Section VI
Schedule of Requirements
IB No. E-003-2024-PB

The delivery schedule expressed as weeks/months stipulates hereafter a delivery date which is the date of delivery to the project site.

Item No.	Item Code	Description	Qty	Delivery Site			Delivery Period and Terms of Payment
				OFFICE	FACILITY	ADDRESS	
1	CL-01-2024	**Supply, Delivery, Installation, Testing Commissioning of a Brand-New Ultra Low Freezer, -80 C** **SPECIFICATIONS:** 1. Capacity: at least 18 cubic feet. 2. Temperature range: -40°C to -86°C. 3. Temperature Chart Recorder. 4. Alarm system: High/Low temperature, sensor alarm. 5. Door: 1-door, self-closing, safety lock, with door handle. 6. At least 3 shelves. 7. Type: Upright. 8. Heavy Duty compressor, HFC refrigerant/ CFC Free. 9. Caster wheels with lock. 10. With LED light. 11. Must come with an AVR at least 5000VA. 12. The unit will be operated at 220 volts, 60hz.. **STANDARD REQUIREMENTS:** 1. The bidder must provide the current and valid certificate of	1	Clinical Laboratory Section	NCMH New Laboratory Building	National Center for Mental Health	a. One-time delivery within **30 calendar days upon receipt of Notice to Deliver** b. Terms of payment at least 30 days from receipt of Sales invoice

		authorized or exclusive distributorship. 2. The bidder must provide the current and valid certificate of Manufacturer's compliance with ISO 13485. 3. The bidder/supplier shall secure a certificate from the manufacturer stating that the equipment is brand new, unused and not a discontinued model or was listed in the market recall. 4. Bidder's certificate guaranteeing the availability on the supply of spare parts ten (10) years from the end of production. A certification must be issued by the manufacturer for the bidder. 5. Certification that the bidder must be in the local market for a minimum of ten (10) years. 6. Certification that the system or machine is not a retrofit solution. 7. The machine should have a US FDA approval. The bidder shall provide the manufacturer's approved US FDA Premarket Notification (PMN) or CE mark approval certificate. 8. The winning bidder shall provide current and valid calibration certificate for each equipment during delivery. 9. Certification that the supplier has the capability or authority for corrective and preventive maintenance of the unit. The certificate must be issued by the manufacturer to the bidder. 10. Certification from the manufacturer authenticated by the Philippine Consulate from					

		the country of origin of the unit that the warranty should not be affected with a change of distributor. 11. Warranty Certification: Certification that the bidder/supplier shall provide a three (3) year warranty for parts and services that includes corrective maintenance, preventive maintenance, and/or calibration. The warranty shall commence upon the acceptance of the end-user. 12. Bidder must provide preventive maintenance and/or calibration schedule within warranty period. 13. Certification that the supplier / bidder shall provide applications training for the users and maintenance personnel of the hospital. 14. Certification to provide manuals; Two (2) sets of service manual in English Language and Two (2) sets of user manual in English Language upon delivery of the equipment. 15. Certification that the brand has been in the local market for at least five (5) years with at least 5 current installations. 16. Certification that the supplier will be responsible for the notification, transportation to the site, delivery, installation and testing on the site (hospital / health facility) expenses for such will be on the account of the supplier. 17. Certification that the supplier / bidder shall provide free installation of the equipment. 18. With 24/7 Technical Support.					

		19. Delivery period –30 calendar days.					
		Supply, Delivery, Installation, Testing and Commissioning of a Brand-New Blood Bank Refrigerator 120 blood bag capacity **SPECIFICATIONS:** 1. Alarm and Safety Function. -Audible and flashing LED visual alarms. - Door alarm and key lock are standard feature. - Built in temperature recorder. 2. Stable temperature control. 3. Effective Capacity: at least 302 liter (10.7 cu ft) which equivalent to 120 bags or higher. 4. Exterior and interior Cabinet: Galvanized steel with baked-on finish. 5. Outer Door: Insulated Steel frame with double layer glass windows. 6. Drawers: at least 5 stainless steels roll out drawers. 7. Caster wheels with lock. . 8. Compressor: 150W hermetically sealed. 9. Refrigerant: HFC. 10. Temperature: 4C (+/-1.5C). 11. Air Circulation: Forced circulation. 12. Temperature control: Microprocessor control. 13. Temperature alarm: High (6C), low (2C) Audible and visual alarm. 14. Power Failure Alarm: Audible and visual alarm. 15. Lighting: 12-15W fluorescent lamp/LED. 16. The unit will be operated at 220-240volts. 17. Must come with an AVR: 2000KVA. **STANDARD REQUIREMENTS:** 1. The bidder must provide the current and	1	Clinical Laboratory Section	NCMH New Laboratory Building	National Center for Mental Health	a. One-time delivery within **30 calendar days upon receipt of Notice to Deliver** b. Terms of payment at least 30 days from receipt of Sales invoice

