HERSTON IMAGING RESEARCH FACILITY

Location: -27.446644542139715, 153.02901282726546
Local Services Offered:
Early Phase, Complementary medicines clinical trials, Clinical trials site, Pre-clinical, Investigator Initiated Trials
Details: In addition to the information contained within csv generated by this website, this PDF provides additional Site information

LSQ Member	No
Sub-Therapeutic areas	Post image processing Image analysis
Does your Clinical Trial Site have the capacity to conduct Clinical Trials involving GMOs?	No
If you selected other in the previous question, please specify which other types of GMOs	N/A
Other Affiliated Research Sites	Yes
Other facility details	Department of Nuclear Medicine, RBWH; Translational Research Institute, PAH; Centre of Advanced Imaging, St Lucia
Is your Facility affiliated with a government agency or part of a government funded health service?	Yes
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?	N/A
What percentage of Clinical trials undertaken on your site do you meet or exceed the recruitment target?	99%
Has your Clinical Trial site been audited?	Yes
If your Clinical Trial Site has been audited, please select all relevant types.	CRO (Clinical Research Organisation)
Has your Clinical Trial Site or Service been accredited?	Yes
If your Clinical Trial Site or Service has been accredited, please select all relevant types	Other

If you selected Other type of Accreditation, please list here	DIAS
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 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	Radiation Dose Risk Assessments are required for any project that requires radiation
Local Lab Usage	No
Does your Facility use private laboratory services?	No
Does your Facility have a written SOP/Policy/Procedure for Informed Consent?	Yes
Does your Facility have a written SOP/Policy/Procedure for Minor Assent for paediatric populations?	Yes
Does your Facility have a written SOP/Policy/Procedure for Other vulnerable 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
Include GCP	Yes
Program course name	It is project specific
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?	Yes
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.)?	No
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?	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), Unix/Linux (Solaris,
What browser does your facility use?	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 the Investigational Product?	No
Does your Facility have the ability to manage on-site or off-site destruction of controlled substances when appropriate?	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?	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?	Yes

Do you have Electronic Health Records (EHR)/ Electronic Medical Records (EMR)?	Yes
EMR/EHR systems	In-house system
Please indicate all equipment that will be available to Monitors	Internet Access