Systematic Review

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Performing a rigorous systematic review is multi-step process, which includes (a) identifying a well-defined focused clinically relevant question, (b) developing a detailed review protocol with strict inclusion and exclusion criteria, (c) systematic literature search of multiple databases and unpublished data, in consultation with a medical librarian, (d) meticulous study identification and (e) systematic data abstraction, by at least two sets of investigators independently, (f) risk of bias assessment, and (g) thoughtful quantitative synthesis through meta-analysis where relevant. Besides informing guidelines, credible systematic reviews and quality of evidence assessment can help identify key knowledge gaps for future studies.

Our systematic review is more structured as at every stage of writing we ensure to critically check the rigour using standard methodological checklists such as CASP, AMSTAR, and ARIF etc, based on the checklist provided. Further the general structure is presented as follows:

- Formulate the research question
- Search for studies
- Selection of studies
- Data collection
- Methodology quality assessment
- Results are analysed and presented
- Findings are interpreted.

Pubrica has done plethora of work in the area of clinical trial audits and monitoring for top pharmaceutical companies. Our CRAs will ensure a thorough review of data, frequent the sites, and perform interim

analysis. All tasks in compliance to ethics committee and regulatory standards such as Schedule Y, study protocol, ICH GCP and the other regulations.

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