Management of haemorrhoids: protocol of an umbrella review of systematic reviews and meta-analyses

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ABSTRACT

Introduction The prevalence of haemorrhoidal diseases was high in general population, and many treatments are proposed for the management of haemorrhoids. The treatments include conservative and surgical interventions; the credibility and strength of current evidence of their effectiveness are not comprehensively evaluated. We aim to evaluate the credibility of systematic reviews and meta-analyses that assess the effectiveness of the treatments for haemorrhoidal diseases through an umbrella review.

Methods and analysis We will search Ovid Medline, Embase, Cochrane library and Web of Science from inception to March 2020 without any language restriction. We will include meta-analyses that examine the effectiveness of treatments in the management of haemorrhoids. Two reviewers will independently screen the titles and abstracts of retrieved articles, and they will extract data from the included meta-analyses. For each meta-analysis, we will estimate the effect size of a treatment through the random-effect model and the fixed-effect model, and we will evaluate between-study heterogeneity (Cochrane’s Q and I² statistics) and small-study effect (Egger’s test); we will also estimate the evidence of excess significance bias. Evidence of each treatment will be graded according to prespecified criteria. Methodological quality of each meta-analysis will be evaluated by using Assessmentof Multiple Systematic Reviews 2. The corrected cover area method will be used to assess the impact of overlap in reviews on the findings of the umbrella review.

Ethics and dissemination We will present the results of the umbrella review at conferences and publish the final report in a peer-reviewed journal. The umbrella review does not require ethical approval.

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INTRODUCTION

Haemorrhoidal disease is one of the most common anorectal conditions encountered in daily practice by general practitioners. Symptoms related to haemorrhoids include bleeding during or after defecation, pain or discomfort, and perianal itch or irritation. According to a national health survey, an estimated 23 million adults were diagnosed with haemorrhoids in the USA, accounting for 13% of the US population. Another study from Australia showed that 38.93% of its general population suffered from haemorrhoidal diseases.1 There were 306,000 hospital dischargers and at least 2.2 million outpatien evaluations for haemorrhoids in the USA,2,3 and the demand for haemorrhoidal therapy is predicted to increase.4 Although the total annual costs for medications being used to treat haemorrhoid are unclear, one of the popular medications for haemorrhoid, Preparation H, was sold for a total cost of US$136 million in 2017.1

Many treatment options are proposed for the management of haemorrhoids. For lower grade of haemorrhoids (grades I–II by a classification system proposed by Sir Goligher5), conservative treatments like dietary interventions, lifestyle modification and medication treatment are usually adopted.5–6 A meta-analysis showed that fibre supplementation could reduce the risk of bleeding after defecation.7 Another meta-analysis showed that fibre supplementation provided consistent beneficial effect on perianal pain and itching.8 Constipation is a known risk factor for the development of haemorrhoids,9 and probiotics and probiotics improve the symptoms of constipation10 and therefore are helpful as supplements for managing haemorrhoids.11 For medication treatment, topical glucocorticoids, vasoconstrictors like phenylephrine-based suppositories, or analgesics may be...
beneficial for temporarily relieving haemorrhoidal symptoms. Herbal remedies are also prepared as suppositories or topical agents for alleviating haemorrhoidal symptoms. Although plentiful options of conservative treatments are available in clinical practice, many of them are not evidence based and the long-term effectiveness of them have not been verified in randomised controlled trials.

For higher grade of haemorrhoids (grades III–IV), surgical interventions are normally suggested; these interventions include rubber band ligation (RBL), stapled haemorrhoidopexy, haemorrhoidectomy and haemorrhoidal artery ligation (HAL). The most commonly used surgical interventions are traditionally excisional haemorrhoidectomy including open (Milligan-Morgan procedure) and closed (Ferguson procedure) haemorrhoidectomy. The most important disadvantage of haemorrhoidectomy is believed to be postoperative complications like postoperative pain and urinary retention. For the consideration of this disadvantage, new surgical procedures like HAL and stapled haemorrhoidopexy are developed to lower the incidence of postoperative complications. However, these new procedures are criticised for high recurrence rate after surgery and high healthcare expenditure, although they reduce the rate of the postoperative complications.

