Royal Brisbane and Women's Hospital - Motor Neurone Disease MND

Local Services Offered:

Clinical trials site, Trial Patient Recruitment, Treatment of patients, Completion of study documentation as per ICH GCP.

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.	https://metronorth.health.qld.gov.au/rbwh/
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
Is your facility/organisation a Life Sciences Queensland (LSQ) Member?	No
Please provide other areas of expertise for your Facility.	Other neurological condition including Pompe disease; Chronic idiopathic demyelinating polyneuropath
Provide the list of Sub-Therapeutic Areas for your Facility.	Neurology; Motor Neurone Disease
IRB/ERB/Ethics Committee:	
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.	Metro North HHS HREC
Other Meeting Frequency	Two meetings per week in NMA
Does the HREC Committee require payment prior to the release of final approval documents?	Yes
Consent:	
Does your Facility have a written SOP/Policy/Procedure for Informed Consent?	Yes

Does your Facility have a written	No
SOP/Policy/Procedure for Minor	
Assent for paediatric populations?	
Does your Facility have a written	Yes
SOP/Policy/Procedure for Other	
vulnerable populations?	
Does your Facility have access to	Yes
translators and translation support for	
study conduct (e.g. consent,	
study-specific instruction)?	
<u>Training:</u>	
Does your Facility have a training program	Yes
for the research staff?	
Does your Facility training course content	Yes
include GCP?	
If your facility uses external program	Caledonian; ARCS and Syneos online
course/s. Please provide the program	
course/s name.	
Do you have a process or program in place	Yes
to retrain research staff when a protocol is	
amended?	~ ~
Does the study staff that prepares or	Yes
transports dangerous goods have training that meets the IATA International Air	
Transport Association (US) or other	
countries hazardous training requirements	
for shipping dangerous goods?	
for simpping aangerous goods.	
Facility And Equipment:	
Facility Capabilities:	
Can your Facility support in-patient	Yes
admissions for research studies?	
Can your Facility support patient	Yes
visits on weekends?	
Is your Facility capable of	Yes
administering infusions?	
Is your Facility adequately staffed to	Yes
support studies with both blinded and	
unblinded Investigational	
Product?	
Does your Facility have the ability to	Yes
collect and store PK/PD	
specimens?	

Does your Facility have the ability to	Yes
collect PK/PD samples beyond	
normal business hours?	
Does your Facility typically allow the	Yes
collection of Pharmacogenomic (PGX)	
samples for research	
purposes?	
Does the Facility have storage space for	Yes
Study-Related materials (e.g.	
Lab Kits, Patient Materials, etc.)?	
Equipment:	
Does your Facility have the necessary	Yes
equipment to treat medical emergencies	
(for example	
crash/code cart)?	
Does your Facility have an SOP or process	Yes
that ensures routine calibration and	
maintenance of general equipment?	
Examples of general equipment include:	
scale, pulse oximeter, stadiometer,	
sphygmomanometer, etc.?	
IT Capabilities:	
Does your Facility have computers that are	No
dedicated to research studies?	
What type of computer operating	Windows
system(s) does your institution use to	
support studies?	
What browser does your facility	Microsoft Explorer 11, EDGE
use?	
Does the Facility have access to	Yes
local IT support?	
Does your Facility limit or prohibit access	Yes
and use of external web- based tools or	
sites for clinical research (E.g. web portals	
to submit documents to sponsors or	
CROs)?	
Labs:	
Is your Facility using a local pathology	Yes
lab?	
Please provide the Local Lab Name.	Pathology Queensland- RBWH

IP Recipient Name	Royal Brisbane and Women's Hospital Pharmacy
Does the Investigational Product Storage Room have back-up power?	Yes
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
Source Documents:	
Does your Facility have patient record archiving on-site?	Yes
Does your Facility have secure storage for patient records?	Yes
Electronic Medical Records (EMR) /Elec	tronic Health Records (EHR):
Do you have Electronic Health Records (EHR)/ Electronic Medical Records (EMR)?	Yes