

Position: Site Report Specialist

Department	Clinical Trials
Basic qualification required	B.Pharm / M. Pharm, M.Sc
Brief JDs	<p>Job Description :</p> <ul style="list-style-type: none"> • Site Report Specialist (SRS)-CRA is defined as a team member from the SRS team. • SRS is formally assigned to a study as a Key Team Member with the primary responsibility of Site Visit Reports review. • The SRS reviews Site Visit Reports to ensure consistent standards of quality implementation. • The SRS focuses on subject safety, data integrity, escalations of issues, timely review and approval of the reports. • SRS Works in compliance with Cliantha SOP, ICH-GCP guidelines, protocol requirements and regulatory compliance.