University of Queensland Centre for Clinical Research (UQCCR)

Location:
27.4487,153.0286
Local Services Offered:
Clinical trials site, Pre-clinical, Phase 1 unit
Details: In addition to the information contained within csv generated by this website, this PDF provides additional site information

LSQ Member	Yes
Sub-Therapeutic areas	Clinical Neurosciences: Language neuroscience; Translational neuroscience and drug repurposing, Queensland drug repurposing initiative; The genomic and epigenomic landscapes of epilepsy; The perinatal brain (the newborn brain injury and repair); Neuro-immunology; Neuro-mental health & dementia research; N of -1 trials and sleep research. Infectious diseases: Antibiotic resistance research; Antimicrobial optimisation; Microbial diagnostic and characterisation; Ear, Nose and Throat research; CRE in redefining antimicrobial use to reduce resistance; Burns trauma and critical care research centre. See attached document. Fertility: Human oocyte research; Epigenetics and oogenesis; Sirtuin-mediated regulation of female fertility; DNA damage and ovarian follicular reserve; Meiotic cell-cycle regulation and chromosome segregation Cancer: Molecular breast pathology; Queensland centre for gynaecological cancer; Exosome biology laboratory
Other Affiliated Research Sites	No
Is your Facility affiliated with a government agency or part of a government funded health service?	No
Are there any notable factors relating to your Patient Population	UQCCR Level 3 The Clinical Research Facility, is an outpatient research facility conducting Phase ll - lV clinical trial activity for healthy volunteers. and a range of health conditions. UQCCR supports a dynamic research culture, and is equipped with a range of facilities, interdisciplinary teams purpose-fit, with proximity to specialty services for clinical trial conduct. Co-location and accessibility to specialist services and expertise facilitates bespoke project delivery within a framework of governance and quality.

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?	Yes
Any Additional Process	No
Does your Clinical Trial site or Service undertake any recruitment?	No
Has your Clinical Trial site been audited?	Yes
If your Clinical Trial Site has been audited, please select all relevant types.	Sponsor
Has your Clinical Trial Site or Service been accredited?	No
HREC Committee Name	MNHHS-RBWH, TPCH, Children's Health Queensland HHS, UQ - UQR & I, Office of Research and Ethics
Does the HREC Committee require payment prior to the release of final approval documents?	No
Does the HREC require contract/budget approval prior to release of final approval documents?	No
Does your Facility have other review boards that need to approve the study prior to HREC (IRB/ERB/Ethics) Committee submission? For example, scientific, radiation safety committees, or others	Yes
If other review boards, please name	Scientific and Radiation Safety Committees. UQ IBC (Institutional Biosafey Committee). UQ Radiation
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

Does your Facility have access to translators and translation support for study conduct (e.g. consent, study-specific instruction)?	No
Training program for the research staff	Yes
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 your Facility have a training program for the research staff?	Yes
Can your Facility support patient visits on weekends?	Yes
Can your Facility support in-patient admissions for research studies?	No
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 PK/PD samples beyond normal business hours?	Yes
Does your Facility have the ability to collect and store PK/PD specimens?	Yes
Does your Facility typically allow the collection of Pharmacogenomic (PGX) samples for research purposes?	Yes
Additional equipment	EEG Suite, Patient Scales (Stand-up digital and chair), Digital Sphygmomanometer, hoist, small bench top
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.?	Yes
Does your Facility have the necessary equipment to treat medical emergencies (for example crash/code cart)?	Yes

Does your Facility have computers that are dedicated to research studies?	Yes
What type of computer operating system(s) does your institution use to support studies?	Windows (Windows XP, Windows 7, Windows 10, etc), Apple/Mac (OS X Snow Leopard, Mountain Lion, El Captain, etc), Unix/Linux (Solaris, Ubuntu, Redhat, etc)
Internet Upload Speed	910 MBPS
Internet Download Speed	780 MBPS
What browser does your facility use?	Internet Explorer, Safari, Firefox, Chrome, Other
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
IP Recipient Name	UQCCR
Storage Room Backup Power	Yes
Does your facility have a written SOP/Policy/Procedure for the destruction of Investigational Product?	No
Handling Investigational Product	Secure, restricted access Investigational Product Storage and destruction facilities.
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?	No
Does the Facility have the ability to handle radio-labelled Investigational Products?	No
Secure Storage Records	Yes
Provide Location name and address of a	Grace Storage: 420 Sherbrooke Road, Willawong Qld
Do you have Electronic Health Records (EHR)/ Electronic Medical Records (EMR)?	Yes
EMR/EHR systems	Other

Medical access limitations	ieMR access - restricted access can be provided to monitor(s) under supervision and with approval from
Please indicate all equipment that will be available to Monitors	Phone, Fax, Copy Machines, Internet Access
EDC Systems	Medidate Rave REDCap