Clinical Research Coordinator | Oncology Trials

FIRSTNAME LASTNAME

Frankfurt am Main, Germany | firstname.lastname@email.com
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Profile

Clinical Research Coordinator with five years of experience at a university hospital trial centre, coordinating phase I to III oncology trials. Registered nurse by training. Handles patient screening, consent support, visits, sample logistics and EDC data entry to ICH-GCP. Main site contact for monitors and sponsors.

Career highlights

- Coordinated 11 trials at the same time with 60 active patients.
- Achieved the highest recruitment in Germany for two sponsor studies.
- Kept query resolution under five working days on average.
- Prepared the site for a sponsor audit that ended with no major findings.

Professional experience

Clinical Research Coordinator **April 2021 to present**
University hospital, clinical trials centre (example) Frankfurt am Main, Germany

- Screen patients, schedule visits and coordinate samples, imaging and pharmacy.
- Enter data in Medidata Rave and other EDC systems and resolve queries.
- Maintain the investigator site file and prepare monitoring visits.

Registered Nurse, Oncology Ward **October 2017 to March 2021**
Hospital (example) Offenbach am Main, Germany

- Cared for patients receiving chemotherapy and immunotherapy.
- Supported study patients on the ward and documented adverse events.
- Mentored nursing students.

Education

Registered Nurse, state examination, 2017 | Certified Clinical Research Coordinator course (study nurse qualification), 2021

Further training

ICH-GCP E6(R3) refresher (2025) | IATA shipping of biological samples (2024)

Languages

German (native) | English (good)

Core competencies

ICH-GCP | Oncology trials | Patient screening | Informed consent support | EDC (Medidata Rave) | Query management | Investigator site file | Sample logistics | IATA shipping | Sponsor audits

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