Clinical Research Coordinator Team

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The Clinical Research Coordinator Team (CRCT) carries through the compliant set-up of the clinical trial and oversees patient and study visit management, scheduling, and assists with general conduct of a clinical trial with PI oversight.

Clinical Research Coordinator Services

- Trained and credentialed clinical research coordinators that assist with patient recruitment, patient and study visit management, scheduling, orders development, facilitating monitoring visits, communication with campus and sponsor contacts and ensuring protocol and regulatory compliance during the conduct and closeout of a project.

Data Management Services

- CRF and eCRF data transcription from medical record and source documents
- Query resolution
- Source Document Verification – a review of the source document templates against the protocol and CRF to assist with protocol-compliant data collection during study visits

Budget and Contract Management

- Contract Office (CTC-SR, TDG, or OCGA) document submission for amendments
- OnCore patient management to facilitate sponsor invoicing
- Manage budget amendments

Contact us to meet with one of our team members to discuss your project needs.

Last updated: 26 Aug 2022

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