Several meta-analyses comparing the effectiveness of different surgical interventions have been published. The quality of these meta-analyses varied across studies. No systematic review of these meta-analyses has been performed to provide an evidence map for the management of haemorrhoids. Umbrella review is a new systematic review method that quantitatively summarises up-to-date evidence of a specific clinical scenario. It provides overview of current evidence and finds out the uncertainty to guide future research. For the large amount of evidence on conservative and surgical treatments for the management of haemorrhoidal diseases, we aim to conduct an umbrella review to summarise the findings of meta-analyses on conservative and surgical treatments for the management of haemorrhoids and evaluate the strength and credibility of the findings.

**METHODS AND ANALYSIS**

**Protocol registration and reporting of findings**

We design the protocol of the review according to the guidelines of Preferred Reporting Items for Systematic Review and Meta-analysis Protocols (PRISMA-P). The results of the review will be reported according to the recommendation of PRISMA. Any amendments to the protocol will be recorded and reported in an article with final results. Figure 1 shows the study process.

**Patient and public involvement statement**

The study is an umbrella review focusing on the management of haemorrhoids. Patients from the inpatient setting of the colorectal department at Hospital of Chengdu University of Traditional Chinese Medicine participated in the design of outcome assessments.

**Inclusion and exclusion criteria**

We will include meta-analyses that examine the effectiveness of lifestyle modification, conservative pharmacological treatments and surgical interventions (vs placebo, sham procedures, usual care or active control) in the management of haemorrhoids. Conservative pharmacological treatments will include oral supplements (insoluble fibres, hydrophilic bulk-forming colloids, prebiotics, probiotics and synbiotics), topical agents (glucocorticoids, vasoconstrictors and analgesics) and phlebotonic drugs comprising flavonoids. We will assess the efficacy and safety of lifestyle modification and conservative pharmacological treatments in the management of grades I–II haemorrhoids. Surgical interventions will include RBL, HAL, sclerotherapy, infrared coagulation, stapled...
haemorrhoidopexy and haemorrhoidectomy. Meta-

analyses of combined treatments (eg, adding a topical bioactive gel to a surgical intervention) will also be eval-

uated. Outcomes of interest are classified as: symptoms related to haemorrhoids (rectal bleeding, defecation pain and perianal itching with a sense of swelling), surgical related conditions (surgical time, postoperative pain, postoperative bleeding and urinary retention), recurrence of haemorrhoidal symptoms (<1 year and >1 year) and patient’s satisfaction (quality of life, time needed for return to usual activities and hospital stay). We will set no restrictions to the age of participants, study settings (eg, only inpatient or outpatient setting) or the language of publications. When multiple meta-analyses on the same topic (with the same treatment and the same outcomes) were retrieved, we will include the most recent one. We will consider inclusion of ongoing meta-analyses under the condition that primary analyses of these meta-analyses have been finished, and we will contact the authors to ask for related data when possible. We will exclude meta-

analyses that report only a summary estimate of effect size (ES) without any further data (eg, mean, SD and the number of events) of the primary studies. We will exclude narrative reviews since they provide no quan-

titative data (eg, means, SD or event rate) for analysis. We will exclude meta-analyses published in the format of letters to the editor since they usually contain little valu-

able information.

Outcome assessments
We will assess the following outcomes: improvement of haemorrhoidal symptoms, surgical related outcomes and patient’s satisfaction. The improvement of haemor-

rhoidal symptoms will be assessed in conservative treat-
ments. The haemorrhoidal symptoms will include rectal bleeding, defecation pain and perianal itching with a sense of swelling. We will adopt the criteria for justifi-

cation of improvement of haemorrhoidal symptoms as reported in each included systematic review. The surgical related outcomes will be assessed in surgical treatments. These outcomes will include surgical time, postoperative pain, postoperative bleeding, urinary retention and recurrence of haemorrhoidal symptoms (<1 year and >1 year). The surgical time refers to the duration needed for surgical procedure in managing haemorrhoids, which will be assessed as defined in each systematic review. Post-

operative pain will be defined as acute pain at days 1–3 after surgical intervention, including pain during or after defecation, pain during body movement or rest pain at rest. The assessment of the patient’s satisfaction includes quality of life, time needed for return to usual activities and hospital stay; it will be assessed in surgical treat-
ments, and it will be assessed at 1 month, 6 months and 12 months after treatment.

