

## REPUBLIC OF THE PHILIPPINES Department of Health

## NATIONAL CENTER FOR MENTAL HEALTH

Nueve de Febrero Street, Mandaluyong City, Philippines



## **BIDS AND AWARDS COMMITTEE**

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## 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.

					<b>Delivery Site</b>	2	Delivery	
Item No.	Description		Qty	OFFICE	FACILITY	ADDRESS	Period and Terms of Payment	
1	CL- 01- 2024	Supply, Delivery, Installation, Testing Commissioning of a Brand-New Ultra Low	1	Clinical Laboratory Section	NCMH New Laboratory Building	National Center for Mental Health	a. One-time delivery within 30 calendar	
		Freezer, -80 C  SPECIFICATIONS: 1. Capacity: at least 18 cubic feet.					days upon receipt of Notice to Deliver	
		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.					b. Terms of payment at least 30 days from receipt of Sales invoice	
		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						

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			1	
L I				
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. Cértification 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		i i		
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				
I mapping Consultate from			1/	

			i.
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)		1	
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		Ý	
supplier / bidder shall			
provide free installation			
of the equipment.			
18. With 24/7 Technical			
Support.			1

19. Delivery period -30					
calendar days.					
	1	Clinical	NCMH New	National	0.0004
Supply, Delivery,	1	Clinical			a. One-time
Installation, Testing and		Laboratory	Laboratory	Center for	delivery
Commissioning of a		Section	Building	Mental	within 30
Brand-New Blood Bank				Health	calendar
Refrigerator 120 blood					days upon
bag capacity					receipt of
					l
SPECIFICATIONS:				0	
1. Alarm and Safety					Deliver
Function.					
-Audible and flashing					b. Terms of
LED visual alarms.					payment at
- Door alarm and key lock					least 30 days
are standard feature.					from receipt
- Built in temperature					of Sales
recorder.					1
2. Stable temperature					invoice
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					
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				1
	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				1
	issued by the				
	manufacturer for the				
	bidder.				
	5. Certification that the				1
	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.		1		
	10. Certification from the				
	manufacturer				
	manutaciulti				
B 0	authenticated by the	1	1		

<u></u>				
	Philippine Consulate from			
	the country of origin of			
	the unit that the warranty			
	should not be affected			
	with a change of			
	distributor.			
	11. Warranty			1
	Certification:			1
	Certification that the			1
	bidder/supplier shall			1
	provide a three (3) year			
	warranty for parts and			
	services that includes			
	corrective maintenance,			1
	preventive maintenance,			
	and/or calibration. The			
	warranty shall commence			
	upon the acceptance of the			
	end-user.			
	12. Bidder must provide			
	preventive maintenance	1		
	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.			
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19. Delivery period – 30					
calendar days.  Supply, Delivery, Installation, Testing and Commissioning of a Brand-New Plasma Thawer	1	Clinical Laboratory Section	NCMH New Laboratory Building	National Center for Mental Health	a. One-time delivery within 30 calendar days upon
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 sensorsno risk of contamination or infection					receipt of Notice to Deliver  b. Terms of payment at least 30 days from receipt of Sales invoice
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.					
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		D.	
	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			1
	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.	1		
	7. The machine should			
	have a US FDA approval.			
	The bidder shall provide			1
	the manufacturer's			
1 1 11	approved US FDA			
	Premarket Notification			
	(PMN) or CE mark			
	approval certificate.			
	8. The winning bidder			
1 1	shall provide current and valid calibration			
1 1 1				
	equipment during			
	delivery.			
	9. Certification that the			
	supplier has the capability			
1 1	or authority for corrective			
	and preventive			
	maintenance of the unit.			
1 1 1	The certificate must be			
1 1 1	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			
1 1 1	distributor.			
	11. Warranty			
	Certification:			
	Certification that the			
	biddet/supplier shall			
	provide a three (3) year			
	warranty for parts and			
	services that includes			
	corrective maintenance,			
	preventive maintenance,			
	and/or calibration. The			
1 1 1	warranty shall commence			
				00

