

## **Health Technology Assessment in Traditional and Complementary Medicine: a Scoping Review of International Activity and Examples of Acupuncture**

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**Short title:** A Scoping Review of HTA in T&CM

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## **Abstract**

**Background:** Traditional therapies are of high value in maintaining and improving human well-being. China's healthcare policymakers are attempting to harness health technology assessment for traditional therapies to inform pricing and reimbursement. The value assessment framework for Chinese patent medicine has been developed and is being adopted and validated widely by research institutions. Subsequently, the healthcare decision-makers particularly hanker for the development of the value framework of traditional non-pharmacological therapies.

**Methods:** To construct a practical value framework for traditional non-pharmacological therapies, a scoping review methodology was adopted to identify the evaluation domains and obstacles. A comprehensive search, screening, and analysis process was conducted according to the Preferred Reporting Items for Systematic Reviews and Meta-Analyses extension for Scoping Reviews. Evidence was retrieved from scientific databases and HTA agencies' websites.

**Results:** The search strategy identified 5 guidelines or standards records and 17 acupuncture HTA reports. By synthesizing the valuable reports of Chinese patent medicine and acupuncture evaluation in major countries and regions, this study found that Mainland China was promoting the comprehensive value assessment of Chinese patent medicine, whereas the United Kingdom, Singapore, Canada, the United States, and Malaysia had carried out the health technology evaluation of acupuncture for various conditions among which chronic pain was the most common condition. UK and Singapore applied the HTA results to support acupuncture reimbursement decisions. Three domains, including safety, effectiveness, and economy, were

commonly adopted by HTA agencies. Currently, the biggest challenge of evaluating traditional non-pharmacological therapies comes from the scarce high-quality clinical evidence, leading to high uncertainty in the evaluation of effectiveness and cost-effectiveness.

**Conclusions:** This study identified value evaluation domains and issues of traditional therapies, and pointed out future research implications, to promote the development value framework of TCIM and evidence-informed policy-making.

**Keywords** Traditional and Complementary Medicine; Health technology assessment; Acupuncture

## 1. Introduction

Traditional, Complementary, and Integrative Medicine (TCIM) has demonstrated its unique advantages and values in improving people's health, and 170 of 194 (88 percent) WHO member states have admitted the formal use of TCIM within their health systems in an investigation<sup>1,2</sup>. TCIM is an integrated conception, encompassing three parts: “traditional” therapy is viewed as a total sum of historical knowledge and beliefs unique to different ethnic groups, “complementary” therapy is defined as a non-mainstream practice used together with conventional medicine, while an “integrative” therapy is a combination of complementary approaches used in conjunction with conventional medicine<sup>3</sup>. Among these, acupuncture has witnessed thriving and prosperous worldwide, though this technique was originally a feature of traditional Chinese medicine. According to the WHO’s survey including 129 countries, 80% of them recognized the use of acupuncture, with 18 countries reimbursing acupuncture in their healthcare service system<sup>1</sup>.

To promote the high-quality development of TCIM, plenty of policies touching on clinical efficacy evaluation and value-based health technology evaluation have been introduced in China. Those evaluation systems are designed to showcase the theories, features, and value of TCIM, and eventually to inform the reimbursement decision-making and pricing of TCIM services<sup>4</sup>. Furthermore, the value assessment framework for Chinese patent medicine has been issued by an academic institution. Whereas, there is no value framework for non-pharmacological therapies up to now. Therefore, Chinese healthcare decision-makers are pretty eager to use such tools to make categorical management policies.

Apparently, to galvanize the reviving of TCIM, it is imperative to incorporate more traditional medicine services into health insurance coverage based on the high-quality value evaluation evidence<sup>5</sup>. Consequently, It is necessary and valuable to conduct a scoping review of HTA in TCIM to have a general understanding of the evaluation dimensions and challenges. Given the wide adoption of acupuncture, the health technology assessment (HTA) reports of acupuncture can be more easily accessible than those of other traditional non-pharmacological therapies. It can bring more practical insights to develop the value assessment framework of traditional non-pharmacological technology by reviewing and synthesizing value domains and weaknesses mentioned in HTA reports of acupuncture treated for various diseases. We hope that the value framework can conduce to elucidate the potential value of traditional non-pharmacological therapies for disease prevention, treatment and rehabilitation, and informing healthcare and reimbursement policy-making in China.

## **2. Methods**

### **2.1 Study Design and Search Strategy**

To outline the evaluation framework of TCIM informing the value assessment and reimbursement decision-making, a scoping review of the HTA guideline for TCIM and HTA reports for acupuncture was conducted, obeying the methodological framework introduced by Arksey and O'Malley and refined by the Joanna Briggs Institute<sup>6-8</sup>.