Study source and selection process
We will electronically search Ovid Medline, Embase, the Cochrane Library and Web of Science from inception to

March 2020 without any language restriction for potentially eligible candidates. The search strategy will be developed in consultation with an experienced librarian (Yu-Lan Ren from Chengdu University of Traditional Chinese Medicine). In developing the search strategy, four specialist-

ists (MC, T-CT, T-HH and YJD) from colorectal depart-

ment in Hospital of Chengdu University of Traditional Chinese Medicine will decide keywords, MeSH (Medical Subject Headings) terms and text words, which will be searched in combination: haemorrhoid, haemorrhoidal, systematic reviews and meta-

analyses (table 1 and online supplementary). Additional search will be performed by manual search of the reference lists of the retrieved arti-

cles and the search for ongoing meta-analyses registered in PROSPERO or the Cochrane Library. Table 1 shows the strategy developed for searching in Ovid Medline, and the search strategy for Embase and the Cochrane library is shown in the online supplementary. Retrieved articles will be imported into Zotero V.5.0.82 for screening. Two reviewers will independently screen titles and abstracts of retrieved articles according to the inclusion criteria. Full-
text copies will be accessed when the reviewers cannot decide the inclusion of a study through title or abstract screening. The excluded studies will be recorded along with the reason for exclusion at each screening stage. We will contact the corresponding authors of the published articles for additional information if necessary. When there is a disagreement between reviewers on the inclu-

sion of a study, we will solve the problem through group discussion.

Data collection and verification
Standardised abstraction forms will be used for data collection. Two reviewers (T-CT and MC) will use the forms to collect the following information: characteristics of included meta-

analyses (name of first author, publication year, name of intervention, the number of studies included in each meta-analysis, total sample size and the number of meta-

analyses), disease conditions (the grading of haemorrhoid (24), intervention and control (name of intervention or control, sample size of each treatment cohort and details of treatment) and outcomes (name and definition of outcome, summary ES and its related

<table>
<thead>
<tr>
<th>No.</th>
<th>Search items</th>
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<tbody>
<tr>
<td>1</td>
<td>hemorrhoid/</td>
</tr>
<tr>
<td>2</td>
<td>(hemorrhoid* or haemorrhoid*).ti,ab,kw.tw.</td>
</tr>
<tr>
<td>3</td>
<td>1 or 2</td>
</tr>
<tr>
<td>4</td>
<td>systematic review/ or meta-analysis/ or systematic review as topic/ or meta-analysis as topic/ or network meta-analysis/</td>
</tr>
<tr>
<td>5</td>
<td>(systematic review or meta-analysis$).ti,ab,kw.tw.</td>
</tr>
<tr>
<td>6</td>
<td>4 or 5</td>
</tr>
<tr>
<td>7</td>
<td>3 and 6</td>
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Table 1 Search strategy (via Ovid Medline)
95% CI and the number of participants included in the outcome assessment). When the data are only provided through plots, we will use Ycasd to determine the ES and its 95% CI; when necessary data were not provided in the article, we will contact the corresponding authors to ask for data. All data will be entered into Epi Info V.7.2 for data analysis. A third reviewer (YJD) will check the completeness and correctness of the extracted data.

Assessment of methodological quality

Methodological quality of the included meta-analyses will be assessed by using the Assessment of Multiple Systematic Reviews 2 (AMSTAR2, an updated version of AMSTAR) tool. AMSTAR2 has 16 domains; 7 were critical domains, on which the quality rating of an individual systematic review depends. Two reviewers (T-HH and DQ) will rate the quality of each meta-analysis as high, moderate, low and critically low based on the overall score of the AMSTAR2.

Data analysis

We will use standardised methods adopted by previous umbrella reviews and state-of-the-art approaches will be used to set criteria to evaluate the credibility of the findings. We will first estimate the summary ES and its related 95% CI using both random-effect and fixed-effect models. Second, we will estimate the 95% prediction interval (95% PI) for the summary estimate based on the random-effect model. The 95% PI specifies the uncertainty for the effect that will be expected in a future study examining the same clinical question. Third, we will evaluate heterogeneity of each meta-analysis by using Cochrane’s Q test (considered significant heterogeneity when p<0.1), and we will classify the degree of heterogeneity into low (I^2<25%), moderate (25%≤ I^2≤50%), large (50%≤ I^2<75%) or very large (I^2≥75%) through I^2 statistics.

Fourth, we will use Egger’s test to evaluate publication bias and small-study effect, and a p value<0.1 in the test confirms the bias and small-study effect. Fifth, we will perform an analysis that examines whether the observed number of original studies with positive findings in each meta-analysis is larger than their expected number to detect excess significance bias. The expected number is calculated as the sum of the statistical power estimates for each original study in a meta-analysis. The power of each study will be calculated through an algorithm using non-central t-distribution; the power calculation depends on the value of true ES. Since the true ES for any meta-analysis is impossible to acquire, we will use the ES from the largest study in a meta-analysis to substitute. We will calculate the ratio of observed number over expected number to evaluate the extent of excess significance bias, and we will claim existence of the bias when a χ^2 test arrives at a level of p<0.1.

