

Main responsibilities of Clinical Trial Participant:

Provide Informed Consent: Read and understand the informed consent document. Ask questions and clarify doubts before agreeing to participate. Voluntarily agree to participate without any coercion

Follow Study Protocol: Follow the study instructions carefully, such as taking medications as prescribed. Attend all scheduled appointments, tests and procedures on time.

Report Health Changes: Provide accurate and honest information about changes in health and any side effects, symptoms experienced.

Maintain Communication: Stay in regular contact with the site team. Inform the team if plan to miss an appointment or decide to leave the study early.

Confidentiality: A trial participant is responsible for keeping their personal and health information private during the trial. Participants should also understand how their information will be used and feel confident that confidentiality measures are in place to protect them throughout the trial

Thus, a trial participant helps advance medical research by providing valuable data, testing new treatments and supporting the development of better healthcare for the future