Gallipoli Medical Research Foundation

Location: -27.5122,153.0454 Local Services Offered:

Early Phase, Clinical trials site, Phase 1 unit, Bench research, 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

Contact Phone Number	0733947297
Life Sciences Queensland (LSQ) Member	No
Facility Department	Medicine
Other areas of expertise	Respiratory Non tuberculosis mycobacterium
Sub-Therapeutic areas	NASH, Solid Tumours
Other Affiliated Research Sites	Yes
Other facility details	Ramsay Clinical Trials Network
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	Choice of Ramsay Queensland HREC and lead to Bellberry
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?	Yes
Audit Type	FDA (Food Drug Administration)
If you selected Other types of Audit, please list here	EMEA
Has your Clinical Trial Site or Service been accredited?	No
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	Governance process to be determined
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?	No
Does your Facility have access to translators and translation support for study conduct (e.g. consent, study-specific instruction)?	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
Include GCP	Yes
Program course name	Praxis, ACRP
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.)?	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?	Yes
Does your Facility typically allow the collection of Pharmacogenomic (PGX) samples for research purposes?	Yes
Additional equipment	Fibroscan for NASH studies
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
Operating Systems Supporting Studies	Windows (Windows XP, Windows 7, Windows 10, etc)
What browser does your facility use?	Safari
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)?	Yes
Does the Facility have access to local IT support?	Yes
IP Recipient Name	Ramsay Oncology Pharmacy
IP Recipient Street Name and Number	Greenslopes Private
Additional Address Info	Cyril Gilbert Cancer Centre, GPH Newdegate
Is the Investigational Product Storage Room secured with controlled access?	Yes
Does your Facility have the ability to manage on-site or off-site destruction of the Investigational Product?	Yes
Does your Facility have the ability to manage on-site or off-site destruction of controlled substances when appropriate?	Yes
Do you provide your Satellite Site(s) with a dedicated inventory of Investigational Product?	Not Applicable
Is your Facility capable of administering infusions?	Yes
Staff Support Studies	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?	No
Secure Storage Records	Grace
Do you have Electronic Health Records (EHR)/ Electronic Medical Records (EMR)?	Yes
Please indicate all equipment that will be available to Monitors	Phone, Fax, Copy Machines, Internet Access
EDC Systems	Oracle Inform, Medidata Rave, Oracle Remote Data Capture (RDC), IBM merge, VIADOCS,