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,	1	Clinical	NCMH_New_	National	a. One-time
Installation, Testing and Commissioning of a		Laboratory	Laboratory Building	Center for Mental	delivery
Brand-New Platelet		Section	Dullullig	Health	within 30
Agitator with incubator					calendar days upon
CDE CIPIC LEICE					receipt of
SPECIFICATIONS: 1. Refrigeration &					Notice to
precision pulse heating					Deliver
for achieving 22°C (±					
1°C)					b. Terms of
2. Forced air circulation for temperature					payment at
uniformity					least 30 days from receipt
·					dom receipt

V					
	3. Powder coated exterior			of Sales	
	with high-grade stainless-			invoice	
	steel interior			, 0,00	
	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				
	Tor 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		1		
	guaranteeing the				
	availability on the supply				
	of spare parts ten (10)				
	years from the end of				
	V	\X			

	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	1 1			
	ten (10) years.				
	6. Certification that the				
	system or machine is not a	1 1			
	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	1			
1 1	issued by the				
	manufacturer to the				
	bidder.				
	10. Certification from the				
	manufacturer				
	authenticated by the		1		
	Philippine Consulate from				
	the country of origin of		1		
	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				ly .
	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		1		
	and/or calibration				
	and/or calibration			1	

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schedule within warranty					
period.  13. Certification that the					
				1	
supplier / bidder shall					
provide applications training for the users and				0	
maintenance personnel of					
the hospital.					1
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,	1	Clinical	NCMH New	National	a. One-time
Installation, Testing and		Laboratory	Laboratory Building	Center for- Mental	delitely
Commissioning of a Brand-New Blood Bag		Section	Building	Health	within 30
Mixer				. 1041011	calendar
					days upon
SPECIFICATIONS:					receipt of
1. Linear and 3d mixing					Notice to
with central support					Deliver
2. Weighing range: 0-					1 70
990ml.					b. Terms of
3. Alarm type: Automatic					payment at
low and high flow alarm,					least 30 days
adjustable. 4. Adjustable maximum					from receipt
donation time: Up to 20					of Sales
minutes.					invoice
5. Protocols: Up to 20					
different working					
programs.					
different working					

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	6. Built-in memory: More					
	than 1000 collection					
	procedure data					
	7. Automatic clamping					
	module					
	module					
	CT LVD LDD					
	STANDARD					
	REQUIREMENTS:					
	1. The bidder must					1
	provide the current and					
	valid certificate of					1
	authorized or exclusive			)		
	distributorship.			ľ		
	2. The bidder must					
						1
	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		i i			
	guaranteeing the					
	availability on the supply					
	of spare parts ten (10)					
	years from the end of					
	production. A					
	certification must be					4
	issued by the					
	manufacturer for the				1	
	bidder.					
	5. Certification that the					
	bidder must be in the local					
	market for a minimum of					
	ten (10) years.					
<u> </u>	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	1				
	supplier has the capability					
						1

