

## Job Details

**Job Title** Research Associate

**Job ID** 55388

**Location** Victoria Hospital

**Full/Part Time** Full-Time

**Regular/Temporary** Temporary

**Favorite Job** ☆

### Posting Period

Open: September 27, 2016

Deadline: October 10, 2016

Non-Union

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

### Department Description

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 of Contract: 12 months

### Qualifications

- Successful completion of a Bachelor's Degree in a health related field
- Two (2) years of clinical research experience (oncology drug trial experience preferred)
- Clinical Research Professional Certification (SOCRA, ACRP)
- Blood processing and phlebotomy Certification
- Transportation of Dangerous Goods/ International Air Transport (TDG/ IATA) Certification
- Current CPR Certification (preferred)
- 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) (preferred)
- Demonstrated knowledge of local REB requirements (preferred)
- Well-developed patient assessment and evaluation skills
- Proficient in Microsoft Office and experience with electronic data entry and databases
- Demonstrated knowledge of drug therapy, management and accountability
- Demonstrated ability to plan, prioritize, execute and manage several research studies
- Excellent interpersonal and communication skills
- Ability to work effectively both independently and as part of a team
- Demonstrated flexibility with a high level of initiative and self-direction
- 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 a 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 vulnerable sector police check (original document) completed in the last 8 months.