Clinical Trial Protocol Peer Reviewer Form

Reviewer Information

Name

Email

Affiliation

Date of Review

Protocol Details

Protocol Title

Protocol Number/ID

Principal Investigator

Sponsor

Review

1. Scientific Rationale and Background

2. Study Design (Objectives, Endpoints, Methodology)

3. Ethical Considerations

4. Inclusion/Exclusion Criteria

5. Statistical Analysis Plan

6. Data Management and Monitoring

7. Safety Assessment and Adverse Event Reporting

8. Informed Consent Process

9. Overall Comments and Recommendations

Overall Assessment

Recommendation

Final Comments