Asia-Pacific Centre for Neuromodulation

Local Services Offered: Clinical trials site
Details: In addition to the information contained within csv generated by this website, this PDF provides additional site information

Life Sciences Queensland (LSQ) Member	No
Other areas of expertise	Movement Disorders Touretts,
Sub-Therapeutic areas	Parkinsons; Touretts; Neurological
Does your Clinical Trial Site have the capacity to conduct Clinical Trials involving GMOs?	No
Other Affiliated Research Sites	Yes
Other facility details	Royal Brisbane and Women's Hospital
Is your Facility affiliated with a government agency or part of a government funded health service?	Don't know
Are there any notable factors relating to your Patient Population	No
Does your Facility perform HREC (IRB/ERB/Ethics) Committee submissions?	Yes
Does your Facility have a dedicated department or group to perform HREC (IRB/ERB/ETHICS) Committee submissions?	No
Any Additional Process	No
Does your Clinical Trial site or Service undertake any recruitment?	Yes
Has your Clinical Trial site been audited?	Not Applicable
Has your Clinical Trial Site or Service been accredited?	Not Applicable
If your Clinical Trial Site or Service has been accredited, please select all relevant types	Yes
Other Meeting Frequency	N/A
Review required	Yes
Does the HREC Committee require payment prior to the release of final approval documents?	Yes

Does the HREC require contract/budget approval prior to release of final approval documents?	Yes
Does your Facility use private laboratory services?	Yes
Does your Facility have a written SOP/Policy/Procedure for Informed Consent?	Yes
Does your Facility have a written SOP/Policy/Procedure for Other vulnerable populations?	No
Does your Facility have a written SOP/Policy/Procedure for Minor Assent for paediatric populations?	No
Will your Facility require language translations for consents?	No
Does your Facility have a training program for the research staff?	No
Does the course content include GCP?	Yes
Do you have a process or program in place to retrain research staff when a protocol is amended?	Yes
Does the study staff that prepares or transports dangerous goods have training that meets the IATA International Air Transport Association (US) or other countries hazardous training requirements for shipping dangerous goods?	Yes
Can your Facility support patient visits on weekends?	Yes
Can your Facility support in-patient admissions for research studies?	Yes
Does your Facility have access to translators and translation support for study conduct (e.g. consent, study-specific instruction)?	Not Applicable
Does the Facility have storage space for Study-Related materials (e.g. Lab Kits, Patient Materials, etc.)?	Yes
Does your Facility have the ability to collect and store PK/PD specimens?	No
Does your Facility have the ability to collect PK/PD samples beyond normal business hours?	No

Does your Facility typically allow the collection of Pharmacogenomic (PGX) samples for research purposes?	No
Identify the Diagnostic Equipment available at or near the Facility to support Research studies?	MRI Magnetic Resonance Imaging
Describe any additional equipment relevant to Clinical Trials	N/A
Does your Facility have an SOP or process that ensures routine calibration and maintenance of general equipment? Examples of general equipment include: scale, pulse oximeter, stadiometer, sphygmomanometer, etc.?	No
Does your Facility have the necessary equipment to treat medical emergencies (for example crash/code cart)?	No
Operating Systems Supporting Studies	Windows (Windows XP, Windows 7, Windows 10, etc), Apple/Mac (OS X Snow Leopard, Mountain Lion, El Captain, etc)
What browser does your facility use?	Safari, Chrome
Does your Facility limit or prohibit access and use of external web-based tools or sites for clinical research (E.g. web portals to submit documents to sponsors or CROs)?	No
Does the Facility have access to local IT support?	Yes
Does your Facility have the ability to manage on-site or off-site destruction of controlled substances when appropriate?	Yes
Is your Facility capable of administering infusions?	No
Is your Facility adequately staffed to support studies with both blinded and unblinded Investigational Product?	Yes

Does your Facility have the ability to manage on-site or off- site destruction of controlled substances when appropriate?	Not Applicable
Does the Facility have the ability to handle radio-labelled Investigational Products?	No
Provide Location name and address of any offsite archives.	Yes
Do you have Electronic Health Records (EHR)/ Electronic Medical Records (EMR)?	No
What EMR/EHR system do you use? Electronic Medical Records (EMR) /Electronic Health Records	In-house
Please indicate all equipment that will be available to Monitors	Phone, Copy Machines, Internet Access