The Preferred Reporting Items for Systematic Reviews and Meta-Analyses extension for Scoping Reviews (PRISMA-ScR) was adopted to elaborate this study. Health technology assessment, traditional medicine, complementary medicine, acupuncture, and other similar

terms were used as keywords to construct the search strategy, retrieving the published literature on health technology evaluation in TCIM from relevant electronic databases, such as Medline (PUBMED), EMBASE, Web of Science, the International HTA database hosted by INAHTA, CNKI, Wanfang, and HTA agencies including National Institute for Health and Care Excellence (NICE), Agency for Care Effectiveness (ACE), Canadian Agency for Drugs and Technologies in Health (CADTH), Institute for Clinical and Economic Review (ICER), Scottish Medicines Consortium (SMC), Swedish Agency for Health Technology Assessment and Assessment of Social Services (SBU) from inception to January 2023. The full search strategy can be found in Supplement 2. The scoping review protocol was not registered in a registry.

## **2.2 Study Selection**

Given the purpose was to put attention to the HTA domains of TCIM and the HTA difference between TCIM and modern medical products, studies focusing on guidelines or standards specific to TCIM were included. In consideration of scarce assessment guidelines of TCIM, HTA reports of acupuncture for any diseases in the context of HTA organizations or agencies or other bodies worldwide were included as well, used to summarize HTA domains and characteristics of TCIM. HTA reports written in Chinese or English and available full texts were included. The detailed inclusion and exclusion criteria were shown in Table 1. The preliminary screening by titles and abstracts as well as the final screening by full texts was performed by two independent researchers to determine the eventually included literature and potential disagreements were solved by a third author.

## **2.3 Data Extraction**

A predefined qualitative data form encompassing report characteristics, HTA agency, country, year of publication, HTA domains, population, intervention and comparator, results, limitations and implications were used to extract key information by two independent researchers.

## **2.4 Data Synthesis**

A descriptive analysis and a narrative synthesis were performed employing tables to synthesize the HTA domains and the main characteristics of each report.

## **3. Results**

### **3.1 Reports Selection**

Overall, of the 993 records retrieved after data deduplication, 971 were screened out since they were studies without acupuncture as an intervention, studies focusing on a narrative review of HTA in traditional medicine, or no guideline or consensus. Finally, 5 guideline or consensus records from China<sup>9-13</sup> and 17 acupuncture HTA reports from international HTA agencies<sup>14-30</sup> were included in the review. The full selection process is depicted in Figure 1.

### **3.2 Description of included studies**

#### **3.2.1 Recapitulation of HTA guidelines or consensus in TCIM**

5 guideline or consensus records<sup>9-13</sup> about value assessment of Chinese patent medicine released by Chinese academic groups were identified, while there were no published HTA guidelines or consensus for TCIM in other countries and regions. In 2021, the China National Health Commission (NHC) issued the Management Guideline for the Clinical Comprehensive



Evaluation of Drugs, which involved six domains (safety, effectiveness, economy, innovation, suitability, accessibility) to measure the value of a drug based on the methods of health technology assessment and multiple-criteria decision analysis to improve clinical rational use of medicines<sup>31</sup>. Since then, China healthcare sectors and academic groups have been promoting clinical comprehensive evaluation of drugs. With that management guideline, in the field of TCIM, the Chinese Association of Traditional Chinese Medicine, the Chinese Academy of Chinese Medical Sciences and other institutions drafted or promulgated technical evaluation guidelines or technical specifications, and carried out clinical comprehensive evaluation of specific Chinese patent medicines, of which the evaluation dimensions were basically consistent with management guidelines issued by NHC<sup>9-13</sup>. Additionally, some institutions added up Chinese medicine-specific indicators, such as the traditional Chinese medicine theory, human use experiences, etc<sup>11, 12</sup>. In another technical guideline developed by the Institute of Clinical Basic Medical Sciences of the Chinese Academy of Chinese Medical Sciences, the three dimensions, namely safety, effectiveness, and economy, were drawn up from the national clinical comprehensive evaluation, and the other dimensions (application, science and standards) reflecting the features of TCM were proposed by Delphi method (see Table 2)<sup>10, 13</sup>.

