Translational Research Institute

Location:
-27.49863116542877, 153.031237842328
Local Services Offered:
Pre-clinical device design, Early Phase, Complementary medicines clinical trials, Clinical trials site, Pre- clinical, GP trials, Phase 1 unit, Bench research, Investigator Initiated Trials, Satellite Site, Trial Patient Recruitment, Treatment of patients, Completion of study documentation as per ICH GCP and contract
Details: In addition to the information contained within csv generated by this website, this PDF provides additional site information

Life Sciences Queensland (LSQ) Member	Yes
Other areas of expertise	N/A
Sub-Therapeutic areas	N/A
Does your Clinical Trial Site have the capacity to conduct Clinical Trials involving GMOs?	Yes
If yes, which of the following? (chose all that apply)	GM products that do not contain live GMOs, DNA vaccines, Clinical trials involving other types of GMOs, Other
If you selected other in the previous question, please specify which other types of GMOs	Live attenuated bacteria and viral agents
Other Affiliated Research Sites	Yes
Other facility details	Brisbane and interstate
Is your Facility affiliated with a government agency or part of a government funded health service?	Yes
Are there any notable factors relating to your Patient Population (e	Our CRF facility is located within the Metro South Hospital and Health Service district which is the largest health service area within Queensland. Our TRIC facility is located on the main Children's Health Service campus which is the main referral centre for children across Queensland.
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	Yes
Details of other steps for HREC (IRB/ERB/Ethics)Committee review and submission	Parallel ethics and research governance submissions. HREC meets monthly.

Does your Clinical Trial site or Service undertake any recruitment?	Yes
What percentage of Clinical trials undertaken on your site do you meet or exceed the recruitment target?	>90%
Has your Clinical Trial site been audited?	No
Has your Clinical Trial Site or Service been accredited?	No
Other Meeting Frequency	N/A
Does the HREC require contract/budget approval prior to release of final approval documents?	No
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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	No
If other review boards, please name	N/A
Local Lab Usage	Yes
Lab Name	Pathology Queensland- Princess Alexandra Hospital
Does your Facility use private laboratory services?	Yes
If you selected 'Yes' on the previous question, please specify here which services	SNP, QML
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?	Yes
Does your Facility have a written SOP/Policy/Procedure for Minor Assent for paediatric populations?	Yes
Does your Facility have access to translators and translation support for study conduct (e.g. consent, study-specific instruction)?	Yes
Training program for the research staff	Yes
Does the course content include GCP?	Yes

Please provide program course/s name	The Global Health Network ICH Good Clinical Practice E6 (R2)
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?	No
Can your Facility support in-patient admissions for research studies?	No
Is your Facility adequately staffed to support studies with both blinded and unblinded Investigational Product?	Yes
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?	Yes
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?	Yes
Additional equipment	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.?	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)
Internet Upload Speed	Unknown
Internet Download Speed	Unknown
What browser does your facility use?	Internet Explorer, Safari, Firefox, 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
IP Storage Location Name	Princess Alexandra Hospital Pharmacy
Is the Investigational Product Storage Room secured with controlled access?	Yes
Do you provide your Satellite Site(s) with a dedicated inventory of Investigational Product?	Yes
Does your facility have a written SOP/Policy/Procedure for the destruction of Investigational Product?	Yes
Is your Facility capable of administering infusions?	Yes
Does your Facility have the ability to manage on-site or off-site destruction of controlled substances when appropriate?	Yes
Does the Facility have the ability to handle radio-labelled Investigational Products?	Yes
Location Information	Yes
Secure Storage Records	N/A
Do you have Electronic Health Records (EHR)/ Electronic Medical Records (EMR)?	Yes
EMR/EHR systems	Other
Location of Monitor	Brisbane or remote access available via QHealth MyApps
Medical access limitations	Annual access fee per monitor per study.
Please indicate all equipment that will be available to Monitors	Phone, Fax, Copy Machines, Internet Access
EDC Systems	Sponsor specific.
Please provide additional information not captured in other sections of the Facility Profile that you feel is important for Sponsors to know about your Facility	N/A