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REVIEWER'S REPORT

Manuscript No.: IJAR-55012

Title: "TO COMPARE THE EFFICACY OF ORAL ACETAZOLAMIDE VS EPLERENONE IN PATIENTS OF CENTRAL SEROUS CHORIORETINOPATHY"

| Recommendation: | Rating | Excel. | Good | Fair | Poor |
|-------------------------------|----------------|----------|------|------|----------|
| Accept as it is | Originality | | | ✓ | |
| ✓ Accept after minor revision | Techn. Quality | | | | √ |
| Do not accept (Reasons below) | Clarity | | | | ✓ |
| | Significance | √ | | | |

Reviewer Name: Dr S. K. Nath

Date: 29.11.25

Detailed Reviewer's Report

Strengths of the Paper

- Clear Comparative Design: The study employs a randomized clinical trial design that effectively compares the efficacy of oral eplerenone and acetazolamide in treating CSCR, providing valuable clinical insights.
- **Defined Objectives and Methodology**: The study's aims are explicitly stated, and the methodology includes appropriate follow-up periods at 2 weeks, 6 weeks, and 3 months, which allow for a thorough assessment of treatment outcomes.
- Quantitative Data Presentation: The paper provides detailed tables with quantitative measurements of visual acuity, leakage on fundus fluorescence angiography, recurrence rates, and complications, facilitating objective analysis.
- **Relevant and Up-to-Date References**: The references cited are appropriate and include recent studies, which supports the credibility of the research.

Weaknesses of the Paper

- **Insufficient Ethical Details**: The paper does not mention whether ethical clearance was obtained or if patient consent was secured, which is a critical requirement for clinical studies.
- **Limited Sample Size**: The total sample size of 50 subjects may limit the generalizability of the findings and the statistical power of the study.
- **Short Follow-Up Duration**: A follow-up period of 3 months may be insufficient to evaluate long-term efficacy and recurrence, especially given the chronic nature of CSCR.
- Lack of Blinding or Placebo Control: The study design does not mention whether blinding or placebo control was used, which could introduce bias.
- **Absence of Detailed Adverse Effects Data**: There is minimal discussion on side effects or adverse reactions associated with each medication, which is important for clinical decision-making.
- **Formatting and Presentation Issues**: Some tables and data points could be better organized for clarity, and minor typographical mistakes are present.
- **Limited Discussion on Pathophysiology**: While the study mentions the complex pathophysiology of CSCR, it does not thoroughly explore how the medications influence the underlying mechanisms.
- Inadequate Explanation of Statistical Methods: The statistical analysis used for data comparison is not detailed, making it difficult to assess the robustness of results.

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Reviewer Comments

- **Ethical Clearance**: The paper should explicitly state whether ethical approval was obtained from an institutional review board or ethics committee. This information appears to be missing.
- **Methodology Issues**: The methodology is generally appropriate but lacks details about randomization procedures, blinding, and statistical analysis, which are essential for evaluating validity.
- **Typographical Mistakes**: Minor typographical issues are scattered throughout, such as inconsistent spacing and abbreviations.
- **Grammar and Language**: The English language usage is generally acceptable, but some sentences could benefit from clearer phrasing and grammatical refinement.
- **Formatting**: Consistency in table formatting, headings, and referencing style needs improvement to enhance readability.
- Clarity of Objectives, Results, and Conclusions: The objectives are clearly presented. The results are well-organized into tables, but a narrative synthesis would improve comprehension. The conclusion accurately reflects the findings.
- **References**: The references are relevant but limited; including more recent and diverse sources would strengthen the background and discussion.
- Additional Information Needed: Details on ethical approval, inclusion/exclusion criteria, adverse effects, and statistical analysis methods should be included to improve transparency and reproducibility.