## Clinical Research Associate – I (Report Reviewer)

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| 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. Oversee Site Visit Reports (SRVs) for assigned studies, ensuring compliance to the protocol, processes, timelines, applicable SOPs, and GCP guidelines.<br>2. Review SVRs to ensure findings requiring corrective and /or preventative action plans are documented and followed up to resolution, to ensure high quality reports.<br>3. Participate on the project team for all SVR review activities and identifies and escalate CRA and /or site issues, relevant trends, and related risk factors to the project team and appropriate parties in a timely manner to optimize quality of project delivery.<br>4. Provide guidance to Clinical Project Managers (CPMs) at project start-up and throughout the study and partner with the project team members to decrease the level of corrections/additions needed on reports by providing insight and training of the SVR annotations.<br>5. Participate in meetings with project team to discuss any SVR review issues illustrative of quality/performance deficiencies across PIs/sites and CRAs. Assist with identification of the trends emerging.<br>6. To observe and escalate safety trends in patients as identified in SVRs. |

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