# Mater Clinical Trials

Local Services Offered:
Clinical Trial Site;Clinical Trial Administration
Details: In addition to the information contained within csv generated by this website, this PDF provides additional site information

| **Facility Details:** | |
|---|---|
| Please provide your Facility Website. | https://www.materresearch.org.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. | Telehealth |
| | |
| **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 |
| What is the meeting frequency of your Local IRB/ERB/Ethics Committee? | Monthly |
| Does the HREC Committee require payment prior to the release of final approval | Yes |
| | |
| **Consent:** | |
| 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? | N/A |
| Does your Facility have a written SOP/Policy/Procedure for Other vulnerable populations? | Yes |

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| Does your Facility have access to translators and translation support for study conduct (e.g. consent, study-specific instruction)? | Yes |
| | |
| **Training:** | |
| Does your Facility have a training program for the research staff? | Yes |
| Does your Facility training course content include GCP? | Yes |
| 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 |
| | |
| **Facility And Equipment:** | |
| **Facility Capabilities:** | |
| 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 hours? | Yes |
| Does your Facility typically allow the collection of Pharmacogenomic (PGX) samples for research purposes? | Yes |

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| Does the Facility have storage space for Study- Related materials (e.g. Lab Kits, Patient Materials, etc.)? | Yes |
| | |
| **Equipment:** | |
| 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 |
| | |
| **IT Capabilities:** | |
| 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 10 |
| What browser does your facility use? | Microsoft Edge |
| Does the Facility have access to local IT | 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)? | No |
| | |
| **Labs:** | |
| Does your Facility use private laboratory services? | Yes |
| If you selected 'Yes' on the previous question, please specify here which services. | Mater Pathology |
| | |
| **IP Storage Details:** | |
| IP Recipient Name | Mater Hospital Pharmacy Dept- Clinical Trials |

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| 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 |
| 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 |
| 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 secure storage for patient records? | Yes |
| Does your Facility have patient record archiving on-site? | Yes |
| Provide Location name and address of any offsite archives. | Grace Document and Record Management, 420 Sherbrooke Road, Willawong QLD |
| | |
| **Electronic Medical Records (EMR) /Electronic Health Records (EHR):** | |
| Do you have Electronic Health Records (EHR)/ Electronic Medical Records (EMR)? | No |