Economic evaluations of AI-assisted technology in healthcare: how are we assessing the cost-effectiveness of these new therapies?

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## Introduction

- > Artificial Intelligence (AI)-assisted technologies are being applied across a wide range of healthcare areas promising to enhance clinical outcomes, efficiency, and decision-making. As the deployment of these technologies accelerates, understanding their economic implications will become increasingly critical.
- > Economic evaluations (EEs) of AI-assisted technologies in healthcare are growing in number, with a parallel increase in systematic literature reviews (SLRs) that synthesize and assess these evaluations. This aligns with recent developments, such as the introduction of the CHEERS-AI (Consolidated Health Economic Evaluation Reporting Standards for AI) guidelines and updates to the NICE (National Institute for Health and Care Excellence) evidence standards framework (ESF) for digital health technologies, which now include evidence standards specific to AIbased technologies.<sup>1,2</sup>
- > The objective of this study was to understand the current value proposition of Alinterventions in healthcare, to identify gaps between existing evaluations and the

## **Results (continued)**

> Six SLRs were included, all published after 2020. The key characteristics are summarised in Table 2.<sup>4-9</sup>

## Table 2. Key characteristics of included studies

Search outcomes	Results (n=6) <sup>4-9</sup>	
Databases searched	Embase (4), Cochrane Central (3), Web of Science (2), Scopus (2), Google search (2), HTA database (1)	
Period searched	From inception (2), 5 years (3), unclear (1)	
Language covered	English (6)	
Funding source	Public organisation (3), No financial support (3); 1 not claimed	
Conflict of interest	All 6 declared no conflict of interest	
Disease area covered	General medicine/healthcare (3), Ophthalmology (3)	
Type of Al-interventions	General medicines/healthcare focused search identified a breadth of AI-intervention including but not limited to pattern recognition, risk prediction models, and monitoring. Ophthalmology only identified EEs related to AI driven diagnosis and screening intervention.	

A total 74 EEs were included in the six SLRs. After removing 21 duplications, 53 unique studies were identified, including 16 trial based and 37 model based EEs.





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requirements outlined by CHEERS-AI and NICE, and to highlight key considerations for future economic evaluations of emerging AI-driven technologies.

## Methods

- > An umbrella review, a comprehensive synthesis of evidence from multiple systematic reviews, was performed on SLRs of EEs of AI-assisted interventions.
- > The ISPOR good practice for critical appraisal for SLRs with cost and costseffectiveness outcomes (SR-CCEO) was followed.<sup>3</sup>
- > Medline, EMBASE, Cochrane Central and NICE HTA databases were searched to identify SLRs of EEs for AI-assisted interventions.
- > Publications were included based on predefined inclusion and exclusion criteria (Table 1).

## Table 1. Inclusion and exclusion criteria

Types of criteria	Inclusion	Exclusion
Population	Actual or hypothetical patients who receive AI-assisted healthcare interventions	Patients receive healthcare without Al- technology involved
Intervention/ Comparator	Al-assisted healthcare interventions (for diagnosis, monitoring, surgery or other treatments) compared with standard/usual care without Al assistance	non-Al intervention; Healthcare intervention not compared to Al-assisted interventions
Outcome	Health economic evaluation methods or outcomes including cost effectiveness, cost utility, cost minimisation, cost benefit, cost consequences, costs, budget impact	Efficacy and/or safety only; No health economic evaluation outcomes
Study design	Systematic review of economic evaluations (model-based or trial-based studies, including cost effectiveness analysis (CEA), cost utility analysis (CUA), cost consequence analysis (CCA), cost minimisation analysis (CMA), budget impact analysis (BIA), health economic models)	Non-systematic reviews; Systematic reviews of clinical studies, abstract, posters, commentaries, letters, editorials and studies with no findings (e.g. protocol, methodology), animal studies, preprints
Study setting	No restrictions on country and healthcare settings	NA
Date of publication	2014 to 20 September 2024	Before 2014
Language of publication	English	Non-English

- > Majority of the EEs were conducted in North America or Europe.
- > Types of the EEs are shown in Figure 2. The category 'Other' (n=3, 6%) included profit analysis, regression analysis and financial performance analysis, which are not cost-effectiveness models.
- > Perspectives taken by the EEs are shown in Figure 3. Decision trees (DT), Markov models or combinations were commonly used (Figure 4) with time horizons ranging from 28 days to lifetime.

