

John Flynn Private Hospital

Local Services Offered:

Clinical trials site, Trial Patient Recruitment, Treatment of patients, Completion of study documenta contract

Details: This PDF complements the information found in the CSV generated by this website with ad

<u>Facility Details:</u>	
Please provide your Facility Website.	https://www.johnflynnprivate.com.au/
Is your Facility affiliated with a government agency or part of a government funded health service?	No
Is your facility/organisation a Life Sciences Queensland (LSQ) Member?	No
Please provide other areas of expertise for your Facility.	Oncology, Haematology, Orthopaedics, Cardiology and Women's Health
Has your Clinical Trial Site been accredited?	Yes
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% to date
<u>IRB/ERB/Ethics Committee:</u>	
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
Does the HREC Committee require payment prior to the release of final approval?	Yes
<u>Consent:</u>	
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?	No
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

<u>Training:</u>	
Does your Facility have a training program for the research staff?	Yes
Does your Facility training course content include GCP?	No
If your facility uses external program course/s. Please provide the program course/s name.	Via Praxis International
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
<u>Facility And Equipment:</u>	
<u>Facility Capabilities:</u>	
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 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	Yes
Does your Facility typically allow the collection of Pharmacogenomic (PGX) samples for research purposes?	Yes
Does the Facility have storage space for Study Related materials (e.g. Lab Kits, Patient Materials, etc.)?	Yes
<u>Equipment:</u>	
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,	Yes
<u>IT Capabilities:</u>	
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
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
Please indicate all equipment that will be available to Monitors	Fax; Copy Machines
<u>Labs:</u>	
Does your Facility use private laboratory services?	Yes
If you selected 'Yes' on the previous question, please specify here which services.	Sullivan Nicolaides Pathology (SNP)
<u>IP Storage Details:</u>	
IP Storage Location Name	John Flynn Private Hospital
Is the Investigational Product Storage area securely constructed?	Yes
Is the Investigational Product Storage Room secured with controlled access?	Yes
Does the Investigational Product Storage Room have back-up power?	Yes
Does your Facility have the ability to manage on-site or off-site destruction of the 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?	Yes

Does your Facility have the ability to manage on-site or off-site destruction of controlled substances when appropriate?	Yes
Do you provide your Satellite Site(s) with a dedicated inventory of Investigational Product?	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
<u>Source Documents:</u>	
Does your Facility have patient record archiving on-site?	Yes
Does your Facility have secure storage for patient records?	Yes
Provide Location name and address of any offsite archives.	ZircoData 772 Boundary Road Richlands QLD 4077 Brisbane
What Electronic Data Capture (EDC) systems has your staff used for clinical trials?	Oracle Inform;Medidata
Please describe Other EDC Systems:	CRF Health ERT Viedoc BMS Mytrials Novartis proprietary (in
<u>Electronic Medical Records (EMR) /Electronic Health Records (EHR):</u>	
Do you have Electronic Health Records (EHR)/ Electronic Medical Records (EMR)?	No
What EMR/EHR system do you use? Electronic Medical Records (EMR) /Electronic Health Records	In-house system; Others

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ditional site information.