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.

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

Are there any other steps that the Sponsor	No
should be aware of for your	
IRB/ERB/Ethics Committee review and	
submission?	
Does the HREC require contract/budget	No
1	110
approval prior to release of final approval	
documents?	
Does the HREC Committee require payment	Yes
prior to the release of final approval	
documents?	
documents.	
Consent:	
Does your Facility have a written	Yes
SOP/Policy/Procedure for Informed Consent?	
Does your Facility have a written	Yes
SOP/Policy/Procedure for Minor Assent for	
1	
paediatric populations?	
Does your Facility have a written	Yes
SOP/Policy/Procedure for Other vulnerable	
populations?	
Does your Facility have access to translators	Yes
and translation support for study conduct (e.g.	
consent, study- specific instruction)?	
Training:	
Does your Facility have a training	Yes
program for the research staff?	
Does your Facility training course content	Yes
include GCP?	
	G
If your facility uses external program course/s.	Sponsor training
Please provide the program course/s name.	
Does the study staff that prepares or transports	Yes
dangerous goods have training that meets the	
IATA International Air Transport Association	
(US) or other countries hazardous training	
requirements for shipping dangerous goods?	
Do you have a process or program in place to	Yes
retrain research staff when a protocol is	
amended?	
amended:	
Facility And Equipment:	
Facility Capabilities:	
Can your Facility support in-patient admissions	No
for research studies?	
101 1050atoti studies:	

Can your Facility support patient visits on weekends? Is your Facility capable of administering infusions? Is your Facility adequately staffed to support studies with both blinded and unblinded Investigational Product? Does your Facility typically allow the collection of Pharmacogenomic (PGX) samples for research purposes? Does your Facility have the ability to collect PK/PD specimens? Does your Facility have the ability to collect PK/PD samples beyond normal business hours? Does the Facility have storage space for Study-Related materials (e.g. Lab Kits, Patient Materials, etc.)? Equipment: Does your Facility have the necessary equipment to treat medical emergencies (for example crash/code cart)? 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.? Please list any additional equipment that your Facility uses for Clinical Trials. IT Capabilities: Does your Facility have computers that are dedicated to research studies? What type of computer operating system(s) does your institution use to support studies? What type of computer operating system(s) does your institution use to support studies? What type of computer operating system(s) does your institution use to support studies? What type of computer operating system(s) does your institution use to support studies? What type of computer operating system(s) does your facility use? Does the Facility have access to local IT support? Please indicate all equipment that will be available to Monitors Labs: Does your Facility use Local Lab Services? No Does your Facility use private laboratory services?		
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Does your Facility use Local Lab Services? No Does your Facility use private Yes		
Does your Facility use Local Lab Services? No Does your Facility use private Yes	Labs:	
Does your Facility use private Yes		No
		Yes
	laboratory services?	

If Facility use private laboratory services,	Sullivan and Nicolaides Pathology
please provide the details.	Sumvan and Tyleolaides I amology
preuse provide the details.	
ID Stayaga Dataila	
IP Storage Details:	Yes
Do you provide your Satellite Site(s) with a	i es
dedicated inventory of	
Investigational Product?	Yes
Is the Investigational Product Storage Room secured with controlled access?	res
	Yes
Does the Investigational Product Storage Room have Backup Power?	i es
Is the Investigational Product Storage area	Yes
securely constructed?	i es
Does your Facility have the ability to manage	Yes
on-site or off-site destruction of Investigational	1 es
Product?	
Does your facility have a written	Yes
SOP/Policy/Procedure for the destruction of	105
Investigational Product?	
Does the Facility have the ability to handle	No
radio-labelled Investigational Products?	
Does your Facility have the ability to manage	Yes
on-site or off-site destruction of controlled	
substances when appropriate?	
Does your Facility have a written	Yes
SOP/Policy/Procedure to ensure that	
Investigational Product is appropriately	
maintained during transportation to Satellite	
Site(s)?	
, ,	
Source Documents:	
Does your Facility have secure storage for	Yes
patient records?	
Does your Facility have patient record	No
archiving on-site?	
Provide Location name and address of	Grace Medical Records 4/20 Sherbrooke
any offsite archives.	Willawong
What Electronic Data Capture (EDC) systems	Oracle Inform; Medidate Rave; Oracle Remote
has your staff used for clinical trials?	Data
Electronic Medical Records (EMR) /Electron	ic Health Records (EHR):
Do you have Electronic Health Records (EHR)/	
Electronic Medical Records	
(EMR)?	
What EMR/EHR system do you use?	In-house system