## > Key findings and limitations of the included SLRs are outlined in Table 3 and Figure 5.



## Table 3. Key findings and limitations of the included SLRs

## Key findings for Al-interventionsLimitations of existing EEs

# Economic Potential of AI for Cost ReductionLow Metherand Efficiency in Screening: AI hasit challengisubstantial potential to lower costs andInadequatimprove accessibility, particularly forhypotheticscreening in ophthalmology and diabeticpractice arretinopathy in low-resource settings. This isLack of Cordue to its ability to reduce the need fortransparentspecialist involvement and enable earlysensitivitydetection.4,5,9makers to

ction
 Low Methodological Quality: Many studies did not follow established best practices, making it challenging to compare results across different evaluations.<sup>6,8</sup>
 Inadequate Real-World and Long-Term Data: All the included EEs relied on short-term or hypothetical models, which might not reflect actual clinical and economic outcomes in practice and failed to accurately assess Al's impact over time.<sup>5,7</sup>
 his is
 Lack of Comprehensive and Transparent Reporting: The SLRs noted that EEs lack transparency and detail, with inadequate reporting on model structure, assumptions, and sensitivity analyses. This can reduce the studies' credibility and make it difficult for decision-makers to use the findings confidently.<sup>4,6</sup>
 Gap Between Al Development and EE: Rapid advancements in Al technology are not matched by the pace of EEs, which can hinder timely policy decisions. A faster, more agile approach to evaluation is recommended to keep up with Al's dynamic development.<sup>7,8</sup>
 cerns, The need for standardization in EEs of Al-interventions with consistent methods and reporting standards are highlighted in several SLRs to allow for more reliable comparisons across studies, ultimately supporting more informed decision-making.<sup>6,7,8</sup>

- > Two independent reviewers conducted the screening, with any discrepancies resolved through discussion or by involving a third reviewer.
- > Data extraction was carried out by one reviewer and verified by a second reviewer.
- > Outcomes of interest included details of the interventions, disease area, EE methods, findings and quality assessment of the EEs included in the SLRs.
- > The ISPOR CiCERO Checklist<sup>3</sup> was used to assess the quality of the included SLRs, and the EEs included in the SLRs.
- > Due to heterogeneity in the SLRs and the original primary studies, narrative synthesis was performed.

## Results

> 781 titles were identified, four removed for duplication, eight publications were screened at full text, with six SLRs being taken forward to data extraction (Figure 1).

## Figure 1. PRISMA diagram

Database search

Implementation Challenges: Despite Al's promise, practical challenges exist, including regulatory barriers, high initial implementation costs, data privacy concerns, and the need for context-specific adaptations, particularly in LMICs.<sup>4,5,8</sup>

## Figure 5. Application of the CiCERO checklist on the identified publications



Q1. Was the review conducted as per the predefined protocol?, Q2. Did the review clearly define the population, outcomes, time horizon, perspective, study design, and interventions/comparators?, Q3. Was a detailed search strategy provided, including the search date?, Q4. Was the search comprehensive and adequate?, Q5. Were the search dates provided, with justification if applicable?, Q6. Were the inclusion criteria appropriate and relevant?, Q7. Was the study selection process appropriate?, Q8. Was the methodological quality of the included studies assessed?, Q9. Was the risk of bias in the included studies considered in the review?, Q10. Were appropriate methods used to combine study results?, Q11. Were the included studies described in adequate detail?, Q12. Was any observed heterogeneity in the results explored and discussed?, Q13. Were biases, conflicts of interest, and reviewer funding discussed?

## Conclusion

- > As AI is increasingly used in healthcare, EEs of AI-assisted interventions and SLRs of these EEs are also increasing, but both are still limited in numbers, coverage, and quality.
- > Although the existing SLRs are not without limitations, they recognize that AI has substantial potential to lower costs and improve accessibility, particularly for screening in ophthalmology and



## diabetic retinopathy in low-resource settings.

- > Common gaps between existing EEs and CHEERS-AI reporting guideline<sup>1</sup> and NICE's ESF<sup>2</sup> included a lack of reporting transparency, inadequate real-world and long-term data, and adherence to standardized evaluation practices.
- > CHEERS-AI reporting guideline<sup>1</sup> and NICE's ESF<sup>2</sup> both directly address weaknesses identified in current AI EEs, offering comprehensive, context-specific, and standardized guidelines that could substantially improve future evaluations' quality, transparency, and relevance in healthcare settings. They should be closely followed to improve the quality of the future EEs and ensure consistent and standardised methodologies, with transparent and reproducible reporting.

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