Townsville University Hospital – Cardiothoracic Research

Local Services Offered:

Clinical trials site, Trial Patient Recruitment, Treatment of patients, Completion of study documentation as per ICH GCP and contract

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.townsville.health.qld.gov.au/
What Department is your Trial Site?	Surgical Service Group- Cardiothoracic
(Queensland Health HHS- See List of	
Services and Levels-Clinical Services	
Capability Framework)	
Is your facility/organisation a Life	No
Sciences Queensland (LSQ) Member?	
Do you have Affiliated Research Sites	Yes
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.	
Please provide other facility details.	Other Townsville HHS facilities
Please provide other areas of expertise	Cardiothoracic Surgery
for your Facility.	Cardiothoracic case management
Provide the list of Sub-Therapeutic	Cardiac surgery for coronary artery bypass grafting,
Areas for your Facility.	Heat valve replacement and repair surgery,
	Adult open-heart surgery,
	Minimally invasive heat valve repair and
	replacement,
	Transcathether aortic valve implantation (TAVI), Extracorporeal membrane oxygenation (ECMO),
	Video-assisted surgery for lung cancers,
	Surgery for chest wall and chest trauma, excessive
	sweating (hyperhidrosis),
	Telehealth clinics and Outreach clinics for Cairns
	and Mackay,
	Cardiothoracic patient education and rehabilitation.
Does your Clinical Trial site undertake	Yes
any patient recruitment?	

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Yes
No
Townsville HHS HREC
One meeting per month in NMA
Yes
Yes
Yes
Yes
Yes
Other
Other
Yes
ARCS online
Sponsor – Led Training/Courses Yes
Yes

Facility And Equipment:	
Facility Capabilities:	
Can your Facility support in- patient	Yes
admissions for research studies?	
Can your Facility support patient visits on weekends?	Yes
Is your Facility capable of	Yes
administering infusions?	
Is your Facility adequately staffed to support studies with both blinded and unblinded Investigational Product?	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
Does the Facility have storage space for Study-Related materials (e.g. Lab Kits, Patient Materials, etc.)?	Yes
Equipment:	
Does your Facility have the necessary equipment to treat medical emergencies (for example crash/code cart)?	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
IT Capabilities:	
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Does your Facility have computers that are dedicated to research studies?	No
What browser does your facility use?	Microsoft Edge

Does the Facility have access to local IT support?	Yes
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
What type of computer operating system(s) does your institution use to support studies?	Windows 10
Labs:	
Is your Facility using a local pathology lab?	Yes
Please provide the Local Lab Name.	Pathology Queensland-Townsville Hospital
IP Storage Details:	
IP Recipient Name	Townsville University Hospital Pharmacy
Does the Investigational Product Storage Room have back-up power?	Yes
Is the Investigational Product Storage Room secured with controlled access?	Yes
Is the Investigational Product Storage Area securely constructed?	Yes
Does your Facility have the ability to manage on-site or off-site destruction of the Investigational Product?	Yes
Does your facility have a written SOP/Policy/Procedure for the destruction of Investigational Product?	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
Do you provide your Satellite Site(s) with a dedicated inventory of Investigational Product?	Yes

Does your Facility have a written SOP/Policy/Procedure to ensure that Investigational Product is appropriately maintained during transportation to Satellite Site(s)?	Yes	
Source Documents:		
Does your Facility have patient record archiving on-site?	Yes Off-site archiving	
Does your Facility have secure storage for patient records?	Yes	
Electronic Medical Records (EMR) / Electronic Health Records (EHR):		
Do you have Electronic Health Records (EHR)/ Electronic Medical Records (EMR)?	Yes	