CASE STUDY

On demand Clinical Research Coordinator Staffing

Client: Large Regional Hospital Research Institute

SITUATION

The staff at a regional hospital research institute contacted Pearl to assist them find sufficient Clinical Research Coordinator (CRC) support. Their oncology division had been trying to fill two CRC positions for several months and needed immediate research support in order to implement their existing protocols. They were in particular need of an individual with experience in managing multi-center pharmaceutical studies.

SOLUTION

Within three weeks, Pearl placed two of their staff with deep CRC experience on-site at the hospital for a two month engagement. Pearl staff was able to quickly come up to speed on multiple protocols, led all aspects of the study, and communicated effectively with the study sponsors.

RESULT

The client was so pleased with the services of one talent that they approached Pearl and asked for a one month extension of the project. Pearl was able to accommodate this request and has an ongoing strong partnership with this regional hospital research institute.

Pearl Pathways™

Pearl Pathways is a comprehensive life science product development services company. Every day we strive to provide our customers top quality service, unyielding ethics, and efficient services through our team of experts. Pearl Pathways supports biopharmaceutical, medical device, and diagnostic companies as well as life science service providers with clinical, regulatory, and quality compliance needs. Our full-service central IRB supports all aspects of human research.

Clinical: Pearl Pathways' expertise in medical practice, science, ethics, and clinical research supports all phases of clinical development. Our expert consultants help implement customized solutions that drive efficient clinical research programs.

Regulatory (drug): Our consultants draw on over 20 years of scientific, industry, and FDA experience to translate your data into strategic drug development pathways, providing guidance from molecule to market while navigating a complex regulatory environment.

Regulatory (device): Delivering superior end-to-end product development and commercialization strategies, our consultants provide regulatory guidance that adds decades of industry and FDA regulatory expertise to your team, helping you balance risk, cost, and time to market.

Quality: From risk-based quality system development to gap analysis and remediation, our partnership helps reduce operational burdens, manufacturing impacts, regulatory agency warnings, and costly recalls—allowing you to focus on ongoing product quality assurance.

IRB: Our accredited independent review board ensures quality research, unyielding ethics, and human subject protection with individualized Expedited and Full Board reviews.

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