.

**Chen 2002b**

**Methods**

RCT duration: not yet known

Randomisation method not stated

Blinding: not yet known

Number of women randomised: n = 360

Number of women analysed: n = 360

Unit of comparison: individuals.

**Participants**

Inclusion criteria: women with cervical erosion without cervical cancer and infusorium and mould infection

Interferon suppository group: n = 120

Microwave group: n = 120

Interferon suppository plus microwave group: n = 120

Grade I. interferon suppository group: 24; microwave group: 24; interferon suppository plus microwave group: 24

Grade II. interferon suppository group: 63; microwave group: 63; interferon suppository plus microwave group: 63

Grade III. interferon suppository group: 33; microwave group: 33; interferon suppository plus microwave group: 33

Source of participants: medical outpatients.

**Interventions**

1. Interferon suppository

**Microwave therapy for cervical ectropion (Review)**

**Chen 2002b** (Continued)

2. Microwave
3. Interferon suppository plus microwave

Outcomes	<p>Efficacy: cure rate (according to appearance of cervix)</p> <p>The time of vaginal fluid and the status of vaginal bleeding</p> <p>The follow-up period: 8 weeks</p>
Notes	<p>Location: China</p> <p>Setting: Department of Gynaecology and Obstetrics, Nanjing first hospital affiliated to Nanjing Medical University</p> <p>Funding: unclear</p> <p>Failed to contact the author, and it is unclear whether the trial is an authentic RCT.</p>

**Chen 2004**

Methods	<p>RCT duration 5 years from Jan 2000 to Dec 2005</p> <p>Randomisation method not stated</p> <p>Single centre</p> <p>Blinding: not yet known</p> <p>Number of women randomised: n = 188</p> <p>Number of women analysed: n = 188</p> <p>Unit of comparison: individuals.</p>
Participants	<p>Inclusion criteria: women with severe cervical erosion without cervical cancer and infusorium, mould infection and venereal disease</p> <p>Microwave group: n = 98</p> <p>Interferon suppository plus microwave group: n = 90</p> <p>Exclusion criteria: excluded women with cervical cancer, diagnosed through Papanicolaou test. Excluded women with infusorium, mould infection and venereal disease</p> <p>Source of participants: medical outpatients.</p>
Interventions	<ol style="list-style-type: none"> <li>1. Microwave group: 45W (power)</li> <li>2. Interferon suppository plus microwave group: after microwave, interferon suppository one pill once every other day, course of treatment for 15 days</li> </ol>
Outcomes	<p>Efficacy: Cure rate (according to appearance of cervix)</p> <p>The follow-up period: not yet known</p>
Notes	<p>Location: China</p> <p>Setting: Department of Gynaecology and Obstetrics of Mudanjiang Railway Central Hospital in Heilongjiang</p> <p>Funding: unclear</p>



**Chen 2004** (Continued)

Failed to contact the author, and it is unclear whether the trial is an authentic RCT.

**Cheng 2000**

Methods	Randomisation method: not yet known Single centre Blinding: not yet known.
Participants	Inclusion criteria: women with severe cervical erosion
Interventions	1. Treatment: microwave group 2. Control: not yet known
Outcomes	Efficacy: cure rate (according to appearance of cervix) The follow-up period: not yet known
Notes	Location: China Setting: Department of Gynaecology and Obstetrics of Yiwu Traditional Chinese Medicine Hospital Funding: unclear Failed to find the full text.

**He 1995**

Methods	RCT duration: not yet known Randomisation method: not yet known Single centre Blinding: not yet known Number of women randomised: n = 260 Number of women analysed: n = 260 Unit of comparison: individuals.
Participants	Inclusion criteria: women with cervical erosion Microwave group: n = 130 Electrocautery group: n = 130 Source of participants: medical outpatients.
Interventions	1. Microwave group 2. Electrocautery group
Outcomes	Efficacy: cure rate (according to appearance of cervix)

**He 1995** (Continued)

	The follow-up period: not yet known
Notes	Location: China Setting: The Third People's Hospital of Xiaoshan City, Zhejiang province Funding: unclear Failed to contact the author, and it is unclear whether the trial is an authentic RCT.

**Li 2004**

Methods	Randomisation method: not yet known Single centre Blinding: not yet known.
Participants	Inclusion criteria: women with severe cervical erosion
Interventions	1. Treatment: microwave group 2. Control: not yet known
Outcomes	Efficacy: cure rate The follow-up period: not yet known
Notes	Location: China Setting: Women and Children's Hospital of Jinzhou City Funding: unclear Failed to find the full text.