Internationally, even though some HTA of TCIM were conducted to provide evidence for medical insurance coverage decision-making, there was currently no published value assessment framework or economic evaluation method aimed at traditional medicine or non-pharmacological therapies in other countries and regions. South Korea, Switzerland, Singapore, and the United Kingdom had attempted to apply HTA for traditional and complementary

medicine to inform medical insurance reimbursement. In Korea, *Hanbang* services have been covered in the benefits schedule in the country's National Health Insurance (NHI) since 1987. Currently, health insurance benefits of *Hanbang* are especially focused on treatments such as acupuncture, moxibustion, and cupping, while herbal medicines coverage is relatively rare. To promote *Hanbang*, it is necessary to prioritize the extension of health benefits to herbal medicines and traditional techniques with a lower rate of coverage. Consequently, the National Development Institute of Korean Medicine (NIKOM) had developed a strategic plan that includes the development of economic valuation guidelines specific to *Hanbang* services under the current HTA system. In line with the evidence-based decision-making and value purchase principles, the government led the formulation of *Hanbang* clinical practice guidelines for services, the first batch of which contained 27 diseases<sup>5</sup>. In Switzerland, reimbursement for traditional and complementary medicine after 2017 will depend on the results of the evaluation projects and international health technology assessments<sup>2</sup>. A systematic review of technical assessments of traditional medicine conducted by HiTAP, which included peer-reviewed articles, grey literature, and regional or international guidelines on the evaluation of herbal products and traditional and complementary medical practices found there was no organized framework to adapt economic evaluation methods to the unique features of TCIM. The reporting of costs and health-related quality of life remained uncommon in TCIM studies. There was a call for updating the guidelines on the evaluation of TCIM and developing better evaluation frameworks<sup>32</sup>.

### 3.2.2 Examples of Health Technology Assessment in Acupuncture

The scoping review identified 17 published acupuncture HTA reports from five countries, including the United Kingdom (8)<sup>22-29</sup>, the United States (3)<sup>15, 16, 21</sup>, Canada (3)<sup>17, 18, 30</sup>, Singapore (1)<sup>14</sup>, Malaysia (1)<sup>20</sup>, and one international academic organization, Cochrane (1)<sup>19</sup>, between 2003 and 2022. The involved conditions were mainly chronic pain, and otherwise included allergic rhinitis, induction, irritable bowel syndrome, stable angina. Usual care or a sham/placebo acupuncture were the usual comparators. Detailed characteristics of all included reports were provided in Table 3.

The evaluation dimensions mainly contained safety, effectiveness, cost-effectiveness, medical resources consumed, and guideline recommendations (see Table 4). Agency for Care Effectiveness (ACE), a Singaporean HTA organization, also took three other factors, the clinical needs and disease characteristics, organizational feasibility, and ethical or social issues, into consideration<sup>14</sup>. The assessment report of the Institute for Clinical and Economic Review (ICER) in US mentions health insurance coverage, other important benefits or risks<sup>21</sup>. Generally speaking, all evaluation bodies attached importance to the quality of the evidence of safety and effectiveness. In Singapore, guided by the medical device technology guidelines, ACE conducted the evaluation of acupuncture, which predicted the resources utilization with the technology's annual cost and the number of patients who may benefit from it. The organizational feasibility assessment aimed to identify barriers and facilitate technology's adaptation into their public healthcare system, and any organizational factors that might influence the technology's performance or use in clinical practice.

The clinical comparative effectiveness results varied from different patient population in the comparison of acupuncture with placebo or sham acupuncture<sup>14-30</sup>. In the time of only synthesizing high-quality clinical evidence, acupuncture was more effective than sham acupuncture or usual care for all of these chronic pain conditions. However, there were plenty of low-quality clinical studies, resulting the inconclusive conclusions. Methodological flaws leading to the generally downgraded evidence quality, was pointed out most frequently. The evidence quality was typically downgraded due to risk of bias and imprecision, like small sample size, inappropriate randomized sequence generation, lack of allocation concealment, inadequate level of blinding, various types of sham acupuncture, considerable heterogeneity among studies, poor reporting of interventions and comparators treatment details, high placebo response, and non-standardized additional treatments, which finally resulted in unclear and inconsistent clinically relevant benefits. Plus, subjective measures also lacked standardization, clinical validation, length of appropriate follow-up, and minimum clinically important difference (MCID). Typically, the continuous outcome results focused only on statistically significant changes, so it was difficult to interpret the clinical significance of average changes in measures of function, quality of life, and pain. Furthermore, in these studies, treatment was for chronic pain but rarely included long-term follow-up.

Otherwise, the adverse events data for these trials was limited as this was generally found in small studies with a short follow up time and so it was unclear whether this was representative of the events expected to be seen in real life practice. And serious adverse reactions were scarce. Considering the beforementioned limitations about clinical effectiveness, there was a limited

evidence base, and thus many extrapolation assumptions applied in the cost-effectiveness analysis models were lack of robust evidence. The cost-effectiveness results of acupuncture compared with no treatment were inconsistent among various studies, and by limiting the treatment session and duration, acupuncture was likely to become a cost-effective intervention within the country's threshold<sup>25,27</sup>.