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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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	)	C.		
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	- 1			
supplier will be				
responsible for the				
notification,				
transportation to the site,				ľ
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.  ITEMS NO. 1 to 5 UNDER extraction chair for gener with the following specific Blood extraction chair: 1. Stainless steel tubular fra 2. With built in stainless dra 3. Upholstered cushion (Pol 4. Adjustable armrest (swin  Blood bag tube stripper: 1. Spring loaded handle 2. Easy grip handle 3. Capable of sealing clip and	al laborations:  me awer w lyureth g-out)	ith at least one ane material)	otomy use and 2		
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	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
		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
	3485.
	The bidder/supplier
	hall secure a certificate
	rom the manufacturer
	tating that the equipment
	s brand new, unused and
	oot a discontinued model
	or was listed in the market
	ecall.
	Bidder's certificate
	guaranteeing the
	vailability on the supply
	of spare parts ten (10)
	ears from the end of
	ertification must be
	ssued by the
	nanufacturer for the
	idder.
	. Certification that the
1 1 1	idder must be in the local
1 1 1	narket for a minimum of
	en (10) years.
	. Certification that the
	ystem or machine is not a
	etrofit solution.
1 1	. The machine should
	ave a US FDA approval.
	The bidder shall provide
t	ne manufacturer's
8	pproved US FDA
I	remarket Notification
	PMN) or CE mark
	pproval certificate.
	. The winning bidder
	hall provide current and
	alid calibration
	ertificate for each
	quipment during
	elivery.
	. Certification that the
	upplier has the capability
	r authority for corrective
	nd preventive
	naintenance of the unit.
	The certificate must be
	ssued by the
	nanufacturer to the
1 1 1	idder.
	0. Certification from the
	o. Certification from the
1 1 1	
	hilippine Consulate from
	ne country of origin of
	ne unit that the warranty
	hould not be affected
L L I	rith a change of
	istributor.
	istributor.

J	03- 2024	Installation, Testing and Commissioning of a	1	Laboratory Section	Laboratory Building	Center for Mental Health	delivery within 30 days upon
3	CL-	calendar days.  Supply, Delivery,	1	Clinical	NCMH New	National	a. One-time
		Support.  19. Delivery period – 30					
		of the equipment.  18. With 24/7 Technical					
		supplier / bidder shall provide free installation					
		17. Certification that the					
		on the account of the supplier.					
		expenses for such will be					
		testing on the site (hospital / health facility)					
		delivery, installation and					
		notification, transportation to the site,					
		responsible for the					
		16. Certification that the supplier will be					
		current installations.					
		market for at least five (5) years with at least five (5)					
		brand has been in the local					
		15. Certification that the					
		Language upon delivery of the equipment.					
		manual in English					
		Two (2) sets of user					
		sets of service manual in English Language and					
		provide manuals; Two (2)					
		the hospital.  14. Certification to					
		maintenance personnel of					
		training for the users and					
		supplier / bidder shall provide applications					
		13. Certification that the					
		period.					
		and/or calibration schedule within warranty					
		preventive maintenance					
		12. Bidder must provide					
		upon the acceptance of the end-user.					
		warranty shall commence					
		and/or calibration. The					
		corrective maintenance, preventive maintenance,					
		services that includes					
		warranty for parts and					
		bidder/supplier shall provide a three (3) year					
		Certification that the					
		Certification:					

Brand-New Hemoglobinometer  SPECIFICATIONS:  1. Compact and lightweight 2. Portable	receipt of Notice to Deliver  b. Terms of payment at least 30 days
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	from receipt of Sales invoice
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	Ŋ				]
l h	ive a US FDA approval.					
	ne bidder shall provide					
th						
ap	proved US FDA					
Pi	emarket Notification					
(F	MN) or CE mark					
	proval certificate.					
	The winning bidder					
sh	all provide current and					
l l va	lid calibration					
	rtificate for each					
	livery.					1
9.	Certification that the					
su	pplier has the capability					
	authority for corrective					
ar						
				,		
	aintenance of the unit.					
T T	ne certificate must be					
iss	sued by the					
m	anufacturer to the					
I I I	dder.					
	Certification from the					1
1 1						
	anufacturer					
au	thenticated by the					
Pl	ilippine Consulate from					
	e country of origin of					1
	e unit that the warranty					1
	ould not be affected					
	th a change of					
di	stributor.					
11	. · Warranty					
	ertification:					
1. 1. 1	ertification that the					
			(1			
	dder/supplier shall					
	ovide a three (3) year					
w	arranty for parts and					
se	rvices that includes					ĺ
II II	rrective maintenance,		l l			
	eventive maintenance,					
				1		
	d/or calibration. The					
	irranty shall commence	-				-
	on the acceptance of the					
en	d-user.					
12	. Bidder must provide					
	eventive maintenance					
	d/or calibration					
					ĺ	
E E	nedule within warranty					
	riod.			4		
13	. Certification that the					
su	oplier / bidder shall		1			
	ovide applications	1				
	ining for the users and					
				1		
	intenance personnel of		li li		(	
the	hospital.		9	1	Al .	
14	. Certification to					
to the state of th		1	i i		a di	
pr	ovide manuals: Two (2)		. 11		1,	V.
	ovide manuals; Two (2)					
se	ovide manuals; Two (2) s of service manual in glish Language and					