		valid certificate of authorized or exclusive distributorship. 2. The bidder must provide the current and valid certificate of Manufacturer's compliance with ISO 13485. 3. The bidder/supplier shall secure a certificate from the manufacturer stating that the equipment is brand new, unused and not a discontinued model or was listed in the market recall. 4. Bidder's certificate guaranteeing the availability on the supply of spare parts ten (10) years from the end of production. A certification must be issued by the manufacturer for the bidder. 5. Certification that the bidder must be in the local market for a minimum of ten (10) years. 6. Certification that the system or machine is not a retrofit solution. 7. The machine should have a US FDA approval. The bidder shall provide the manufacturer's approved US FDA Premarket Notification (PMN) or CE mark approval certificate. 8. The winning bidder shall provide current and valid calibration certificate for each equipment during delivery. 9. Certification that the supplier has the capability or authority for corrective and preventive maintenance of the unit. The certificate must be issued by the manufacturer to the bidder. 10. Certification from the manufacturer authenticated by the					

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		19. Delivery period – 30 calendar days.					
		Supply, Delivery, Installation, Testing and Commissioning of a Brand-New Plasma Thawer **SPECIFICATIONS:** 1. Thaws FFP bags and warms blood, erythrocyte concentrates (EC), cryopreserved preparation, cryopreserved stem cells (HPC) and infusion solutions 2. Thaws at least 4 bags at a time 3. Heating element not in direct contact to bags 4. Leak detection sensors-no risk of contamination or infection 5. User programmable modes 6. Large display LCD display. 7. With cushions conforms to bags 8. Adjustable temperature range from 37C to 45C 9. Visual and acoustic signal after heating duration is complete or in case of error 10. Power supply: 220-240V; 60Hz **STANDARD REQUIREMENTS:** 1. The bidder must provide the current and valid certificate of authorized or exclusive distributorship.	1	Clinical Laboratory Section	NCMH New Laboratory Building	National Center for Mental Health	a. One-time delivery within **30 calendar days upon receipt of Notice to Deliver** b. Terms of payment at least 30 days from receipt of Sales invoice
		2. The bidder must provide the current and valid certificate of Manufacturer's compliance with ISO 13485. 3. The bidder/supplier shall secure a certificate from the manufacturer stating that the equipment is brand new, unused and not a discontinued model or was listed in the market recall.					

		4. Bidder's certificate guaranteeing the availability on the supply of spare parts ten (10) years from the end of production. A certification must be issued by the manufacturer for the bidder. 5. Certification that the bidder must be in the local market for a minimum of ten (10) years. 6. Certification that the system or machine is not a retrofit solution. 7. The machine should have a US FDA approval. The bidder shall provide the manufacturer's approved US FDA Premarket Notification (PMN) or CE mark approval certificate. 8. The winning bidder shall provide current and valid calibration certificate for each equipment during delivery. 9. Certification that the supplier has the capability or authority for corrective and preventive maintenance of the unit. The certificate must be issued by the manufacturer to the bidder. 10. Certification from the manufacturer authenticated by the Philippine Consulate from the country of origin of the unit that the warranty should not be affected with a change of distributor. 11. Warranty Certification: Certification that the bidder/supplier shall provide a three (3) year warranty for parts and services that includes corrective maintenance, preventive maintenance, and/or calibration. The warranty shall commence					

		upon the acceptance of the end-user. 12. Bidder must provide preventive maintenance and/or calibration schedule within warranty period. 13. Certification that the supplier / bidder shall provide applications training for the users and maintenance personnel of the hospital. 14. · Certification to provide manuals; Two (2) sets of service manual in English Language and Two (2) sets of user manual in English Language upon delivery of the equipment. 15. Certification that the brand has been in the local market for at least five (5) years with at least five (5) current installations. 16. Certification that the supplier will be responsible for the notification, transportation to the site, delivery, installation and testing on the site (hospital / health facility) expenses for such will be on the account of the supplier. 17. Certification that the supplier / bidder shall provide free installation of the equipment. 18. With 24/7 Technical Support. 19. Delivery period – 30 calendar days.					
		Supply, Delivery, Installation, Testing and Commissioning of a Brand-New Platelet Agitator with incubator **SPECIFICATIONS:** 1. Refrigeration & precision pulse heating for achieving 22°C (± 1°C) 2. Forced air circulation for temperature uniformity	1	Clinical Laboratory Section	NCMH New Laboratory Building	National Center for Mental Health	a. One-time delivery within **30 calendar days upon receipt of Notice to Deliver** b. Terms of payment at least 30 days from receipt