## **4. Discussion**

### **4.1 Summary of Main Findings**

Overall, there were still no other HTA guideline or consensus of TCIM besides clinical comprehensive evaluation guideline published by academic agencies in China. Typically, the evaluation of TCIM adhered to general guidelines drafted by HTA agencies. The evaluation domains included safety, effectiveness, economy (involving cost effectiveness, consumed medical resources), innovation, suitability, accessibility, specialties of TCM, application, science, standards, guidelines recommendations, clinical needs and disease features, organizational feasibility, health insurance coverage, among which safety, effectiveness, and economy had the biggest frequency.

The role and function of TCIM HTA in policy-making varied across different countries. Despite the fact that forty-five of the WHO member states currently had health insurance coverage for traditional medicine and T&CM practices (e.g., acupuncture and chiropractic) <sup>2</sup>. Few countries or regions applied HTA to support medical insurance access to traditional medical services<sup>33</sup>, and countries with well-developed HTA knowledge translation mechanism such as the United Kingdom, Singapore and Switzerland are exploring the use of HTA to

support medical insurance access for acupuncture<sup>2,14, 22-29</sup>. With the need to cover more traditional medical technologies in insurance coverage, evidence-based assessment of its budget impact and value for money would become a matter of growing importance. There was an appeal to focus on developing better frameworks for evaluating TCIM in order to obtain regulatory approval and/or reimbursement<sup>32</sup>.

Compared with other narrative reviews about the status and challenges of Health technology assessment and economic evaluation in traditional Chinese medicine and acupuncture, this study qualitatively synthesized the value evaluation domains and issues by systematic retrieving evaluation guidelines of TCIM and HTA reports of acupuncture according to the methodology framework of scoping review. It was the first time to synthesize the value evaluation domains of TCIM, even though those domains were mostly consistent with general drug or medical device technology evaluation domains. However, some researchers were appealing and making efforts to add traditional medical related factors to value framework<sup>10-13, 32</sup>. Currently, it was evident that economic evaluation played a limited role in reimbursement decisions for acupuncture<sup>34</sup>. Given that specific healthcare system, costing, healthcare service pathway, and population health preferences, it was difficult to generalize the economic evaluation results to other countries. In terms of the issues, apart from the study design, lack of long-term effectiveness, and poor reporting quality, previous studies also depicted other challenges, like cost measurement, which was difficult to accurately collect due to dosage and duration adjustment based on the symptoms and characteristics of the patient at each visit, self-pricing and unpublished price, and PICO structuring, among which “Intervention” was not easy

to be determined because the number and dosage of traditional medical service were adjusted based on patients' characteristics and doctors' experience<sup>33, 35</sup>.

#### **4.2 Limitations**

There were also several limitations in this article. First of all, studies might have been missed due to several peer-reviewed databases and influential HTA agencies websites were searched, and peer-reviewed publications or HTA reports released by HTA agencies were considered only if they were written in English or Chinese. Therefore, the included literatures might not be completely representative of the research available. In addition, the study limitations were mainly due to the applicability of generalizing acupuncture assessment domains to traditional non-pharmacological technology assessment, especially considering the specialities of other Chinese medical technology, like tuina, acupotomy and so on. Furthermore, some included HTA reports might contain inadequate information, leading to insufficient data extraction of the study characteristics of interest, which also restricted the extrapolation of results.

#### **4.3 Future Research and implication**

It can be expected value domains of TCIM could expedite the generation and accumulation of high-quality primary evidence. As human-use experience plays an extremely important role in the development of TCIM, real-world diagnosis and treatment data have significant advantages in the effectiveness evaluation of TCIM, which also conduce to address the questions about blindness in RCT and complex peculiarity of traditional non-pharmacological technology to enhance the quality of clinical effectiveness evidence. The pragmatic randomized

clinical trials and other observational studies are very suitable to consider the features of TCIM, which can not only standardize, but also reflect the characteristics of dialectical treatment of Chinese medicine. Recently, eRCT-pRCT-RCDOS-REGOS<sup>\*</sup>, stepwise progressive clinical effectiveness evaluation system of TCIM was proposed, aimed to enhance study quality by broadening the study populations and interventions to assess the net benefits in more contexts, such as practitioner proficiency, equipment quality, etc<sup>36</sup>. For continuous outcomes, when using a measurement, researchers should report both statistical and clinical significance, taking MCID into consideration. Considering the widely usage of subject clinical outcomes in the clinical value assessment of TCIM, the study regarding MCID of clinical outcome measures should be encouraged. Categorical measures reporting the proportion of patients achieving a clinically meaningful improvement in function, quality of life, and pain are more useful and should be reported in addition to average group changes. Those factors are beneficial for generating high-quality value evidence for TCIM.

