

Toowoomba Hospital - Cancer Care Services- Oncology Haematology

Local Services Offered:

Clinical trials site;Investigator Initiated Trials;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.

<u>Facility Details:</u>	
Please provide your Facility Website.	https://www.darlingdowns.health.qld.gov.au/hospitals-and-health-centres/toowoomba-hospital
What Department is your Trial Site? (Queensland Health HHS- See List of Services and Levels-Clinical Services Capability Framework)	Medical Services
Is your Facility affiliated with a government agency or part of a government funded health service?	Yes
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.	N/A
Is your facility/organisation a Life Sciences Queensland (LSQ) Member?	No
Please provide other areas of expertise for your Facility.	General Medicine; ATSI; CNS; ED; Geriatric; imaging; Renal; Respiratory; Dietetics; Nervous System Diseases;Prostate Cancer;Sarcoma;Ovarian Cancer;Chronic Lymphocytic Leukaemia;Lymphocytic

Provide the list of Sub-Therapeutic Areas for your Facility.	Bladder cancer;Bone Cancer;Brain Cancer;Breast Cancer; Cervical Cancer;Colorectal Cancer; CRPC; Follicular Lymphoma;Gastro Intestinal Solid Tumours; Head and Neck Cancer; Hepatocellular Carcinoma; Hereditary Angioedema;HR Prostate Cancer;Islet Cell Tumours;Liver Cancer;Lung Cancer; Lymphoma; Malignant Pleural Mesothelioma;Melanoma;Multiple Myeloma;Neuroma;Non- Hodgkin's Lymphoma;Non-Small Cell Lung Cancer;Ovarian Epithelial Carcinoma; Pancreatic Cancer;Renal Cell Carcinoma;Soft Tissues Sarcomas;Squamous & Non Squamous Sarcomas; Thyroid Cancer;Throat Cancer;Acquired Haemophilia; Acute Lymphoblastic Leukaemia;Acute Myelogenous Leukaemia;Chronic Myelogenous Leukaemia
Does your Clinical Trial site undertake any patient recruitment?	Yes
<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
HREC Committee Name.	Darling Downs HHS HREC
What is the meeting frequency of your Local IRB/ERB/Ethics Committee?	Monthly
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 Committee require payment prior to the release of final approval documents?	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?	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
<u>Training:</u>	
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	Caledonian; ARCS; Syneos online, Praxis
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?	No
Is your Facility capable of administering infusions?	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?	No
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
<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, etc.?	Yes
Please list any additional equipment that your Facility uses for Clinical Trials.	Ultrasounds, CT Scan, Bone densitometry, ECG, Echo, Fluoroscopy, Mammography, MRA, MRI, PET scan, X-Ray,
<u>IT Capabilities:</u>	
Does your Facility have computers that are dedicated to research studies?	No
What type of computer operating system(s) does your institution use to support studies?	Windows
What browser does your facility use?	Edge
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	Phone;Copy Machines;Internet Access
<u>Labs:</u>	

Is your Facility using a local pathology lab?	Yes
Please provide the Local Lab Name.	Pathology Queensland- Toowoomba Hospital
<u>IP Storage Details:</u>	
IP Recipient Name	Toowoomba Hospital Pharmacy
Is the Investigational Product Storage Room secured with controlled access?	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 the Investigational Product?	Yes
Does your facility have a written SOP/Policy/Procedure for the destruction 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
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
<u>Electronic Medical Records (EMR) /Electronic Health Records (EHR):</u>	
Do you have Electronic Health Records (EHR)/ Electronic Medical Records (EMR)?	No