Standard Operating Procedure: Clinical Trial Management (SOP-CTM-001)

1. Purpose

To define the responsibilities of the trial team.

2. Responsibilities

The Investigator must be qualified by education. (Note: Missing 'training and experience')

The Investigator shall maintain a delegation log.

3. Safety Reporting

Any SAE must be reported within 24 hours.

(Note: Missing the requirement for 'detailed written follow-up reports')

4. Compliance

The trial must be conducted according to the protocol.

(Note: Missing the prohibition on deviations without approval)