CASE STUDY

IRB Software System QC Services

Client: Large Hospital Healthcare System

SITUATION

A large hospital healthcare system was transferring IRB files from a legacy software system to a new cloud-based system. The IRB leader approached Pearl Pathways to provide quality compliance services (QC) for the transfer.

SOLUTION

Pearl's staff ensured all source data and files were mapped in the old system. Further, once the files for the old system were transferred, Pearl's staff executed a QC check to confirmed data integrity.

RESULT

Pearl was able to identify files that had not been transferred properly, managed the needed resolution while meeting an aggressive timeline. Pearl was able to deliver the QC services on time before access to the legacy system was discontinued, therefore achieving the client's goals. This large hospital healthcare system continues to be a loyal client utilizing a breadth of Pearl Pathways' serves.

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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