



PROFESSIONAL SUMMARY

Seasoned Clinical Research Coordinator with over 14 years of experience leading complex trials from initiation through close-out. Proven track record in multi-site management, regulatory compliance, and patient-centric care.

EDUCATION

2005 - 2009

Bachelor of Science in Public Health

San Diego State University / CA

Certifications

- **Certified Clinical Research Coordinator-** (CCRC) - ACRP (Renewed in 2024)
- **Certified in Clinical Research Administration** - The Joint Commission (Renewed in 2023)

SKILLS

- Multi-site trial management **Expert**
- Budgeting and contract negotiation **Expert**
- GCP, FDA, and ICH guidelines **Expert**
- Patient retention strategies **Expert**

EXPERIENCE



2015 - Now

Senior Clinical Research Coordinator
UC San Diego Health / San Diego, CA

- Lead the coordination of multi-site Phase II-IV clinical trials, focusing on oncology and immunotherapy.
- Manage study start-up activities, including regulatory submissions-, contract negotiations, and budget tracking.
- Oversee patient recruitment and follow-up, ensuring retention and high data quality.
- Train and supervise junior coordinators, providing guidance on GCP compliance and regulatory requirements.



2010 - 2015

Clinical Research Coordinator
Scripps Research Institute / San Diego, CA

- Coordinated clinical trials for rare diseases and immunology studies.
- Monitored data quality and conducted regular source verification (SDV).
- Liaised with CROs, sponsors, and investigators to ensure adherence to study timelines and milestones.

PRESENTATIONS



- "Effective Strategies for Multi-Site Clinical Trial Coordination." Presented at the 2023 ACRP Global Conference, Las Vegas, NV.
- "Innovations in Patient Retention Techniques for Oncology Trials."- Presented at the 2022 Society for Clinical Research Professionals Annual Meeting, San Diego, CA.