



What is a Living Meta-Analysis?

A living meta-analysis is a systematic review and meta-analysis continually updated as new evidence becomes available, rather than being a one-time publication.

The ongoing "Living" network meta-analysis

periodically evaluates newly published data on Varithena* compared to endovenous thermal ablation in treating chronic venous insufficiency. provides more precise and generalizable evidence on the relative effectiveness and safety of Varithena and thermal treatments.

SR / MA with large sample sizes are considered Level 1A evidence – informing guidelines, payer policies and more.

*Varithena, the only FDA approved commercially available non-compounded polidocanol 1% endovenous microfoam ablation, from other foam scientificany options



Regular systematic searches of databases

Screening of new studies against inclusion

Data extraction from new eligible studies

Statistical updates incorporating new data

Regular quality assessment Version control of all updates

(eg, quarterly)

criteria

Living Meta Analyses Have Increased Utility

TIME LEDGE ATES	Traditional meta-analyses become outdated quickly in rapidly evolving fields Particularly valuable in fast-moving areas like emerging treatments
TICAL TAGES	Reduces research waste by avoiding multiple overlapping meta-analyses Maintains an up-to-date evidence base for clinical decision- making
IICAL LERS	Modern technology and databases make continuous updates more feasible Online platforms allow for easier collaboration and version control.
lity Ement	Allows for continuous refinement of methods Enables cumulative learning as evidence builds

Living Meta-Analysis A continuous Process

A Living Network Meta-Analysis

The Comparative Effectiveness of Non-compounded Polidocanol 1% Endovenous Microfoam (Varithena) Ablation versus Endovenous Thermal Ablation

"Overview

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The orgoing network meta-analysis (NMA) evaluates the evidence for Varithena (PEM) evaluates the evidence for Varithena (PEM) (CTI) in ret length of the evidence of the statistic field (CVI). By distinguishing Varithena, the only FDA-approved, commercially available noncompounded polidoconal 1% endowenous microfeam ablation, from other for an sclerotherapy options, this NMA provides more precise and generalizable evidence on the relative effectiveness and safety of these treatments."



48 publications Zero papers with Varithena cohort No change to LNMA

Original NMA publication date: April 26, 2024* Most Recent Literature Review : October 31, 2024



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