Clinical Manager for an Exciting and Growing

Medical Device Company located in the Caesarea area

Job Description:

- Work on the frontline of communication with clinical sites
- Ensuring study timelines
- Conduct and report all types of onsite monitoring visits
- Build and maintain relationships with clinical sites and investigators
- Be involved in study startup (if applicable)
- Perform CRF review, source document verification and query resolution
- Be responsible for site communication and management
- Supervise study activities, timelines, and schedules on the country level
- Be a point of contact for in-house support services and vendors
- Be involved in quality control, such as compliance monitoring and reports review
- Support regulatory team in preparing documents for study submissions
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Qualifications

- Degree in Life Sciences or an equivalent combination of education, training & experience
- On-site monitoring experience
- Experience in all types of monitoring visits in Phase II and/or III
- Participation in clinical projects as a Lead/Senior Monitor
- Experience monitoring urology - advantage
- Full working proficiency in English
- Proficiency in MS Office applications
- Ability to plan, multitask and work in a dynamic team environment
- Communication, collaboration, and problem-solving skills
- Ability to travel (once a month)

Please send your CV to Katzav.hagit@gmail.com