## 5. Conclusion

This scoping review provided an overview of the HTA guideline of TCIM and acupuncture HTA across multiple countries. With China vigorously promoting the inheritance and innovation of traditional Chinese medicine and increasingly emphasizing evidence-based decision-making, the demand for health technology evaluation of TCIM is on the rise, providing evidence for the price adjustment of traditional medical services and the coverage of Chinese

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\* eRCT, explanatory randomized controlled trial;  
pRCT, pragmatic randomized controlled trial;  
RCDOS, observational studies using routinely collected data, observational studies using routinely collected data;  
REGOS, observational studies using patient registry data, is an observational study using patient registry data.



patent medicine in national basic medical insurance. Currently, domestic studies about comprehensive value assessment of Chinese patent medicine are on the rise, while few studies address the question of how to assess the value of traditional non-pharmacological technology in order to pursue the decision-makers to purchase traditional and complementary medical services widely. International HTA agencies had published several evaluation reports of acupuncture for various conditions, and the results were used to inform medical insurance reimbursement decisions. The most commonly assessment domains are clinical effectiveness, safety and economy. In the process of evidence synthesis, however, some issues, such as poor clinical evidence quality, sophisticated costing, lack of long-term effectiveness, and so on, set obstacles to conclude robust value evaluation results for traditional non-pharmacological technology. In the future, more efforts should be put on how to transform tons of human-usage experience in real-world clinical practice into high-quality evidence, and it is a valuable topic to determine if economic evaluation reference case is needed for traditional non-pharmacological technology considering these unique features like short-term intervention and potential long-term benefits for physical and mental health, in order to improve study quality and inform scientific decision-making.

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### **Conflict of interest**

The authors declare that they have no conflicts of interest.

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### **Ethical statement**

Not applicable.

### **Data availability**

Not applicable.

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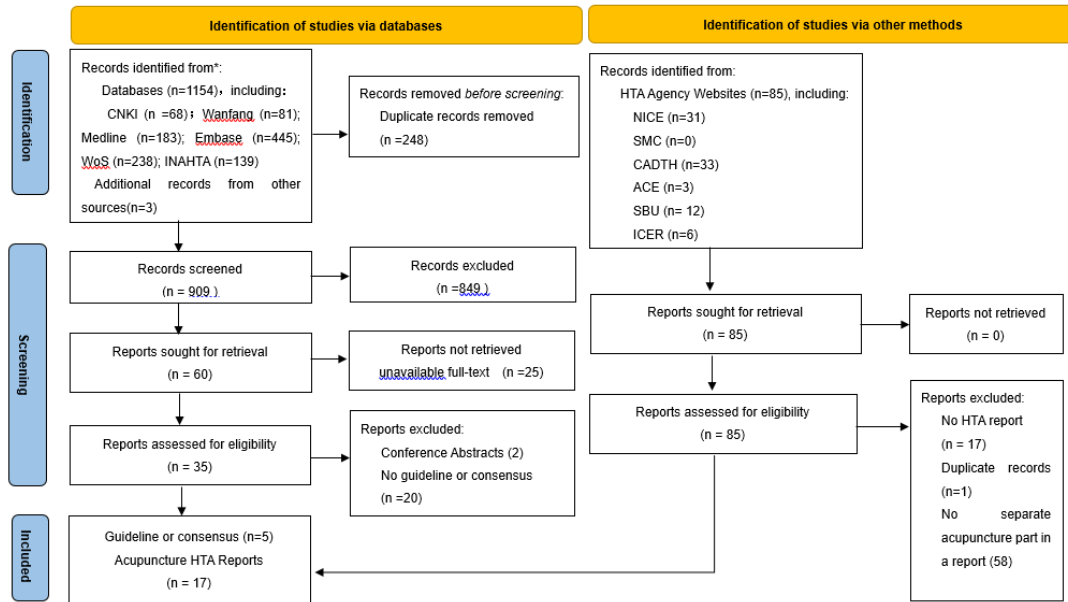
SUPPLEMENTARY FILES

Supplement 1: PRISMA-ScR Checklist

Supplement 2: Search Strategy

FIGURES

Figure 1: PRISMA flow diagram



## TABLES

Table 1 Inclusion and Exclusion Criteria

Table 2 Information table for comprehensive evaluation of Chinese patent medicine

Table 3 Summary Characteristics of Included Acupuncture HTA Reports

Table 4 HTA domains of acupuncture evaluation

Table 1 Inclusion and Exclusion Criteria

	Inclusion criteria	Exclusion criteria
Intervention	<ul style="list-style-type: none"> <li>● Guideline: specific to Traditional, Complementary, and Integrative Medicine</li> <li>● Case study: Acupuncture or dry needling</li> </ul>	General guidelines
Study type	<ul style="list-style-type: none"> <li>● Guidelines or consensus standards for TCIM</li> <li>● HTA reports for acupuncture</li> </ul>	Systematic reviews or Economic evaluations or for single traditional technology
Context	HTA agencies or bodies worldwide	-
Language	Chinese, English	Any other languages

