Clinical Research Associate – I (Central Monitoring)

Department	Clinical Trial
Designation	Clinical Research Associate - I
Basic qualification required	B.Pharm / M.Pharm / M.Sc
Experience	1 - 2 Years
Location	Ahmedabad / Work from Home
Brief JD	1. Contribute to the development and use of Central Monitoring Plans and/or RBM specific tools and templates and/or other study specific plans. 2. Support project management team to develop monitoring strategy including monitoring triggers/thresholds. 3. With guidance, provide Inputs to clinical study teams, key decision makers, and internal team members to manage continuous process improvements, issue escalation, workload projections. 4. Perform centralized monitoring activities on assigned projects and evaluate the quality and integrity of the study as per the protocol, SOPs & respective regulation and guidelines. 5. Preparation of Site Performance Matrix (SPM) Tool for respective sites for assigned study (ies). 6. Escalate quality issues pertaining to site and/or subject to respective stakeholder within the project team. 7. Perform Subject Level Data Review that require further investigation with the clinical site to determine overall accuracy (inclusion & exclusion criteria/ IP/AE/ Labs/EOT/EOS/ End points/SAEs etc.) 8. Conduct periodic review of KRIs, Trend Analysis points and share the output with Project Team. 9. Monitor site performance and make recommendations for timely corrective actions (e.g. Site Telephone Contact or Triggered Onsite Monitoring Visit) 10. Perform Source Data Verification (SDV) as per defined scope for the study.

Interested candidates can send their resume at **rchourey@cliantha.com**