



## Job Description

Department	Medical Affairs
Designation Location	Full Time Nurse Practitioner Mississauga, Ontario
Basic qualification required	<ol> <li>Current Nurse Practitioner license in the Province of Ontario</li> <li>Completion of a Narcotics and Controlled Substances Prescribing Course</li> <li>Certification in ACLS preferred</li> <li>Demonstrate knowledge of or participation in a course on ICH GCP Guidelines is an asset</li> <li>Strong and accurate diagnostic skills</li> <li>Be able to work in a fast-paced environment as well as maintain productivity during less demanding times.</li> <li>Must be able to work and learn independently and maintain strong analytical and problem-solving skills.</li> <li>Must be detail oriented and self-motivated; and have excellent work ethic, organizational skills, efficiency, and ability to multi-task and prioritize effectively</li> <li>Excellent interpersonal and leadership skills</li> <li>Strong written and verbal communication skills</li> <li>Due to the nature of the research industry, the candidate must be able to accommodate flexible hours and manage changes to work schedule.</li> </ol>
Experience	1-2 Years
Brief JD	<ol> <li>Reviewing the study protocol and informed consent form</li> <li>Performs medical procedures as outlined in the protocol (i.e. Physical exams. MMSE, Pap tests)</li> <li>Administer any form of Investigational Product as specified in the protocol</li> <li>Evaluate subject's eligibility for a clinical trial by reviewing the medical history, physical examination, laboratory values, ECG and diagnostic findings, spirometry/ PFT and the study inclusion and exclusion criteria</li> <li>Monitor and medically manage subject's health status and progress during the clinical trial according to the protocol, Cliantha Research Limited SOPs, and sound clinical judgment.</li> <li>Assist and/or direct the study coordinator, study manager, or other study personnel in the functions of a clinical trial</li> <li>Participate in any external or internal audit of a clinical trial</li> <li>Policy and Standards of Practice/ Standard Operating Procedure (SOP) creation and revision</li> <li>Training of clinical staff</li> <li>Be on-call after hours if required</li> </ol>