Table 2 Information table for comprehensive evaluation of Chinese patent medicine

Author, year	Safety	Effectiveness	Economy	Innovation	Suitability	Accessibility	Specialties of TCM	Application	Science	Standards
YUAN, 2023 <sup>9</sup>	✓	✓	✓	✓	✓	✓				
ZHANG, 2021 <sup>11, 12</sup>	✓	✓	✓	✓	✓	✓	✓			
ZHANG, 2022 <sup>10, 13</sup>	✓	✓	✓					✓	✓	✓

Table 3 Summary Characteristics of Included Acupuncture HTA Reports

Country	Year	HTA organization	Population	Intervention	Comparator	Outcome Main results	challenges and issues
UK <sup>25</sup>	2021	NICE	Chronic primary pain	<ul style="list-style-type: none"> <li>Acupuncture/dry needling</li> <li>Electro acupuncture</li> </ul>	<ul style="list-style-type: none"> <li>Placebo/sham</li> <li>Usual care</li> </ul>	-The recommendation was caveated with information on the number of hours of staff time, and banding of staff that would make the delivery of acupuncture cost-effective.	<ul style="list-style-type: none"> <li>-Low to very low evidence quality.</li> <li>-Lack of blinding</li> <li>-Mostly subjective outcomes resulted in a high risk of performance bias.</li> <li>-Uncertainties about the cost-effectiveness of acupuncture, as there was a limited evidence base and there were uncertainties around the cost of the intervention.</li> <li>-High to very low evidence quality due to inconsistency and imprecision.</li> <li>-Inconsistent types of sham acupuncture between studies, increasing the uncertainty of the results.</li> <li>-Complex nature of acupuncture treatment: the effect of the relationship between the health care professional and the person with osteoarthritis.</li> </ul>
UK <sup>28</sup>	2022	NICE	Osteoarthritis	<ul style="list-style-type: none"> <li>Acupuncture/dry needling</li> <li>Electro-acupuncture</li> </ul>	<ul style="list-style-type: none"> <li>Sham</li> <li>No intervention</li> </ul>	-The adverse events data for these trials was limited as this was generally found in small studies with a short follow-up time.	<ul style="list-style-type: none"> <li>-Complex nature of acupuncture treatment: the effect of the relationship between the health care professional and the person with osteoarthritis.</li> </ul>

UK <sup>27</sup>	2021	NICE	Primary headaches	<ul style="list-style-type: none"> <li>• Acupuncture</li> <li>• Sham</li> </ul>	<p>-Some evidence for improvements in headache days and responder rate.</p> <p>-Treatment reactions after acupuncture needling were common. The risk of serious side effects was low.</p> <p>-Acupuncture was cost-effective when compared to no treatment in people with migraine or tension-type headache (£ 12,381/QALY) from a published literature based on an RCT.</p> <p>-An original cost-effectiveness analysis showed that acupuncture was more cost-effective than no treatment when 10 or fewer sessions were provided.</p>	<p>-Low and very low-quality evidence</p> <p>-Economic evidence had minor limitations and partial applicability.</p>
UK <sup>24</sup>	2017	NICE	Irritable bowel syndrome (IBS)	<ul style="list-style-type: none"> <li>• Single acupuncture</li> <li>• Combination acupuncture</li> <li>• Sham/placebo</li> <li>• Other type of treatment</li> </ul>	<p>-No significant effect of acupuncture on IBS global symptoms, pain, and quality of life compared with placebo.</p>	<p>-Limited evidence of potentially serious adverse effects associated with acupuncture treatments.</p>
UK <sup>26</sup>	2021	NICE	Women undergoing induction	<ul style="list-style-type: none"> <li>• Acupuncture</li> <li>• Usual care</li> </ul>	<p>-Evidence was insufficient to determine the effectiveness of acupuncture in cervical priming/induction of labor.</p> <p>-The evidence was mixed: One RCT showed some improvement in angina and exercise test variables when compared to tablet placebo. However, there was no improvement</p>	<p>-Insufficient evidence</p>
UK <sup>23</sup>	2016	NICE	Stable angina	<ul style="list-style-type: none"> <li>• Acupuncture</li> <li>• Sham</li> </ul>	<p>-The evidence was mixed: One RCT showed some improvement in angina and exercise test variables when compared to tablet placebo. However, there was no improvement</p>	<p>-Small sample size (&lt;50 patients)</p> <p>-Without long-term follow-up outcomes.</p>

