Austrials

Local Services Offered: Clinical trials CRO
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?	Yes
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 the Investigational Product?	Yes
Is your Facility capable of administering infusions?	Yes
What browser does your facility use?	Internet Explorer
Does the Facility have access to local IT support?	Yes
Please list any access limitations/requirements for the Electronic Medical Records	Yes
Does your Facility have the ability to manage on-site or off-site destruction of the Investigational Product?	Not Applicable
Other facility details	Yes
Does your Facility have patient record archiving on-site?	No

Does the Facility have storage space for Study-Related materials (e.g. Lab Kits, Patient Materials, etc.)?	Yes
Can your Facility support in-patient admissions for research studies?	No
Does your Facility have the ability to collect and store PK/PD specimens?	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
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
Does your Facility have the ability to collect PK/PD samples beyond normal business hours?	Yes
Is your Facility adequately staffed to support studies with both blinded and unblinded Investigational Product?	Yes
Study Space	Yes
Can your Facility support patient visits on weekends?	No
Does the course content include GCP?	Yes
Is your Facility using a local pathology lab?	No
Do you have Electronic Health Records (EHR)/ Electronic Medical Records (EMR)?	No
LSQ Member	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
Please provide program course/s name	IATA; IQVIA and Whitehall
IP Recipient Name	Westside Hospital-Jemimah Sguena
Does the HREC Committee require payment prior to the release of final approval documents?	Yes

Will your Facility require language translations for consents?	No
Do you provide your Satellite Site(s) with a dedicated inventory of Investigational Product?	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
Does your Facility have a training program for the research staff?	Yes
Controlled Access in Storage Room	Yes
What type of computer operating system(s) does your institution use to support studies?	Windows (Windows XP; Windows 7; Windows 10; etc)
Sub-Therapeutic areas	Endometriosis;depression and anxiety.
Provide Location name and address of any offsite archives.	Grace
EDC Systems Description	CLIMCAL INK
Please describe Other EDC Systems:	Oracle Inform;Medidate Rave;Oracle Remote Data Capture (RDC);Others
Please indicate all equipment that will be available to Monitors	Phone;Fax;Copy Machines;Internet Access
Has your Clinical Trial Site been accredited?	No
If your Clinical Trial Site has been audited, please select all relevant types.	Yes
Audit Type	FDA;CRO;Sponsor
What is your facility's Internet download speed?	100/100
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?	80%
For Facilities with satellite sites, where is the monitor required to access source documents, please details the location of the monitor?	At site

What is your facility's internet upload speed?	Fibre 100/100
Additional General information	We provide a coordinating investigator to support our network at multiple sites
Does your Facility use private laboratory services?	Yes
Please list any access limitations/requirements for the Electronic Medical Records	No CRA access to eHR - print off a paper source
Patient Population Comments	General practice population
Additional Address Info	Taringa
Does the HREC require contract/budget approval prior to release of final approval documents?	No
Any Additional Process	No