

Director of Clinical Operations

Responsible for clinical operations for effective execution of company clinical trials

Seek, evaluate, and open sites in various geographies (travel included)

Oversee sites management

Communication, and oversight of clinical CROs/CRAs/study coordinators and other relevant clinical personnel

Support the training of internal staff, study investigators and other site personnel on the proper conduct of the trial

Preparation and maintenance of clinical study documents and clinical SOPs

Responsible for overseeing clinical trial monitoring at sites

Responsible for database management

Assist with all company's activities and regulatory submissions, as requested

QUALIFICATIONS AND ESSENTIAL REQUIREMENTS

Education: Life Science / Bio-Medical Engineering degree -advantage

At least 4 years of experience in clinical operations for early phase studies, preferable in cardiovascular indications and class III devices, potentially as CRA or clinical coordinator

Good knowledge of GCP and regulatory framework for clinical development

Excellent communication skills in English (both written and spoken)

Goal orientation and ability to comprehend and work within complex clinical trials

Professional knowledge of common software packages (Word, PowerPoint, Excel) as well as clinical development tools and databases International travelling required

International travelling required.

michal@insigthr.co.il