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**Type of Article**: Original Research, Review Article, Case Report, Short Commentary [These are paper types. Please choose one and delete the rest.]

**Title:**

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**Introduction:**

**Methods:**

**Results:**

**Conclusion:**

**Keywords**: [Choose three to six keywords. They should not repeat words given in the title.]

**Clinical Trial Registration:** [For clinical trial and meta-analysis only]

**Introduction**

[Please note that citations are indicated by superscript numerals following the punctuation. All abbreviations are defined in full at their first occurrence.]

**Methods**

[A structured approach is recommended, which might include the following components:]

**Overview**

**Inclusion and Exclusion Criteria**

**Procedures, Definitions, Interventions, etc., as per the study design**

**Outcomes**

**Statistical Methods]**

**Results**

[Ensure that all tables and figures are appropriately cited]

**Discussion**

[1 The first paragraph usually summarizes the findings, highlighting how they align or contrast with previous studies and emphasizing any novel discoveries.

2. Include a section on limitations.

3 Avoid presenting results in this section.]

**Conclusion**

**Acknowledgments**

[Its purpose is to thank all of the people who helped with the research but did not qualify for authorship]

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For example: O.O. contributed to the study search, quality check, data extraction, and drafting. As principal investigators, O.O. and O.O. worked on the study search, quality check, data extraction, and analysis. O.O., O.O., and O.O. worked on the interpretation of data and the revision process. All authors have read the manuscript and agree with the content and data.

**Data availability:**

Please add “The datasets generated during and/or analysed during the current study are available in the [NAME] repository, [PERSISTENT LINK TO DATASETS]” or “The data supporting this study’s findings are available from the corresponding author upon reasonable request.”

**Ethical Statement:**

Please add “Informed consent was obtained from all subjects involved in the study.” OR “IRB approved this study of name and ID” or “Patient consent was waived due to REASON (please provide a detailed justification).” OR “Not applicable.”

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**References**

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Table 1 [Table titles do not end with a full stop.]

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[Notes: Abbreviations: AUC, area under the curve; LS, least squares; NE, not estimable. These are examples of the format.]

Table 2 [Table titles do not end with a full stop.]

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[Notes: Abbreviations: AUC, area under the curve; LS, least squares; NE, not estimable. These are examples of format.]

**Figure and legend**

Figure 1 [Title of figure and ends in a full stop. Please see Figures and tables for our Figure requirements.]

Figure 2 [Title of figure and ends in a full stop. Please see Figures and tables for our Figure requirements.]