		3. Powder coated exterior with high-grade stainless-steel interior 4. LCD display of temperature, real date, real time, temperature histogram etc. 5. Auto stop / start feature pauses agitation when incubator door is opened 6. Intuitive thermostat to prevent temperature fluctuations in additional to microprocessor controller 7. Swivel lockable castors with floor standing jacks 8. Triple pane tempered glass door agitator 9. Built in stopper prevents accidental drawing out 10. Each drawer can be operated independently from other drawers 11. Motor cooling done by fan (cooler) 12. Capacity: 60 bags 13. Power supply: 220V 14. Back up battery in case of power failure 15. Monitor and printer for weekly printout **STANDARD REQUIREMENTS:** 1. The bidder must provide the current and valid certificate of authorized or exclusive distributorship. 2. The bidder must provide the current and valid certificate of Manufacturer's compliance with ISO 1348. 3. The bidder/supplier shall secure a certificate from the manufacturer stating that the equipment is brand new, unused and not a discontinued model or was listed in the market recall. 4. Bidder's certificate guaranteeing the availability on the supply of spare parts ten (10) years from the end of					of Sales invoice

		production. A certification must be issued by the manufacturer for the bidder. 5. Certification that the bidder must be in the local market for a minimum of ten (10) years. 6. Certification that the system or machine is not a retrofit solution. 7. The machine should have a US FDA approval. The bidder shall provide the manufacturer's approved US FDA Premarket Notification (PMN) or CE mark approval certificate. 8. The winning bidder shall provide current and valid calibration certificate for each equipment during delivery. 9. Certification that the supplier has the capability or authority for corrective and preventive maintenance of the unit. The certificate must be issued by the manufacturer to the bidder. 10. Certification from the manufacturer authenticated by the Philippine Consulate from the country of origin of the unit that the warranty should not be affected with a change of distributor. 11. Warranty Certification:					
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		schedule within warranty period. 13. Certification that the supplier / bidder shall provide applications training for the users and maintenance personnel of the hospital. 14. Certification to provide manuals; Two (2) sets of service manual in English Language and Two (2) sets of user manual in English Language upon delivery of the equipment. 15. Certification that the brand has been in the local market for at least five (5) years with at least five (5) current installations. 16. Certification that the supplier will be responsible for the notification, transportation to the site, delivery, installation and testing on the site (hospital / health facility) expenses for such will be on the account of the supplier. 17. Certification that the supplier / bidder shall provide free installation of the equipment. 18. With 24/7 Technical Support. 19. Delivery period – 30 calendar days.					
		Supply, Delivery, Installation, Testing and Commissioning of a Brand-New Blood Bag Mixer **SPECIFICATIONS:** 1. Linear and 3d mixing with central support 2. Weighing range: 0-990ml. 3. Alarm type: Automatic low and high flow alarm, adjustable. 4. Adjustable maximum donation time: Up to 20 minutes. 5. Protocols: Up to 20 different working programs.	1	Clinical Laboratory Section	NCMH New Laboratory Building	National Center for Mental Health	a. One-time delivery within **30 calendar days upon receipt of Notice to Deliver** b. Terms of payment at least 30 days from receipt of Sales invoice

		6. Built-in memory: More than 1000 collection procedure data 7. Automatic clamping module **STANDARD REQUIREMENTS:** 1. The bidder must provide the current and valid certificate of authorized or exclusive distributorship. 2. The bidder must provide the current and valid certificate of Manufacturer's compliance with ISO 13485. 3. The bidder/supplier shall 'secure a certificate from the manufacturer stating that the equipment is brand new, unused and not a discontinued model or was listed in the market recall. 4. Bidder's certificate guaranteeing the availability on the supply of spare parts ten (10) years from the end of production. A certification must be issued by the manufacturer for the bidder. 5. Certification that the bidder must be in the local market for a minimum of ten (10) years. 6. Certification that the system or machine is not a retrofit solution. 7. The machine should have a US FDA approval. The bidder shall provide the manufacturer's approved US FDA Premarket Notification (PMN) or CE mark approval certificate. 8. The winning bidder shall provide current and valid calibration certificate for each equipment during delivery. 9. Certification that the supplier has the capability					

