QIMR Berghofer Medical Research Institute

Location:
-27.4491356580931, 153.02725959838406
Local Services Offered:
Early Phase, Pre-clinical, Bench research, Investigator Initiated Trials
Details: In addition to the information contained within csv generated by this website, this PDF provides additional site information

Is your Facility affiliated with a government agency or part of a government funded health service?	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?	Yes
Any Additional Process	Yes
Details of other steps for HREC (IRB/ERB/Ethics)Committee review and submission	Details of other steps for HREC (IRB/ERB/Ethics) Committee review and submission: HREC review includes a staged review, first scientific review (clinical trial protocol committee or human scientific sub-committee) and then ethics review
Does your Clinical Trial site or Service undertake any recruitment?	No
Has your Clinical Trial site been audited?	Not Applicable
Has your Clinical Trial Site or Service been accredited?	Not Applicable
Other Meeting Frequency	9 times per year
Does the HREC require contract/budget approval prior to release of final approval documents?	No
Does the HREC require contract/budget approval prior to release of final approval documents?	Yes
Does your Facility have other review boards that need to approve the study prior to	Yes
If other review boards, please name	Human scientific subcommittee and Clinical trial protocol committee. Safety/Biosafety committee meets monthly
Local Lab Usage	No

Does your Facility use private laboratory services?	Yes
If you selected 'Yes' on the previous question, please specify here which services	Q-Gen Cell Therapeutics - Microbiological, Immunological
Does your Facility have a written SOP/Policy/Procedure for Informed Consent?	No
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)?	Yes
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 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?	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 typically allow the collection of Pharmacogenomic (PGX) samples for research purposes?	Yes
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)?	No
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	100 Mbps
Internet Download Speed	100 Mbps
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 Recipient Name	QIMR Berghofer Medical Research Institute
Storage Room Backup Power	Yes
Does your Facility have the ability to manage on-site or off-site destruction of the Investigational Product?	No
Do you provide your Satellite Site(s) with a dedicated inventory of Investigational Product?	Not Applicable
Does your facility have a written SOP/Policy/Procedure for the destruction of Investigational Product?	Not Applicable

Is your Facility capable of administering infusions?	No
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?	No
Do you have Electronic Health Records (EHR)/ Electronic Medical Records (EMR)?	No