Tanyia Verser

Clinical Project Manager





• 123 Main St, Kansas City, MO 64108

Education

Master of Science in Clinical Research Management at Washington University in St. Louis, MO

Sep 2014 - May 2018

Relevant Coursework: Clinical Trial Design and Conduct, Biostatistics, Regulatory Affairs, Ethics in Clinical Research, Data Management and Analysis, Medical Writing, Pharmacovigilance, Quality Assurance, and Project Management.

Links

linkedin.com/in/tanyiaverser

Skills

Budgeting

Gantt-charting

Risk-assessment

Stakeholder-engagement

Protocol-development

Data-management

Regulatory-compliance

Languages

English

Mandarin

Profile

A dedicated Clinical Project Manager with 5 years of experience driving clinical trial success through meticulous planning, resource management, and cross-functional collaboration. Proficient in streamlining processes, ensuring regulatory compliance, and fostering strong relationships with key stakeholders. Demonstrates a consistent track record of delivering projects on time and within budget, while maintaining a keen focus on quality and patient safety.

Employment History

Clinical Project Manager at PRA Health Sciences, MO

Mar 2023 - Present

- Successfully led a team of 15 clinical research associates and coordinated with over 30 clinical sites to complete a Phase III clinical trial, resulting in FDA approval of a novel oncology drug.
- Reduced study start-up timelines by 25% through the implementation of streamlined processes and efficient communication strategies, enabling timely initiation of 10 clinical trials within a one-year period.
- Oversaw the management of a \$5 million clinical trial budget, ensuring financial targets were met and resources were effectively allocated across various project phases.
- Implemented quality control measures that led to a 20% reduction in data discrepancies and improved overall data integrity, contributing to successful regulatory submissions and approvals.

Associate Clinical Project Manager at, MO

Sep 2018 - Feb 2023

- Successfully managed a clinical trial with 300+ patients, completing the project 2 months ahead of schedule and saving the company \$200,000 in operational costs.
- Achieved a 95% patient retention rate across all clinical projects, resulting in high-quality data and faster study completion times.
- Implemented a streamlined communication process between the clinical team and external vendors, reducing response times by 30% and improving overall project efficiency.
- Led a cross-functional team of 15 members in the successful completion of a Phase II clinical trial, resulting in a positive FDA review and paving the way for Phase III trials.

Certificates

Project Management Professional (PMP)

Mar 2022

Certified Clinical Research Associate (CCRA)

Oct 2020