Clinical Trial Progress Monitoring Report Form

Trial Title

Protocol Number

Sponsor

Site Name/Location

Principal Investigator

Date of Visit

Type of Visit

Summary of Activities Performed

Subject Enrollment Status

Investigational Product Accountability

Informed Consent Documentation

Protocol Deviations/Violations

Adverse Events/Safety Issues

Data Quality (CRFs, Source Documentation)

Action Items/Recommendations

Monitor Name

Monitor Signature

Date