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		(hospital / health facility) expenses for such will be on the account of the supplier. 17. Certification that the supplier / bidder shall provide free installation of the equipment. 18. With 24/7 Technical Support. 19. Delivery period – 30 calendar days.					
		ITEMS NO. 1 to 5 UNDER LOT BIDDING. The winning bidder shall provide 2 units of blood extraction chair for general laboratory phlebotomy use and 2 units blood bag tube stripper with the following specifications: **Blood extraction chair:** 1. Stainless steel tubular frame 2. With built in stainless drawer with at least one (1) layer 3. Upholstered cushion (Polyurethane material) 4. Adjustable armrest (swing-out) **Blood bag tube stripper:** 1. Spring loaded handle 2. Easy grip handle 3. Capable of sealing clip and cutting tubes					
2	**CL-02-2024**	**Supply, Delivery, Installation, Testing and Commissioning of a Brand-New Blood Donor Chair** **SPECIFICATIONS:** 1. With optional apparatus tray and adjustable IV stand 2. Adjustable headrest 3. Quick fold arm rest 4. With one-touch rocker switch enabling quick donor's head lowering (Trendelenburg position) 5. Fiber reinforced plastic base 6. Four caster wheels **STANDARD REQUIREMENTS:** 1. The bidder must provide the current and valid certificate of authorized or exclusive distributorship. 2. The bidder must provide the current and valid certificate of Manufacturer's	1	Clinical Laboratory Section	NCMH New Laboratory Building	National Center for Mental Health	a. One-time delivery within **30 calendar days upon receipt of Notice to Deliver** b. Terms of payment at least 30 days from receipt of Sales invoice

		compliance with ISO 13485. 3. The bidder/supplier shall secure a certificate from the manufacturer stating that the equipment is brand new, unused and not a discontinued model or was listed in the market recall. 4. Bidder's certificate guaranteeing the availability on the supply of spare parts ten (10) years from the end of production. A certification must be issued by the manufacturer for the bidder. 5. Certification that the bidder must be in the local market for a minimum of ten (10) years. 6. Certification that the system or machine is not a retrofit solution. 7. The machine should have a US FDA approval. The bidder shall provide the manufacturer's approved US FDA Premarket Notification (PMN) or CE mark approval certificate. 8. The winning bidder shall provide current and valid calibration certificate for each equipment during delivery. 9. Certification that the supplier has the capability or authority for corrective and preventive maintenance of the unit. The certificate must be issued by the manufacturer to the bidder. 10. Certification from the manufacturer authenticated by the Philippine Consulate from the country of origin of the unit that the warranty should not be affected with a change of distributor.					

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3	**CL-03-2024**	**Supply, Delivery, Installation, Testing and Commissioning of a**	1	Clinical Laboratory Section	NCMH New Laboratory Building	National Center for Mental Health	a. One-time delivery within **30 days upon**

		Brand-New Hemoglobinometer **SPECIFICATIONS:** 1. Compact and lightweight 2. Portable 3. With digital display 4. Finger prick, spot testing analyzer with micro cuvette technology 5. Should have LED/LCD display of hemoglobin in g/l or g/dl 6. Instrument should be able to work in hot climate up to 45 degrees Celsius 7. The winning bidder shall provide startup kits for at least 50 tests **STANDARD REQUIREMENTS:** 1. The bidder must provide the current and valid certificate of authorized or exclusive distributorship. 2. The bidder must provide the current and valid certificate of Manufacturer's compliance with ISO 13485. 3. The bidder/supplier shall secure a certificate from the manufacturer stating that the equipment is brand new, unused and not a discontinued model or was listed in the market recall. 4. Bidder's certificate guaranteeing the availability on the supply of spare parts ten (10) years from the end of production. A certification must be issued by the manufacturer for the bidder. 5. Certification that the bidder must be in the local market for a minimum of ten (10) years. 6. Certification that the system or machine is not a retrofit solution.					**receipt of Notice to Deliver** b. Terms of payment at least 30 days from receipt of Sales invoice

		7. The machine should have a US FDA approval. The bidder shall provide the manufacturer's approved US FDA Premarket Notification (PMN) or CE mark approval certificate. 8. The winning bidder shall provide current and valid calibration certificate for each equipment during delivery. 9. Certification that the supplier has the capability or authority for corrective and preventive maintenance of the unit. The certificate must be issued by the manufacturer to the bidder. 10. Certification from the manufacturer authenticated by the Philippine Consulate from the country of origin of the unit that the warranty should not be affected with a change of distributor. 11. Warranty Certification: Certification that the bidder/supplier shall provide a three (3) year warranty for parts and services that includes corrective maintenance, preventive maintenance, and/or calibration. The warranty shall commence upon the acceptance of the end-user. 12. Bidder must provide preventive maintenance and/or calibration schedule within warranty period. 13. Certification that the supplier / bidder shall provide applications training for the users and maintenance personnel of the hospital. 14. Certification to provide manuals; Two (2) sets of service manual in English Language and					

