Sunshine Coast University Private Hospital - Clinical Trials Unit

Location:

-26.744073 - 153.114366

Local Services Offered:

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

Details: In addition to the information contained within csv generated by this website, this PDF provides additional site information

Does your Facility perform HREC (IRB/ERB/Ethics) Committee submissions?	Yes
Does your Facility have a written SOP/Policy/Procedure for Informed Consent?	Yes
Does your Facility have a dedicated department or group to perform HREC (IRB/ERB/ETHICS) Committee submissions?	No
Does your Facility have the necessary equipment to treat medical emergencies (for example crash/code cart)?	Yes
Is your Facility affiliated with a government agency or part of a government funded health service?	No
Does your Facility typically allow the collection of Pharmacogenomic (PGX) samples for research purposes?	Yes
Does your Facility have computers that are dedicated to research studies?	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?	No
Is your Facility capable of administering infusions?	Yes
What browser does your facility use?	Internet Explorer

Yes
Yes
Yes
Yes
Ramsay Healthcare
No
Yes
No
Yes
Not Applicable
Yes
Yes
Yes
Yes
No
I
Yes
Yes

Do you have Electronic Health Records (EHR)/ Electronic Medical Records (EMR)?	No
Does your Facility have a written	No
SOP/Policy/Procedure for Minor Assent	
for paediatric populations?	
Does your Facility have a written	No
SOP/Policy/Procedure for Other	
vulnerable populations?	
Please provide program course/s name	The Global Health Network Online; IQVIA
IP Recipient Name	Sunshine Coast University Private Hospital Pharmacy
Does the HREC Committee require	Yes
payment prior to the release of final	
approval documents?	
Does your Facility have access to	Yes
translators and translation support for	1 65
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study conduct (e.g. consent, study-	
specific instruction)?	
Do you provide your Satellite Site(s)	Yes
with a dedicated inventory of	
Investigational Product?	
Do you have a process or program in	Yes
place to retrain research staff when a	
protocol is amended?	
Does your Facility have a training	Yes
program for the research staff?	
Storage Room Backup Power	Yes
Is the Investigational Product Storage	Yes
Room secured with controlled access?	
Storage area securely constructed	Yes
Other review boards	No
Additional equipment	Ultrasound
What type of computer operating	Phone;Copy Machines;Internet Access
system(s) does your institution use to	
support studies?	
Has your Clinical Trial Site been	No
accredited?	
	No
Has your Clinical Trial site been audited?	INO
Does your Clinical Trial site undertake	Yes
any patient recruitment?	
Location of Monitor	In Clinical Trials Unit
Additional General information	SSA and approval by dedicated Advisory Committee
Any Additional Process	No S
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