Caboolture Hospital - Women's and Family

Local Services Offered:
Clinical trials site;Satellite 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:	
What is your Facility ID? **(Queensland Health only)**	30
Please provide your Facility Website.	https://metronorth.health.qld.gov.au/caboolture/
What Department is your Trial Site? (**Queensland Health HHS**- *See List of Services and Levels-Clinical Services Capability Framework)*	Medical Services
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
Is your facility/organisation a Life Sciences Queensland (LSQ) Member?	No
Please provide other areas of expertise for your Facility.	General Paediatrics; Maternity; Special Care Nursery; Integrated Care; Neurodevelopmental conditions.; Women and family; Cerebral Palsy; Hemeplagia
Provide the list of Sub-Therapeutic Areas for your Facility.	Womens; Family; Gynaecology; Obstetrics; Paediatrics
Does your Clinical Trial site undertake any recruitment?	Yes
What percentage of Clinical trials undertaken on your site do you meet or exceed the recruitment target?	80-90%
Are there any notable factors relating to your Patient Population?	Neonates; ATSI;
IRB/ERB/Ethics Committee:	

Does your Facility have a dedicated department or group to perform HREC (IRB/ERB/ETHICS) Committee submissions?	No
If your facility uses other HREC Commitee, please provide the HREC details here.	TPCH and QCH
What is the meeting frequency of your Local IRB/ERB/Ethics Committee?	Two meetings per week in NMA
Does your Facility have other review boards that need to approve the study prior to IRB/ERB/Ethics Committee submission? *For example, scientific, radiation safety committees, or others.*	Paediatrics through QCH; Adults through RBWH; TPCH
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 Other vulnerable populations?	Yes
Will your Facility require language translations for consents?	Yes
Training:	
Does your Facility have a training program for the research staff?	Yes
If your facility uses external program course/s. Please provide the external program course/s name.	N/A
Does your facility training course content include GCP?	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
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 patient visits on weekends?	Yes
Can your Facility support in-patient admissions for research studies?	Yes
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 the Facility have storage space for Study-Related materials (e.g. Lab Kits, Patient Materials, etc.)?	Yes
Is the Storage area for study related materials securely constructed?	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
Do you provide your Satellite Site(s) with a dedicated inventory of Investigational Product?	Yes
Equipment:	
Does your Facility have the necessary equipment to treat medical emergencies (for example crash/code cart)?	Yes
Please provide detail of the additional equipment that your facility uses for clinical trials.	Ultrasounds iMED - radiology imaging based in the hospital
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)
What browser does your facility use?	Internet Explorer
Does the Facility have access to local IT support?	Yes
Please indicate all equipment that will be available to Monitors	Phone;Copy Machines;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)?	Yes
Lab:	
Is your Facility using a local pathology lab?	Yes
Please provide the Local Lab Name.	Pathology Queensland-Caboolture Hospital
Does your Facility use private laboratory services?	Yes
IP Storage Details:	
Please provide the IP Recipient Name.	Caboolture Hospital
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 controlled substances when appropriate?	Yes
Does your facility have a written SOP/Policy/Procedure for the destruction of Investigational Product?	Yes
Source Documents:	
Does your Facility have patient record archiving on-site?	Yes
For Facilities with satellite sites, where is the monitor required to access source documents, please details the location of the monitor?	With researchers Metro North