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4	**CL-04-2024**	**Supply, Delivery, Installation, Testing and Commissioning of a Brand-New Agglutination viewer** **SPECIFICATIONS:** 1. Designed for use in blood typing, crossmatching, prothrombin time, kahn and other agglutination, flocculation, and serodiagnosis test. 2. Illuminates test tube contents through the use of an 8-watt bulb from above and a magnifying mirror from below. 3. Flexible neck allows for easy manipulation of light source. 4. Include Bulb, magnifying mirror, Flexible mirror, side mounted on/off switch, flexible neck, mainly operator. 5. Hertz : 50/60Hz 6. Voltage : 220v 7. Weight : 3kg 8. Amps : 36Ma	1	Clinical Laboratory Section	NCMH New Laboratory Building	National Center for Mental Health	a. One-time delivery within **30 days upon receipt of Notice to Deliver** b. Terms of payment at least 30 days from receipt of Sales invoice

		9. Temperature : 18C to 33C **STANDARD REQUIREMENTS:** 1. The bidder must provide the current and valid certificate of authorized or exclusive distributorship. 2. The bidder must provide the current and valid certificate of Manufacturer's compliance with ISO 13485. 3. The bidder/supplier shall secure a certificate from the manufacturer stating that the equipment is brand new, unused and not a discontinued model or was listed in the market recall. 4. Bidder's certificate guaranteeing the availability on the supply of spare parts ten (10) years from the end of production. A certification must be issued by the manufacturer for the bidder. 5. Certification that the bidder must be in the local market for a minimum of ten (10) years. 6. Certification that the system or machine is not a retrofit solution. 7. The machine should have a US FDA approval. The bidder shall provide the manufacturer's approved US FDA Premarket Notification (PMN) or CE mark approval certificate. 8. The winning bidder shall provide current and valid calibration certificate for each equipment during delivery. 9. Certification that the supplier has the capability or authority for corrective and preventive maintenance of the unit.					

		The certificate must be issued by the manufacturer to the bidder. 10. Certification from the manufacturer authenticated by the Philippine Consulate from the country of origin of the unit that the warranty should not be affected with a change of distributor. 11. Warranty Certification: Certification that the bidder/supplier shall provide a three (3) year warranty for parts and services that includes corrective maintenance, preventive maintenance, and/or calibration. The warranty shall commence upon the acceptance of the end-user. 12. Bidder must provide preventive maintenance and/or calibration schedule within warranty period. 13. Certification that the supplier / bidder shall provide applications training for the users and maintenance personnel of the hospital. 14. Certification to provide manuals; Two (2) sets of service manual in English Language and Two (2) sets of user manual in English Language upon delivery of the equipment. 15. Certification that the brand has been in the local market for at least five (5) years with at least five (5) current installations. 16. Certification that the supplier will be responsible for the notification, transportation to the site, delivery, installation and testing on the site (hospital / health facility) expenses for such will be					

		on the account of the supplier. 17. Certification that the supplier / bidder shall provide free installation of the equipment. 18. With 24/7 Technical Support. 19. Delivery period – 30 calendar days.					
5	**CL-05-2024**	**Supply, Delivery, Installation, Testing and Commissioning of a Brand-New Clinical Centrifuge, table top, 24 placer** **SPECIFICATIONS:** 1. 24 placer 2. Adjustable RPM from 500-4000 3. Max. RCF: 2500×g 4. Speed Accuracy: ±100rpm 5. Swing-out and fixed-angle rotor options 6. Lid lock protection 7. Brushless DC motor 8. Large LCD display 9. Safety Device: Automatic internal, Diagnosis 10. Power: Single phase, 100V-240V, 50Hz/60Hz **STANDARD REQUIREMENTS:** 1. The bidder must provide the current and valid certificate of authorized or exclusive distributorship. 2. The bidder must provide the current and valid certificate of Manufacturer's compliance with ISO 13485. 3. The bidder/supplier shall secure a certificate from the manufacturer stating that the equipment is brand new, unused and not a discontinued model or was listed in the market recall. 4. Bidder's certificate guaranteeing the availability on the supply of spare parts ten (10)	1	Clinical Laboratory Section	NCMH Modular Hospital II (Temporary Lab Building)	National Center for Mental Health	a. One-time delivery within **30 calendar days upon receipt of Notice to Deliver** b. Terms of payment at least 30 days from receipt of Sales invoice

		years from the end of production. A certification must be issued by the manufacturer for the bidder. 5. Certification that the bidder must be in the local market for a minimum of ten (10) years. 6. Certification that the system or machine is not a retrofit solution. 7. The machine should have a US FDA approval. The bidder shall provide the manufacturer's approved US FDA Premarket Notification (PMN) or CE mark approval certificate. 8. The winning bidder shall provide current and valid calibration certificate for each equipment during delivery. 9. Certification that the supplier has the capability or authority for corrective and preventive maintenance of the unit. The certificate must be issued by the manufacturer to the bidder. 10. Certification from the manufacturer authenticated by the Philippine Consulate from the country of origin of the unit that the warranty should not be affected with a change of distributor. 11. Warranty Certification: Certification that the bidder/supplier shall provide a three (3) year warranty for parts and services that includes corrective maintenance, preventive maintenance, and/or calibration. The warranty shall commence upon the acceptance of the end-user. 12. Bidder must provide preventive maintenance					