Many systematic reviews and meta-analyses focusing on a similar topic include a different number of primary studies; the overall results and conclusions of an umbrella review might therefore be biased. To assess the potential impact of the overlap in the inclusion of the same primary studies, the degree of overlap within and between reviews was measured using the validated corrected cover area (CCA) method. A CCA score of 0–5 indicates slight overlap, 6–10 moderate, 11–15 high and >15 very high.

Criteria for evaluating credibility of evidence

We will use the following criteria to evaluate the credibility of the included meta-analyses: (1) having p<10^-6 on the basis of the random-effect model; (2) having >1000 participants in a single meta-analysis; (3) having low or moderate heterogeneity (I^2<50%); (4) having 95% PI that excludes the null value; (5) having no evidence of small-study effect; (6) having no evidence of excess significance bias. Meta-analysis that meets criteria (1)–(6) will be classified as convincing evidence (not suggestive of bias; class I evidence); meta-analysis that meets criteria (1)–(4) will be classified as highly suggestive evidence (class II evidence); meta-analysis that meets criteria (2) and has p<0.001 will be classified as suggestive evidence (class III evidence); meta-analysis that has only p<0.05 will be classified as weak evidence (class IV evidence).

DISCUSSION

Regarding the high prevalence of haemorrhoids in the general population and its heavy socioeconomic impact, we believe that it is important to provide an evidence map of treatments for haemorrhoids for clinical practitioners and patients, especially when plentiful systematic reviews and meta-analyses are available. Haemorrhoidal disease is one of the most common clinical conditions in practice. Multiple treatments including conservative and surgical treatments are available for the management of different grades of haemorrhoids. Conservative treatments are usually prescribed for grades I–II haemorrhoids, and surgical treatments are for grades II–IV haemorrhoids. Numerous clinical studies have been performed to study the effect of conservative and surgical treatments on haemorrhoids, and many systematic reviews and meta-analyses were conducted to evaluate and confirm the effectiveness of these treatments. However, an overview of the systematic reviews and meta-analyses is still lacking. An evidence map of treatments for the management of haemorrhoids is necessary, to facilitate the decision-making of clinical practitioners and participants. Current guidelines usually focus on the evidence of surgical treatments but neglect the evidence of conservative treatments, which also warrants an overview of current evidence.

Umbrella review is a review of systematic reviews and meta-analyses, which is viewed as one of the four next-generation meta-analyses. An umbrella review is able to quantitatively synthesise information from all systematic reviews and meta-analyses on a given topic. Being different from narrative reviews of systematic reviews, the umbrella review recalculates the ES of a treatment, evaluates the
credibility of the evidence by estimating excess significance bias and small-study effect and further classifies the confidence of the evidence into four levels.29 41 Our review will adopt the method of umbrella review, to re-evaluate the credibility of the evidence of treatments for haemorrhoidal diseases, especially the conservative treatments. To the best of our knowledge, the review will be the first to adopt the method of umbrella review in assessing the credibility of current evidence of haemorrhoidal management. The umbrella review is initially designed for confirmation of risk factors in the development of a specific disease condition,28 and it is used for evaluating the effectiveness of multiple treatments for a disease in the recent 5 years.29–31 We therefore assume that it is essential to use this method to screen and find out treatments that are with convincing evidence for haemorrhoidal management.

The result of this review will be published in a peer-reviewed journal, and we believe that the result will benefit clinical practitioners, patients and policy-makers.

**Ethics and dissemination**

The study is an umbrella review, which requires no ethical approval. We will present the results of the umbrella review at conferences and publish the final report in a peer-reviewed journal.

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**Contributors** MC and HZ designed the study and will perform data analysis, DQ developed the search strategy. MC and T-CT will perform literature search, screen the eligibility of the retrieved articles. T-CT and DQ will develop the information for data extraction. MC wrote the first draft of the protocol, and all authors read the article and approve it for publication.

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**Disclaimer** Funders and sponsors have no role in the design of this protocol.

**Competing interests** None declared.

**Patient and public involvement** Patients and/or the public were involved in the design, or conduct, or reporting, or dissemination plans of this research. Refer to the Methods section for further details.

**Patient consent for publication** Not required.

**Provenance and peer review** Not commissioned; externally peer reviewed.

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