						in angina or exercise test variables in two RCTs that compared acupuncture to sham	-The methodology of the trials was not well reported and the derived data was not analyzable.
						-Acupuncture was more effective than sham acupuncture, and was better than standard medical care for all of these chronic pain conditions when based on high-quality trial evidence.	
UK <sup>22</sup>	2017	NIHR	<ul style="list-style-type: none"> <li>•Osteoarthritis</li> <li>•Headaches</li> <li>•Shoulder pain</li> <li>• Back or neck pain</li> </ul>	<ul style="list-style-type: none"> <li>• Acupuncture</li> </ul>	<ul style="list-style-type: none"> <li>• Sham</li> <li>• non-acupuncture</li> </ul>	<p>-Acupuncture was also cost-effective if only high-quality trials were analyzed. When synthesizing all trials, including both low- and high-quality trials, transcutaneous electrical nerve stimulation was cost-effective.</p>	-Limited data regarding the long-term effects of many non-pharmacological interventions
Singapore <sup>14</sup>	2020	ACE	<ul style="list-style-type: none"> <li>•Low back pain and neck pain</li> </ul>	<ul style="list-style-type: none"> <li>• Acupuncture ± standard care</li> </ul>	<ul style="list-style-type: none"> <li>• Standard care</li> <li>• Sham</li> <li>• No treatment</li> </ul>	<p>- The effectiveness of acupuncture was unclear for chronic arthritis, osteoarthritis, shoulder-arm syndrome, and epicondylitis when compared with alternative treatments.</p> <p>-The overall applicability of economic evaluation estimates to the local context was unclear</p>	<ul style="list-style-type: none"> <li>-Poor quality of evidence</li> <li>-Small sample size</li> <li>-Unclear clinically relevant benefits, studies</li> <li>-Considerable heterogeneity among studies</li> <li>-Poor reporting of acupuncture treatment details</li> </ul>
Canada <sup>18</sup>	2019	CADTH	<ul style="list-style-type: none"> <li>• Chronic non-cancer pain</li> </ul>	<ul style="list-style-type: none"> <li>• Acupuncture</li> </ul>	<ul style="list-style-type: none"> <li>• Pharmacological interventions</li> </ul>	<p>- The clinical effectiveness results and recommendations were variable depending on the patient population. Acupuncture decreased</p>	<ul style="list-style-type: none"> <li>- Few high-quality primary studies and many low-quality primary studies.</li> </ul>



				<ul style="list-style-type: none"> <li>• No treatment</li> <li>• Usual care</li> </ul>	<p>pain intensity in chronic prostatitis/chronic pelvic pain syndrome, chronic headache, chronic neck pain, chronic shoulder pain, sciatica, myofascial pain, hip osteoarthritis, knee osteoarthritis, and osteoarthritis when compared to sham. For other pain conditions, the evidence regarding the clinical effectiveness was mixed.</p> <p>- No firm conclusions can be made regarding the relative cost-effectiveness.</p>	<p>- High risk of bias: non-randomized or unconcealed allocation, lack of blinding or maintenance, heterogeneous population or treatment, non-standardized additional treatments, or even insufficient reporting of risk of bias items.</p> <p>- Outcomes lack of standardization, clinical validation, length of appropriate follow-up, and minimum clinically important difference.</p> <p>-Economic studies from outside of Canada might be less applicable within Canada.</p>
Canada <sup>17</sup>	2016	CADTH	<ul style="list-style-type: none"> <li>• Musculoskeletal pain</li> <li>• Joint disorders</li> <li>• Joint pain</li> <li>• Derangement of joints</li> <li>• Chronic tendinosis</li> <li>• Tendinopathy</li> </ul>	<ul style="list-style-type: none"> <li>• Dry needling ± injection</li> <li>• Usual care</li> </ul>	<p>No comparator</p> <p>- Insufficient evidence on the effects of dry needling on other outcomes, such as range of movement or quality of life.</p>	<p>-Most methodologically flawed primary studies: inadequate blinding, underpowered to accurately indicate treatment effects, heterogeneous patient populations, various outcome measures and length of follow-up, and non-standardized treatment and comparator interventions</p>