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6	**CL-06-2024**	**Supply, Delivery, Installation, Testing and Commissioning of a Brand-New Multiplex PCR system capable of panel testing** **SPECIFICATIONS:** 1. Molecular Multiplex PCR Machine 2. Capable of running Pneumonia, blood culture, meningitis, gastrointestinal, respiratory and joint infection panels Respiratory panel: Can detect Chlamydia pneumonia.	1	Clinical Laboratory Section	NCMH Modular Hospital II (Temporary Lab Building)	National Center for Mental Health	a. One-time delivery within **30 calendar days upon receipt of Notice to Deliver** b. Terms of payment at least 30 days from receipt of Sales invoice

		Meningitis Panel: Can detect Cytomegalovirus (CMV) Pneumonia Panel: Can run sputum/ETA/BAL samples, includes detection of AMR genes Blood Culture: Includes detection of yeast, includes detection of AMR genes. 3. Panels are FDA approved. 4. Integrates sample preparation, nucleic acid extraction, amplification, and detection in 60 minutes testing time. 5. Can detect bacteria, viruses, yeasts and antimicrobial resistant genes. 6. LIS ready **STANDARD REQUIREMENTS:** 1. The bidder must provide the current and valid certificate of authorized or exclusive distributorship. 2. The bidder must provide the current and valid certificate of Manufacturer's compliance with ISO 13485. 3. The bidder/supplier shall secure a certificate from the manufacturer stating that the equipment is brand new, unused and not a discontinued model or was listed in the market recall. 4. Bidder's certificate guaranteeing the availability on the supply of spare parts ten (10) years from the end of production. A certification must be issued by the manufacturer for the bidder. 5. Certification that the bidder must be in the local market for a minimum of ten (10) years.					

		6. Certification that the system or machine is not a retrofit solution. 7. The machine should have a US FDA approval. The bidder shall provide the manufacturer's approved US FDA Premarket Notification (PMN) or CE mark approval certificate. 8. The winning bidder shall provide current and valid calibration certificate for each equipment during delivery. 9. Certification that the supplier has the capability or authority for corrective and preventive maintenance of the unit. The certificate must be issued by the manufacturer to the bidder. 10. Certification from the manufacturer authenticated by the Philippine Consulate from the country of origin of the unit that the warranty should not be affected with a change of distributor. 11. Warranty Certification: Certification that the bidder/supplier shall provide a three (3) year warranty for parts and services that includes corrective maintenance, preventive maintenance, and/or calibration. The warranty shall commence upon the acceptance of the end-user. 12. Bidder must provide preventive maintenance and/or calibration schedule within warranty period. 13. Certification that the supplier / bidder shall provide applications training for the users and maintenance personnel of the hospital.					

		14. Certification to provide manuals; Two (2) sets of service manual in English Language and Two (2) sets of user manual in English Language upon delivery of the equipment. 15. Certification that the brand has been in the local market for at least five (5) years with at least five (5) current installations. 16. Certification that the supplier will be responsible for the notification, transportation to the site, delivery, installation and testing on the site (hospital / health facility) expenses for such will be on the account of the supplier. 17. Certification that the supplier / bidder shall provide free installation of the equipment. 18. With 24/7 Technical Support. 19. Delivery period – 30 calendar days.					
7	**CL-07-20241**	**Supply, Delivery, Installation, Testing and Commissioning of a Brand-New Cartridge Based PCR Analyzer** **SPECIFICATIONS:** 1. Cartridge Based Nucleic Acid Amplification Test 2. Real- Time PCR technology integrating sample preparation, amplification and detection 3. Small foot-print and can be used as point-of-care equipment 4. Detect an expanded number of genetic targets (microbes, antimicrobial resistance genes, disease loci, oncogenes) and capable to run and simultaneously detects MTB complex and extensive drug resistance associated mutations,	1	Clinical Laboratory Section	NCMH Modular Hospital II (Temporary Lab Building)	National Center for Mental Health	a. One-time delivery within **30 calendar days upon receipt of Notice to Deliver** b. Terms of payment at least 30 days from receipt of Sales invoice

