JULIAN MERCER

CLINICAL RESEARCH COORDINATOR

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PROFESSIONAL SUMMARY

Clinical Research Coordinator with 6+ years managing Phase II and III trials at academic medical centers. Skilled in patient recruitment, informed consent, protocol compliance, and sponsor communication, with a record of strong enrollment and clean monitoring visits.

PROFESSIONAL EXPERIENCE

Senior Clinical Research Coordinator May 2023 - Present
MD Anderson Cancer Center, Houston, TX

- Coordinated six concurrent oncology trials, managing visit schedules, source documentation, and data entry for 85 active participants.
- Screened electronic records and clinic lists to identify eligible patients, enrolling 34 participants across two studies in 14 months.
- Led informed consent discussions and maintained regulatory binders, achieving zero major findings across four sponsor monitoring visits.
- Resolved 95% of data queries within five business days by working directly with investigators and data managers.
- Trained three new coordinators on protocol amendments, adverse-event reporting, and IRB submission steps.
- Screened and enrolled participants for two oncology trials, meeting the site's monthly enrollment targets for six straight months.
- Prepared regulatory binders and IRB amendments with the principal investigator, supporting a clean sponsor monitoring visit.

Clinical Research Coordinator Aug 2020 - Apr 2023
Johns Hopkins Hospital, Baltimore, MD

- Managed recruitment and follow-up for a cardiology trial, retaining 91% of 120 participants through study completion.
- Prepared IRB submissions, continuing reviews, and amendment packages that were approved without revision requests.
- Reported adverse events and protocol deviations within required timelines using the sponsor's electronic data capture system.
- Built a visit-reminder workflow that lowered missed study visits from 12% to 5%.
- Maintained regulatory binders and delegation logs for three cardiology studies, passing two sponsor monitoring visits without major findings.
- Screened clinic schedules and referrals to enroll 41 participants across two studies, meeting site enrollment targets in five of six quarters.
- Scheduled research visits and reminded participants by phone and text, supporting steady retention through long follow-up periods.

CORE SKILLS

Clinical trials | Patient recruitment | Informed consent | Protocol compliance | ICH-GCP | IRB submissions | Adverse event reporting | Source documentation | REDCap | Medidata Rave | Regulatory binders | Data query resolution | Sponsor communication | Participant retention

EDUCATION

B.S. in Health Sciences, University of Pennsylvania, Philadelphia, PA

CERTIFICATIONS

ACRP Certified Clinical Research Coordinator (CCRC) | CITI GCP Certification