US <sup>21</sup>	2017	ICER	<ul style="list-style-type: none"> <li>• Chronic low back or neck pain</li> </ul>	<p>Cognitive and mind-body therapies, including:</p> <ul style="list-style-type: none"> <li>• Acupuncture</li> <li>• CBT</li> <li>• MBSR</li> <li>• Yoga</li> <li>• Tai chi</li> </ul>	<ul style="list-style-type: none"> <li>• Usual care</li> <li>• Sham/placebo</li> </ul>	<p>-Small to moderate improvements in function and pain compared with usual care immediately following the completion of therapy.</p> <p>-The differences in outcomes were smaller and often non-significant clinically when compared to sham acupuncture, suggesting that much of the benefit may be from the placebo effect.</p> <p>- The incremental cost of achieving one case of improved pain over the five-year time horizon relative to usual care ranged from \$660 for yoga to approximately \$15,800 for CBT (&lt;\$100,000/QALY).</p>	<p>-MCID was not defined, and the results focused only on statistically significant changes.</p> <p>-Difficult to interpret the clinical significance of average changes in continuous measures of function, quality of life, and pain.</p> <p>-Did not model varying treatment effectiveness over time due to the deficit of long-term trial data, did not model subsequent lines of intervention for individuals who experienced a recurrence of low back pain due to a lack of published evidence on this estimate.</p>
Malaysia <sup>20</sup>	2020	MaHTAS	<p>Neurological disorders (including headache, refractory neuralgia, Bell's palsy,</p>	<ul style="list-style-type: none"> <li>• Acupuncture</li> <li>• Traditional body needling</li> <li>• Moxibustion</li> <li>• Electro-acupuncture</li> </ul>	<ul style="list-style-type: none"> <li>• Sham/placebo</li> <li>• Scalp acupuncture</li> <li>• Other conventional treatment</li> </ul>	<p>-None of the clinical trials reported any serious adverse events.</p> <p>-Acupuncture was suggested to be clinically relevant benefit and cost-effective in certain sectors.</p>	<p>-Poor quality of the included trials: high risk of bias due to inappropriate randomized sequence generation, lack of allocation concealment, inadequate level of blinding, and poor description of patient withdrawals</p>

			post-stroke, Guillain Barre and transverse myelitis)	• Laser acupuncture			from the studies.
Canada <sup>30</sup>	2022	WorkSafeBC	Lateral Elbow Pain	• Dry Needling	• Sham • Other active treatments • No comparator	- No high-quality evidence supporting the efficacy and/or effectiveness of dry needling,	-Most studies with very low to low study quality: methodological limitations and imprecision, low numbers of participants, and inadequate blinding.
International organization <sup>19</sup>	2018	Cochrane	Carpal tunnel syndrome	• Acupuncture • Acupuncture-related interventions	• Sham • Other active treatments	-Acupuncture was associated with no serious adverse events, or reported discomfort, pain, local anesthesia and temporary skin bruises.	-Low or very low evidence -Most evidence was short-term.
UK <sup>29</sup>	2007	WMHTAC	Allergic rhinitis	• Acupuncture	• Sham ± standard care	-Insufficient evidence on the clinical effectiveness to support or refute its use in patients with allergic rhinitis. -Acupuncture was not associated with any additional adverse events.	- Poor quality and the inconclusive results
US <sup>15</sup>	2003	AHRQ	Fibromyalgia	• Acupuncture • Electro-acupuncture	• Sham	-Insufficient evidence to conclude that acupuncture had efficacy for the treatment of fibromyalgia.	- Small sample size - Inappropriateness of outcome measures -Complexities of acupuncture: different types of acupuncture, different systems for choosing sites,

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						and variability in the technique of needle insertion and manipulation
						- Small sample size
						- Underpowered to measure statistically significant differences between the interventions
US <sup>16</sup>	2003	AHRQ	Osteoarthritis	<ul style="list-style-type: none"> <li>• Acupuncture</li> <li>• Electro-acupuncture</li> </ul>	<ul style="list-style-type: none"> <li>• Sham</li> <li>• Other active treatments</li> </ul>	<ul style="list-style-type: none"> <li>-Most studies did not find a benefit for acupuncture compared to sham acupuncture.</li> <li>-Lack of blinding</li> <li>-Lack of description of the handling of dropouts and withdrawals</li> <li>-No formal test statistics</li> </ul>

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ACE, Agency for Care Effectiveness; AHRQ, Agency for Healthcare Research and Quality; CADTH, Canadian Agency for Drugs and Technologies in Health; ICER, Institute for Clinical and Economic Review; MaHTAS, Malaysian Health Technology Assessment Section; NICE, National Institute for Health and Care Excellence; NIHR, National Institute for Health Research; WMHTAC, West Midlands Health Technology Assessment Collaboration.

Table 4 HTA domains of acupuncture evaluation

HTA organization	National health system	Safety	Efficacy/ effectiveness	Cost effectiveness	Consumed medical resources	Guidelines recommendations	Clinical needs and disease features	Organizational feasibility	Health insurance coverage
ACE <sup>10</sup>	Public	✓	✓	✓	✓	-	✓	✓	-
NICE <sup>19-24</sup>	Public	✓	✓	✓	✓	✓	-	-	-
CADTH <sup>13, 14</sup>	Public	✓	✓	✓	-	✓	-	-	-
ICER <sup>17</sup>	Private	✓	✓	✓	✓	✓	-	-	✓
MaHTAS <sup>16</sup>	Public	✓	✓	✓	-	-	-	-	-