**Shi 2000**

Methods	Randomisation method: not yet known Single centre Blinding: not yet known.
Participants	Inclusion criteria: women with severe cervical erosion
Interventions	1. Treatment: microwave group 2. Control: not yet known
Outcomes	Efficacy: cure rate (according to appearance of cervix) The follow-up period: not yet known
Notes	Location: China Setting: Women and Children's Hospital of Nantong City

**Microwave therapy for cervical ectropion (Review)**

**Shi 2000** (Continued)

Funding: unclear  
Failed to find the full text.

**Sun 2001**

Methods	<p>RCT duration 2.5 years from Jul 1997 to Dec 1999</p> <p>Randomisation method not stated</p> <p>Single centre</p> <p>Blinding: not yet known</p> <p>Number of women randomised: n = 150</p> <p>Number of women analysed: n = 150</p> <p>Unit of comparison: individuals.</p>
Participants	<p>Inclusion criteria: women with cervical erosion without cervical cancer and infusorium, neisseria gonorrhoeae vaginitis and mould infection</p> <p>Microwave group (n = 106):</p> <p>Grade I: 18; Grade II: 47; Grade III: 25</p> <p>Age: 32y (20 to 40)</p> <p>Electrocautery group (n = 90):</p> <p>Grade I: 12; Grade II: 31; Grade III: 17</p> <p>Age: 33y (19 to 56)</p> <p>Source of participants: medical outpatients.</p>
Interventions	<p>1. Microwave group</p> <p>2. Electrocautery group</p> <p>The follow-up period: 8 and 36 months. Once a week after treatment, once 4 weeks, once 8 weeks, then once 3 months for going on 1 year. And then once half yearly.</p>
Outcomes	<p>Efficacy: cure rate (according to appearance of cervix)</p> <p>The time of vaginal fluid and the status of vaginal bleeding</p> <p>Bleeding during operation and after operation, more bleeding during decrustation than menstruation</p>
Notes	<p>Location: China</p> <p>Setting: The first hospital affiliated to Qiqihaer Medical College of Heilongjiang</p> <p>Funding: unclear</p> <p>Failed to contact the author, and it is unclear whether the trial is an authentic RCT.</p>

**Yang 2004**

Methods	Randomisation method: not yet known  Single centre  Blinding: not yet known
Participants	Inclusion criteria: women with severe cervical erosion
Interventions	1. Treatment: microwave group  2. Control: not yet known
Outcomes	Efficacy: cure rate  The follow-up period: not yet known
Notes	Location: China  Setting: Department of Gynaecology and Obstetrics of Shuangya People's Hospital  Funding: unclear  Failed to find the full text.

**ADDITIONAL TABLES**
**Table 1. Questions used to identify an authentic RCT**

Process	Questions
Question 1	Introduce myself and purpose: How do you do? I was a student of Lanzhou University, was doing a review about microwave tissue coagulation in the treatment of cervical ectropion. Purpose of my study was to compare the effects of variety randomisation methods. I had searched out a paper that published in (time, journal) written by you. Could you please tell me what method to be used in this trial?
Question 2	If the subject cannot describe the method clearly, change the question like this: could you please tell me when a new participant enrolled, how did you decide which group the participant should be allocated to?
Question 3	If there was any problem about the first author, the second author or others should be interviewed.
Question 4	Next two questions aim to understand the category of support source for investigated study, I selected one of them or both: (1) Was your study funded by government or any other source? (2) Was your study for new drug development?
Question 5	Judgment should be made immediately for whether it was a real RCT or not. If it be judged as real RCT, next questions aim to understand the status of allocation concealment: (1) Do you know allocation concealment? If so, please clarify. (2) Did you use any method to mask allocation sequence? If any, please clarify.
Question 6	Next question aims to understand the validity of blinding particularly pay attention to whether the detector was blinded or not: Please tell me who were blinded in your trial?
Question 7	We said "thank you" to author.

**Table 1. Questions used to identify an authentic RCT** (Continued)

Question 8	Finally, We needed to judge whether the subject knew well the trial principle or not: if anyone persisted in the method of randomisation as correct but actually wrong or ineligible, it should be judged didn't know; if anyone claimed that "I knew we performed not good enough" or "it was impossible to perform a completely correct RCT" and so on, it should be judged as knew well about the principle of trial but violated knowingly claimed the non-RCT as RCT.
Question 9	Put all of record in the form.

