

# Icon Cancer Care -Gold Coast Private Hospital

## Local Services Offered:

Early Phase;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.

<b>Facility Details:</b>	
Please provide your Facility Website.	<a href="http://www.iconcancercentre.com.au">www.iconcancercentre.com.au</a>
Is your Facility affiliated with a government agency or part of a government funded health service?	No
Do you have Affiliated Research Sites or Satellite Sites/Clinics? <i>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.</i>	Yes
Please provide other facility details.	Icon Cancer Centres
Is your facility/organisation a Life Sciences Queensland (LSQ) Member?	Yes
Provide the list of Sub-Therapeutic Areas for your Facility.	Radiation Oncology
Has your Clinical Trial Site been accredited?	Yes
If your Clinical Trial Site has been accredited, please select all relevant types.	ISO; NATA
Does your Clinical Trial site undertake any patient recruitment?	Yes
What percentage of Clinical trials undertaken on your site do you meet or exceed the recruitment target?	90%
Patient Population Comments	Icon is a network of private cancer treatment centres and the majority of patients required private health
<b>IRB/ERB/Ethics Committee:</b>	
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
Review Board Name	Bellberry

Are there any other steps that the Sponsor should be aware of for your IRB/ERB/Ethics Committee review and submission?	No
Does the HREC require contract/budget approval prior to release of final approval documents?	No
Does the HREC Committee require payment prior to the release of final approval documents?	Yes
<b>Consent:</b>	
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
<b>Training:</b>	
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.	Sponsor training
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
<b>Facility And Equipment:</b>	
<b>Facility Capabilities:</b>	
Can your Facility support in-patient admissions for research studies?	No

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 Facility typically allow the collection of Pharmacogenomic (PGX) samples for research purposes?	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 the Facility have storage space for Study-Related materials (e.g. Lab Kits, Patient Materials, etc.)?	Yes
<b>Equipment:</b>	
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.	Ultrasound
<b>IT Capabilities:</b>	
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?	Chrome
Does the Facility have access to local IT support?	Yes
Please indicate all equipment that will be available to Monitors	Copy Machines;Internet Access
<b>Labs:</b>	
Does your Facility use Local Lab Services?	No
Does your Facility use private laboratory services?	Yes

If Facility use private laboratory services, please provide the details.	Sullivan and Nicolaides Pathology
<b><u>IP Storage Details:</u></b>	
Do you provide your Satellite Site(s) with a dedicated inventory of Investigational Product?	Yes
Is the Investigational Product Storage Room secured with controlled access?	Yes
Does the Investigational Product Storage Room have Backup Power?	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 Investigational Product?	Yes
Does your facility have a written SOP/Policy/Procedure for the destruction of Investigational Product?	Yes
Does the Facility have the ability to handle radio-labelled Investigational Products?	No
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 to ensure that Investigational Product is appropriately maintained during transportation to Satellite Site(s)?	Yes
<b><u>Source Documents:</u></b>	
Does your Facility have secure storage for patient records?	Yes
Does your Facility have patient record archiving on-site?	No
Provide Location name and address of any offsite archives.	Grace Medical Records 4/20 Sherbrooke Willawong
What Electronic Data Capture (EDC) systems has your staff used for clinical trials?	Oracle Inform;Medidate Rave;Oracle Remote Data
<b><u>Electronic Medical Records (EMR) /Electronic Health Records (EHR):</u></b>	
Do you have Electronic Health Records (EHR)/ Electronic Medical Records (EMR)?	No
What EMR/EHR system do you use?	In-house system