

Job Details

Job Title Research Associate - Uro-oncology Clinical Trials Group

Job ID 56050

Location University Hospital

Full/Part Time Full-Time

Regular/Temporary Temporary

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Posting Period

Open: January 21, 2017

Deadline: January 27, 2017

Non - Union

Different terms and conditions of employment may apply to externally funded positions.

Department Name

Lawson Health Research Institute - Uro-oncology Clinical Trials Group

The Clinical Research Associate is required to conduct industry-sponsored, academic and investigator-initiated clinical trials in GU Cancer and Urologic Diseases. This includes interventional drug trials, medical device trials, as well as quality of life and observational studies. The successful candidate will be responsible for research ethics board submissions, required institutional and departmental submissions, participant screening and enrolment, and overall study conduct including data collection. The position requires working collaboratively with a study team and reporting to the Unit Manager.

Rate of Pay: To commensurate with experience

Hours of Work: 37.5 hours per week

Duration: 12 months

Qualifications

- Minimum of Bachelor's Degree in health related field
- Minimum of two years clinical research experience and must demonstrate full responsibility for implementing drug and/or medical device trials (oncology drug trial experience an asset)
- Clinical Research Professional certification required (SOCRA, ACRP)
- Blood processing and phlebotomy certification required
- Transportation of Dangerous Goods/ International Air Transport (TDG/IATA) certification required
- Current CPR certification an asset
- Demonstrated knowledge of current regulations and guidelines for conducting clinical trials (ICH GCP, USA FDA 21 CFR, Tri-Council Policy Statement) and privacy legislation (PIPEDA, PHIPA)
- Knowledge of local REB requirements an asset
- Well-developed patient assessment and evaluation skills
- Demonstrated knowledge of drug therapy, management and accountability
- Demonstrated ability to plan, prioritize, execute and manage several research studies
- Demonstrated computer proficiency in Microsoft Office and experience with electronic data entry (eCRFs) and databases
- Excellent verbal and written communication skills in English. Demonstrated ability to communicate effectively and professionally with co-workers and study participants
- Demonstrated cooperation in a team environment
- Highly motivated and self-directed
- Demonstrated knowledge of and commitment to the principles of patient and family centred care
- Demonstrated knowledge of and commitment to patient and staff safety at LHSC
- Demonstrated ability to attend work on a regular basis

London Health Sciences Centre fosters a culture of patient and staff safety whereby all employees are guided by LHSC's Mission, Vision, Values and Code of Conduct.

We are committed to providing a safe, healthy and inclusive work environment that inspires respect. LHSC encourages applications from persons with disabilities and we are committed to providing accommodations upon request.

As part of the assessment process applicants may be required to complete a written examination or test. Please be advised that an internal reference check may be conducted as part of the selection process.

Your interest in this opportunity is appreciated. Only those applicants selected for an interview will be contacted. Successful candidates, as a condition of job offer, would be required to provide a satisfactory police check (original document) completed in the last 8 months.