RCTs: randomised controlled trials

**Table 2. Characteristics of the first RCT**

Study	Song 1994
Methods	Randomization method not stated. Parallel, single centre, single blinded study. Number of women randomised: n = 300; Number of women analysed: n = 300. Units of comparison: individuals. Source of funding: unclear.  RCT duration: from Jan 1991 to May 1992
Participants	Inclusion criteria: women with cervical ectropion. Macroscopic examination: discriminating acute with chronic cervicitis. Age: MTC group (n = 154): 19 to 57 years (mean age 33); Laser group (n = 146): 28 to 54 years (mean age 35). Average gravidity: MTC group: 2.2; laser group: 1.9. Average parity: MTC group: 1.0; laser group: 1.0. Grade II: MTC group: 32; laser group: 43; Grade III: MTC group: 122; laser group: 103. Symptom: unclear. Exclusion criteria: excluded women with cervical intraepithelial neoplasia (CIN1, 2, 3), cancer in situ or early cervical cancer which were diagnosed through Papanicolaou test and/or histological examination or colposcopy. Excluded women with vaginal or cervical infection. Source of participants: medical outpatients.
Interventions	Treatment: MTC; Control: CO2 laser. Timing of treatment: during 3 to 7 days after menstrual period (once).
Outcomes	Efficacy: cure rate (according to appearance of cervix). Adverse effect: bleeding during operation, bellyache during operation, bleeding post-operation, topical infection bleeding post-operation. The follow-up period: 8-12 weeks.
Note	Location: China  Setting: Department of Gynaecology and Obstetrics of Chaoyang Hospital  Funding: unclear

 MTC: microwave tissue coagulation  
 CIN: cervical intraepithelial neoplasia

**Table 3. Characteristics of the second RCT**

Study	Wang 2003
Methods	Randomisation: random digits table; Parallel, single centre, no blinding. Number of women randomised: n = 120; Number of women analysed: n = 120. Units of comparison: individuals. Source of funding: unclear.  RCT duration: from Aug 2001 to Oct 2001.
Participants	Inclusion criteria: women with cervical ectropion. Age: MTC group (n = 60): unclear; Laser group (n = 60): unclear. Grade I: MTC group: 8; laser group: 9; Grade II: MTC group: 27; laser group: 28; Grade III: MTC group: 25; laser group: 23; Simple pattern: MTC group: 42; laser group: 43; Granular pattern or Mammilla pattern: MTC group: 18; laser group: 17. Symptom: unclear. Exclusion criteria: Excluded women with cervical intraepithelial neoplasia (CIN1, 2, 3), cancer in situ or early cervical cancer. Excluded women with vaginal or cervical infection. Source of participants: not stated.
Interventions	Treatment: MTC;  Control: CO <sub>2</sub> laser. Timing of treatment: during 3 to 7 days after menstrual period (once).
Outcomes	Efficacy: Cure rate (According to appearance of cervix); Adverse effect: vaginal discharge post-treatment. The follow-up period: 2,3,4,6,8,12 weeks.
Note	Location: China.  Setting: Shiyan Maternal and Child Health Hospital, Hubei.  Funding: unclear.

MTC: microwave tissue coagulation  
 CIN: cervical intraepithelial neoplasia

**Table 4. Characteristics of the third RCT**

Study	Xu 2004
Methods	Randomisation: random digits table; Parallel, single centre, no blinding. Number of women randomised: n = 120; Number of women analysed: n = 120. Units of comparison: individuals. Source of funding: Unclear. RCT duration: from Jun 2000 to May 2002.
Participants	Inclusion criteria: women with cervical ectropion. Papanicolaou test: Grade I or II. Age: 23 to 50 years (mean age 33). Grade I: MTC group: 15; Laser group: 13; Grade II: MTC group: 27; Laser group: 28;

**Table 4. Characteristics of the third RCT** *(Continued)*

	Grade III: MTC group: 19; Laser group: 18. Symptom: unclear. Exclusion criteria: not reported. Source of participants: medical outpatient.
Interventions	Treatment: MTC (once); Control: interferon-alpha suppository (aoping suppository): 6000 U once/two days; duration: 6-10 days. (Repeat it after next menstrual period). Timing of treatment: during 3 to 7 days after menstrual period.
Outcomes	Efficacy: cure rate (according to appearance of cervix); Adverse effect: vaginal discharge post-treatment. The follow-up period: 8, to 12 weeks.
Note	Location: China  Setting: Department of Gynaecology and Obstetrics of The First Hospital of Nanping, Fujian.  Funding: unclear

MTC: microwave tissue coagulation  
 CIN: cervical intraepithelial neoplasia

**Table 5. Quality table**

Study ID	Song 1994	Xu 2004	Wang 2003
Design	Parallel single-centre.	Parallel single-centre.	Parallel single-centre.
Adequate sequence generation?	Yes.	Yes(random number table).	Yes(random number table).
Allocation concealment?	Yes(envelopes).	No.	No.
Blinding?	Yes (Single: participants).	No.	No.
Incomplete outcome data addressed?	Unclear.	Unclear.	Unclear.
Free of selective reporting?	Unclear.	Unclear.	Unclear.
Free of other bias?	Unclear.	Unclear.	Unclear.
Power calculation	Unclear.	Unclear.	Unclear.

## APPENDICES

### Appendix 1. MEDLINE search strategy

1 exp uterine cervical erosion/ or exp uterine cervicitis/.

2 (cervi\$ adj2 erosion\$).tw

3 cervicitis.tw.