		HIV viral load, HPV Genotype, CT/NG, BCR-ABL, MRSA, Sars CoV-2, Flu, etc. 5. Unit/System with 4 indepently controlled reaction sites 6. Solid state heater and forces-air cooling at each site 7. Interoperability with existing cartridge-based PCR machine 8. Reaction chamber thermistors calibrated using National Institute of Standards and Technology (NIST) traceable standards 9. With Desktop/Computer System 10. With Barcode Scanner for Cartridge and/or Sample ID 11.Power supply: 100-240 V (Auto volt): 50-60Hz: Single phase 12. With UPS compatible with the unit **STANDARD REQUIREMENTS:** 1. The bidder must provide the current and valid certificate of authorized or exclusive distributorship. 2. The bidder must provide the current and valid certificate of Manufacturer's compliance with ISO 13485. 3. The bidder/supplier shall secure a certificate from the manufacturer stating that the equipment is brand new, unused and not a discontinued model or was listed in the market recall. 4. Bidder's certificate guaranteeing the availability on the supply of spare parts ten (10) years from the end of production. A certification must be issued by the					

		manufacturer for the bidder. 5. Certification that the bidder must be in the local market for a minimum of ten (10) years. 6. Certification that the system or machine is not a retrofit solution. 7. The machine should have a US FDA approval. The bidder shall provide the manufacturer's approved US FDA Premarket Notification (PMN) or CE mark approval certificate. 8. The winning bidder shall provide current and valid calibration certificate for each equipment during delivery. 9. Certification that the supplier has the capability or authority for corrective and preventive maintenance of the unit. The certificate must be issued by the manufacturer to the bidder. 10. Certification from the manufacturer authenticated by the Philippine Consulate from the country of origin of the unit that the warranty should not be affected with a change of distributor. 11. Warranty Certification: Certification that the bidder/supplier shall provide a three (3) year warranty for parts and services that includes corrective maintenance, preventive maintenance, and/or calibration. The warranty shall commence upon the acceptance of the end-user. 12. Bidder must provide preventive maintenance and/or calibration schedule within warranty period.					

		13. Certification that the supplier / bidder shall provide applications training for the users and maintenance personnel of the hospital. 14. Certification to provide manuals; Two (2) sets of service manual in English Language and Two (2) sets of user manual in English Language upon delivery of the equipment. 15. Certification that the brand has been in the local market for at least five (5) years with at least five (5) current installations. 16. Certification that the supplier will be responsible for the notification, transportation to the site, delivery, installation and testing on the site (hospital / health facility) expenses for such will be on the account of the supplier. 17. Certification that the supplier / bidder shall provide free installation of the equipment. 18. With 24/7 Technical Support. 19. Delivery period – 30 calendar days.					
8	**CL-08-2024**	**Supply, Delivery, Installation, Testing and Commissioning of a Brand-New Incubator, Microbiological/General Incubator** **SPECIFICATIONS:** 1. At least 150 liters capacity 2. Natural convection air flow 3. Digital control display of temperature and time 4. Adjustable temperature from ambient +5C to at least 60C 5. Tempered safety glass door 6. Touch button operating panel	1	Clinical Laboratory Section	NCMH Modular Hospital II (Temporary Lab Building)	National Center for Mental Health	a. One-time delivery within **30 calendar days upon receipt of Notice to Deliver** b. Terms of payment at least 30 days from receipt of Sales invoice

		7. Stainless steel chamber and shelves material 8. At least 2 shelves 9. Alarms and self diagnostic function 10. Power: Single phase, 100V-240V, 50Hz/60Hz 11. With 2KVA Automatic Voltage Regulator **STANDARD REQUIREMENTS:** 1. The bidder must provide the current and valid certificate of authorized or exclusive distributorship. 2. The bidder must provide the current and valid certificate of Manufacturer's compliance with ISO 13485. 3. The bidder/supplier shall secure a certificate from the manufacturer stating that the equipment is brand new, unused and not a discontinued model or was listed in the market recall. 4. Bidder's certificate guaranteeing the availability on the supply of spare parts ten (10) years from the end of production. A certification must be issued by the manufacturer for the bidder. 5. Certification that the bidder must be in the local market for a minimum of ten (10) years. 6. Certification that the system or machine is not a retrofit solution. 7. The machine should have a US FDA approval. The bidder shall provide the manufacturer's approved US FDA Premarket Notification (PMN) or CE mark approval certificate. 8. The winning bidder shall provide current and valid calibration					

