Research Disease Site Specialist (RN), Orthopedics

Contact: NORTHSIDE HOSPITAL

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DetailsApplyPosition DescriptionThe Research Disease Site Specialist will be the subject matter expert for their disease site or intervention type. This role will be responsible for guiding research activities for the major disease sites in which the Northside Hospital is focused. The Research Disease Site Specialist will work with the lead research nurse/ coordinator and their supervisor/ manager to guide the conduct of research protocols and provide training and education to clinical research staff. The Disease Site Specialist will also be responsible for identifying and reviewing new projects for their disease areas and support study start up processes. Support study selection and start up for assigned disease site.Identify and review potential protocolsComplete feasibility questionnairesSupport study start up processCreate source documents, review OnCore calendars and review pharmacy ordersHelp secure vendor services, as neededWork with the regulatory team to track regulatory approvalsFacilitate site initiation visitsSupport development of the medicare coverage analysisWork closely with research nurses and coordinators to guide protocol executionEstablish in depth understanding of assigned protocolsWork with research nurses and coordinators to centrally track and screen potential patients for assigned protocolsReview patient eligibilitySupport research nurses and coordinators and procedure schedulingWork with research nurses and coordinators to maintain thorough knowledge of patients on study and visits scheduledServe as the touch point for dose modificationsTrend the AEs/ SAEs for assigned protocols and provide training as appropriateProvide training on ongoing protocol changes, and related trends/ issues identifiedReview amendments and update formsAttend and/or present at tumor boardsSupport study closureFollow up with Regulatory team to ensure closureEnsure OnCore closureSupport leadership in the development and implementation of new or updated processes, and helps maintain, and evaluate the SOPs for administering clinical researchRegularly report pertinent information on areas of improvement to research leadership.Position RequirementsRequired: Nursing degree (Bachelors or Masters), and a minimum of five (5) years experience in a clinical research setting OR Licensed Practical Nurse, Diploma or Associates Nursing degree and a minimum of eight (8) years experience in a clinical research setting. Certification as a Certified Clinical Research Coordinator (CCRC) or Certified Clinical Research Professional (CCRP) must be completed within 1 year of hire date. Experience working with Clinical Research Nurses (CRN), Clinical Research Coordinators (CRC) and investigators or systems that relate to clinical research.Knowledge of clinical trial management systems and/or technology solutions used to manage clinical research. Knowledge of clinical trial policies and regulations, IRB and Clinical Practice (GCP). Previous

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