Gold Coast Private Hospital

Local Services Offered:
Early Phase, Clinical trials site
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.	www.goldcoastprivatehospital.com.au
What Department is your Trial Site? (Queensland Health HHS- See List of Services and Levels-Clinical Services Capability Framework)	Oncology
Please provide other areas of expertise for your Facility.	Endometriosis
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.	All Healthscope Hospitals across Australia
Is your Facility affiliated with a government agency or part of a government funded health service?	No
Does your Clinical Trial site or Service undertake any recruitment?	Yes
Has your Clinical Trial Site or Service been accredited?	Yes
If your Clinical Trial Site has been accredited, please select all relevant types	NSQHS (National Safety and Quality Health Service) Standards
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?	No
Types of HREC (IRB/ERB/ETHICS) Committee that are used	Sponsor Provided Central
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
Any Additional Process	No
Does the HREC Committee require payment prior to the release of final approval documents?	No

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?	No
Does your Facility have access to translators and translation support for study conduct (e.g. consent, study-specific instruction)?	No
Training:	
Does your Facility have a training program for the research staff?	No
Does the course content include GCP?	Yes
If your facility uses external program course/s. Please provide the external program course/s name.	NIDA
Do you have a process or program in place to retrain research staff when a protocol is amended?	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
Facility And Equipment:	
Facility Capabilities:	
Can your Facility support in-patient admissions for research studies?	Yes
Can your Facility support patient visits on weekends?	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
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
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
Please list any additional equipment that your Facility uses for Clinical Trials.	Cardiac Catheter/Hybrid Lab
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
What browser does your facility use?	Chrome
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)?	No
Please indicate all equipment that will be available to Monitors	Phone, Fax, Copy Machines, Internet Access
Labs:	
Local Lab Usage	No
Does your Facility use private laboratory services?	Yes
If you selected 'Yes' on the previous question, please specify here which services	Other
Lab Name	QML, ACL and SydPath
IP Storage Details:	
IP Recipient Name	HPS Pharmacy-Gold Coast Private Hospital
Is the Investigational Product Storage Room secured with controlled access?	Yes
Does the Investigational Product Storage Room have Backup Power?	Yes

Does your Facility have the ability to manage on-site or off-site destruction of the Investigational Product?	Yes
Do you provide your Satellite Site(s) with a dedicated inventory 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
Electronic Medical Records (EMR) /Electronic Health Records (EHR):	
Do you have Electronic Health Records (EHR)/ Electronic Medical Records (EMR)?	Yes