Marguerite Delacroix

Clinical Research Coordinator

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SUMMARY

Clinical research coordinator with 4 years on Phase II and III cardiology and oncology trials at an academic medical center. Manages a caseload of 35 active subjects across 6 protocols, maintains a zero-major-finding audit record over 3 FDA inspections, and coordinates with sponsors, CRAs and the IRB to keep studies on schedule.

EXPERIENCE

Clinical Research Coordinator II — Magnolia Heart Institute

Birmingham, AL · Mar 2023 – Present

- Coordinate 6 active Phase II/III cardiology trials with a combined caseload of 35 subjects, managing visit scheduling, consent and data entry.
- Screened 220 potential subjects over 18 months, enrolling 60 and keeping screen failures documented to sponsor standards with zero query escalations.
- Maintained regulatory binders and submissions for 6 studies, passing 3 sponsor and 1 FDA audit with zero major findings.
- Reduced average case report form entry time from 48 hours to 12 hours per visit by building standardized source document templates.

Clinical Research Coordinator I — Vulcan Oncology Research Group

Birmingham, AL · Jan 2021 – Feb 2023

- Supported 4 Phase I/II oncology trials, obtaining informed consent from 90 subjects and tracking adverse events per protocol.
- Coordinated with 8 CRAs across 3 sponsors to resolve 150 data queries within a 5-business-day average turnaround.
- Shipped and tracked 400+ biological specimens to central labs with zero chain-of-custody discrepancies over 2 years.

SKILLS

Regulatory: GCP, IRB submissions, informed consent, FDA audit preparation, protocol deviation reporting

Systems: REDCap, Medidata Rave, CTMS, EPIC research module, eConsent platforms

Clinical: Vital sign collection, phlebotomy, specimen processing, adverse event tracking

Coordination: Sponsor and CRA communication, subject recruitment, source document creation

CERTIFICATIONS

ACRP Certified Clinical Research Coordinator (CCRC), 2022

EDUCATION

B.S. Biology, University of Alabama at Birmingham, 2020

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