		certificate for each equipment during delivery. 9. Certification that the supplier has the capability or authority for corrective and preventive maintenance of the unit. The certificate must be issued by the manufacturer to the bidder. 10. Certification from the manufacturer authenticated by the Philippine Consulate from the country of origin of the unit that the warranty should not be affected with a change of distributor. 11. Warranty Certification: Certification that the bidder/supplier shall provide a three (3) year warranty for parts and services that includes corrective maintenance, preventive maintenance, and/or calibration. The warranty shall commence upon the acceptance of the end-user. 12. Bidder must provide preventive maintenance and/or calibration schedule within warranty period. 13. Certification that the supplier / bidder shall provide applications training for the users and maintenance personnel of the hospital. 14. Certification to provide manuals; Two (2) sets of service manual in English Language and Two (2) sets of user manual in English Language upon delivery of the equipment. 15. Certification that the brand has been in the local market for at least five (5) years with at least five (5) current installations. 16. Certification that the supplier will be					

		responsible for the notification, transportation to the site, delivery, installation and testing on the site (hospital / health facility) expenses for such will be on the account of the supplier. 17. Certification that the supplier / bidder shall provide free installation of the equipment. 18. With 24/7 Technical Support. 19. Delivery period – 30 calendar days.					
9	**CL-09-2024**	**Supply, Delivery, Installation, Testing and Commissioning of a Brand-New Incubator, CO2** **SPECIFICATIONS:** 1. At least 50 liters capacity 2. LCD Touch Panel Controller/ Touchscreen controller (deliver full control over different protocols) 3. Integrated tray catches to accommodate more culture containers 4 Self- diagnostic system for errors with visual and audible alarms 5. Notification and alarm Functions: • Temperature setpoint deviations • CO2 concentration setpoint deviations • Other deviations outside normal machine operations 6. With temperature safety device 7. With chamber controller for controls of the Temperature in °C and Carbon Dioxide concentration 9. Stainless steel chamber and shelves material 10. Adjustable temperature from ambient +5C to at least 60C	1	Clinical Laboratory Section	NCMH Modular Hospital II (Temporary Lab Building)	National Center for Mental Health	a. One-time delivery within **30 calendar days upon receipt of Notice to Deliver** b. Terms of payment at least 30 days from receipt of Sales invoice

		11. Temperature control uniformity - as per manufacturer's standard 12. CO2 control range and deviation - 0 % to 20% 13. Vertical/natural air flow 14. Automatic decontamination protocol 15. Must come with CO2 tank filled, regulator and tubing 16. Power: Single phase, 100V-240V, 50Hz/60Hz 17. With 2KVA Automatic Voltage Regulator **STANDARD REQUIREMENTS:** 1. The bidder must provide the current and valid certificate of authorized or exclusive distributorship. 2. The bidder must provide the current and valid certificate of Manufacturer's compliance with ISO 13485. 3. The bidder/supplier shall secure a certificate from the manufacturer stating that the equipment is brand new, unused and not a discontinued model or was listed in the market recall. 4. Bidder's certificate guaranteeing the availability on the supply of spare parts ten (10) years from the end of production. A certification must be issued by the manufacturer for the bidder. 5. Certification that the bidder must be in the local market for a minimum of ten (10) years. 6. Certification that the system or machine is not a retrofit solution. 7. The machine should have a US FDA approval. The bidder shall provide the manufacturer's					

		approved US FDA Premarket Notification (PMN) or CE mark approval certificate. 8. The winning bidder shall provide current and valid calibration certificate for each equipment during delivery. 9. Certification that the supplier has the capability or authority for corrective and preventive maintenance of the unit. The certificate must be issued by the manufacturer to the bidder. 10. Certification from the manufacturer authenticated by the Philippine Consulate from the country of origin of the unit that the warranty should not be affected with a change of distributor. 11. Warranty Certification: Certification that the bidder/supplier shall provide a three (3) year warranty for parts and services that includes corrective maintenance, preventive maintenance, and/or calibration. The warranty shall commence upon the acceptance of the end-user. 12. Bidder must provide preventive maintenance and/or calibration schedule within warranty period. 13. Certification that the supplier / bidder shall provide applications training for the users and maintenance personnel of the hospital. 14. Certification to provide manuals; Two (2) sets of service manual in English Language and Two (2) sets of user manual in English Language upon delivery of the equipment.					

		15. Certification that the brand has been in the local market for at least five (5) years with at least five (5) current installations. 16. Certification that the supplier will be responsible for the notification, transportation to the site, delivery, installation and testing on the site (hospital / health facility) expenses for such will be on the account of the supplier. 17. Certification that the supplier / bidder shall provide free installation of the equipment. 18. With 24/7 Technical Support. 19. Delivery period – 30 calendar days.					
CONFORME:							
(Company Name)							
(Name and Signature of Authorized Representative)							