Herston Imaging Research Facility (HIRF)

Local Services Offered:
Early Phase, Complementary medicines clinical trials, Clinical trials site, Pre-clinical, Investigator Initiated Trials
Details: This PDF complements the information found in the CSV generated by this website with additional site information.

Facility Details:	
Please provide your Facility Website.	https://www.hirf.com.au/
Is your facility/organisation a Life Sciences Queensland (LSQ) Member?	No
Is your Facility affiliated with a government agency or part of a government funded health service?	Yes
Do you have Affiliated Research Sites or Satellite Sites/Clinics? A Satellite Site is a secondary location where the investigator sees clinical trial subjects. Usually this is the same investigator who sees subjects at the primary site location.	Yes
Please provide other facility details.	Department of Nuclear Medicine, RBWH; Translational Research Institute, PAH; Centre
Provide the list of Sub-Therapeutic Areas for your Facility.	Post image processing Image analysis
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 or Service been accredited?	Yes
If your Clinical Trial Site or Service has been accredited, please provide the type accreditation.	DIAS
IRB/ERB/Ethics Committee:	
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
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

Are there any other steps that the Sponsor should be aware of for your IRB/ERB/Ethics Committee review and submission?	Radiation Dose Risk Assessments are required for any project that requires radiation
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
Consent:	
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:	
Does your Facility have a training program for the research staff?	Yes
Does your Facility training course content include GCP?	Yes
If your facility uses external program course/s. Please provide the program course/s name.	It is project specific
Do you have a process or program in place to retrain research staff when a protocol is amended?	Yes
Facility And Equipment:	
Facility Capabilities:	
Can your Facility support in-patient admissions for research studies?	Yes
Can your Facility support patient visits on weekends?	No
Is your Facility capable of administering infusions?	Yes
Is your Facility adequately staffed to support studies with both blinded and unblinded Investigational Product?	Yes
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
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 the Facility have storage space for Study-Related materials (e.g. Lab Kits, Patient Materials, etc.)?	No
Equipment:	
Does your Facility have the necessary equipment to treat medical emergencies (for example crash/code cart)?	Yes
IT Capabilities:	
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 the Facility have access to local IT support?	Yes
Please indicate all equipment that will be available to Monitors	Internet Access
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
Labs:	
Does your Facility use Local Lab Services?	No
Does your Facility use private laboratory services?	No
IP Storage Details:	
Does your Facility have the ability to manage on-site or off-site destruction of the Investigational Product?	No
Does your facility have a written SOP/Policy/Procedure for the destruction of Investigational Product?	Not Applicable

Does the Facility have the ability to handle radio-labelled Investigational Products?	Yes
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
Source Documents:	
Electronic Medical Records (EMR) /Electronic Health Records (EHR):	
Do you have Electronic Health Records (EHR)/ Electronic Medical Records (EMR)?	Yes
What EMR/EHR system do you use?	In-house system