4 (cervi\$ adj2 ectop\$).tw.

- 5 (cervi\$ adj2 ectrop\$).tw.
- 6 columnar ectop\$.tw.
- 7 (cervi\$ adj2 infect\$).tw.
- 8 or/1-7
- 9 exp Microwaves/
- 10 Microwave\$.tw.
- 11 ultrahigh frequency wave\$.tw.
- 12 radio wave\$.tw.
- 13 ehf wave\$.tw.
- 14 or/9-13
- 15 8 and 14
- 16 randomised controlled trial.pt.
- 17 controlled clinical trial.pt.
- 18 randomized.ab.
- 19 placebo.tw.
- 20 clinical trials as topic.sh.
- 21 randomly.ab.
- 22 trial.ti.
- 23 (crossover or cross-over or cross over).tw.
- 24 or/16-23
- 25 (animals not (humans and animals)).sh.
- 26 24 not 25
- 27 26 and 15

## **Appendix 2. EMBASE search strategy**

- 1 exp uterine cervicitis/ or exp uterine cervix erosion/
- 2 (cervi\$ adj2 erosion\$).tw.
- 3 cervicitis.tw.
- 4 (cervi\$ adj2 ectop\$).tw.
- 5 (cervi\$ adj2 ectrop\$).tw.
- 6 columnar ectop\$.tw.
- 7 (cervi\$ adj2 infect\$).tw.
- 8 or/1-7
- 9 exp Microwave Radiation/
- 10 Microwave\$.tw.
- 11 ultrahigh frequency wave\$.tw.

### **Microwave therapy for cervical ectropion (Review)**



12 radio wave\$.tw.

13 ehf wave\$.tw.

14 or/9-13

15 8 and 14

16 Clinical Trial/

17 Randomized Controlled Trial/

18 exp randomisation/

19 Single Blind Procedure/

20 Double Blind Procedure/

21 Crossover Procedure/

22 Placebo/

23 Randomized controlled trial\$.tw.

24 Rct.tw.

25 random allocation.tw.

26 randomly allocated.tw.

27 allocated randomly.tw.

28 (allocated adj2 random).tw.

29 Single blind\$.tw.

30 Double blind\$.tw.

31 ((treble or triple) adj blind\$).tw.

32 placebo\$.tw.

33 prospective study/

34 or/16-33

35 case study/

36 case report.tw.

37 abstract report/ or letter/

38 or/35-37

39 34 not 38

40 39 and 15

### **Appendix 3. The Cochrane Library search strategy**

1 exp uterine cervical erosion/ or exp uterine cervicitis/

2 (cervi\$ adj2 erosion\$).tw.

3 cervicitis.tw.

4 (cervi\$ adj2 ectop\$).tw.

5 (cervi\$ adj2 ectrop\$).tw.

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#### **Microwave therapy for cervical ectropion (Review)**

6 columnar ectop\$.tw.

7 (cervi\$ adj2 infect\$.tw.

8 or/1-7

9 exp Microwaves/

10 Microwave\$.tw.

11 ultrahigh frequency wave\$.tw.

12 radio wave\$.tw.

13 ehf wave\$.tw.

14 or/9-13

15 8 and 14

## WHAT'S NEW

Date	Event	Description
20 September 2010	Amended	Contact details updated.
20 March 2010	Review declared as stable	The conclusions of this review are believed to be well established, therefore there is no plan to update this review in the future.

## HISTORY

Protocol first published: Issue 4, 2006

Review first published: Issue 4, 2007

Date	Event	Description
31 May 2009	New search has been performed	Converted to new review format.
3 August 2007	New citation required but conclusions have not changed	conclusions not changed

## CONTRIBUTIONS OF AUTHORS

Yali Liu and Taixiang Wu were responsible for protocol and review development, Kehu Yang was responsible for organising the research team and review development.

Data Collection for the review was carried out independently by LYL,YK, ZP and TJY.

Entering data into RevMan: LYL, TJ, MB.

Analysis of data: WT, LYL,YH, LJ.

Interpretation of data: WT, LYL, YH.

Writing the protocol and review: LYL, YK.

Providing general advice on the protocol and review: WT, HR.

Helen Roberts was the English and western medical clinical contributor

## DECLARATIONS OF INTEREST

None known

## Microwave therapy for cervical ectropion (Review)

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## SOURCES OF SUPPORT

### Internal sources

- No sources of support found, Not specified.

### External sources

- No sources of support supplied

## DIFFERENCES BETWEEN PROTOCOL AND REVIEW

Ya Li Liu is the first author.

## INDEX TERMS

### Medical Subject Headings (MeSH)

Chronic Disease; Microwaves [\*therapeutic use]; Randomized Controlled Trials as Topic; Uterine Cervicitis [\*radiotherapy]

### MeSH check words

Female; Humans