4	CL- 04- 2024	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 Agglutination viewer  SPECIFICATIONS:  1. Designed for use in blood, typing, crossmatching, prothrombin time, kahn and other agglutination,	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
		flocculation, and serodiagnosis test.					from receipt of Sales
		2. Illuminates test tube					invoice
		of an 8-watt bulb from					- Maria Maria Maria
		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,					

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			3	
(PMN) or CE mark				
approval certificate.				
8. The winning bidder				
shall provide current and				
valid calibration				
certificate for each				
equipment during	1			
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				1
bidder.				
10. Certification from the				1
manufacturer				
authenticated by the				
Philippine Consulate from				
the country of origin of				
the unit that the warranty				1
should not be affected				
with a change of				
distributor.				
11. Warranty				1
Certification:				
Certification that the		1		l)
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			1	
and/or calibration				
schedule within warranty				
period.				
13. Certification that the				
supplier / bidder shall				
provide applications				
training for the users and	8			
maintenance personnel of		-		
the hospital.		1		
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			J	
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				
* *				
responsible for the				
notification,				
transportation to the site,				
delivery, installation and				
testing on the site				
(hospital / health facility)				
expenses for such will be				

	C	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.			NGMI	National	
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.	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
		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)					

		III.		
	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			1
	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	1		
	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	j i		
	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: Certification that the	į.		
	bidder/supplier shall			
	provide a three (3) year	le l		
	warranty for parts and	Y .		
	services that includes		-	
	corrective maintenance,			
	preventive maintenance,			
	and/or calibration. The			
	warranty shall commence		1	
	upon the acceptance of the			
	end-user.			
	12. Bidder must provide			
	preventive maintenance			
A				

		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)					
1		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			c		
	li .	testing on the site					
		(hospital / health facility)				l,	
		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					
-	CL-	calendar days.  Supply, Delivery,	9	Clinical	NCMH	National	a. One-time
6	06-	Supply, Delivery, Installation, Testing and	1	ACTIVITY OF THE PARTY OF	Modular_	Center for	The second secon
	2024	Commissioning of a		Laboratory	Hospital II	Mental	delivery
	2027	Brand-New Multiplex		Section	(Temporary	Health	within 30 calendar
		PCR system capable of			Lab		
		panel testing			Building)		days upon
					J.		receipt of
		SPECIFICATIONS:					Notice to
		1. Molecular Multiplex					Deliver
		PCR Machine					
		2. Capable of running					b. Terms of
		Pneumonia, blood culture,					payment at
		meningitis,					least 30 days
		gastrointestinal,					from receipt
		respiratory and joint					of Sales
		infection panels					invoice
		Respiratory panel: Can detect Chlamydia					
		pneumonia.			,		
		pheumoma.					

	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
M 10.	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.
	ion (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.	
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(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.	
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.	
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or authority for corrective and preventive maintenance of the unit.  The certificate must be issued by the manufacturer to the bidder.	
and preventive maintenance of the unit. The certificate must be issued by the manufacturer to the bidder.	
maintenance of the unit. The certificate must be issued by the manufacturer to the bidder.	
The certificate must be issued by the manufacturer to the bidder.	
issued by the manufacturer to the bidder.	
manufacturer to the bidder.	
bidder.	
10. Certification from the	
manufacturer	1
authenticated by the	
Philippine Consulate from	
the country of origin of	
the unit that the warranty	
should not be affected	1
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) 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					
		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					
		supplier. 17. Certification that the supplier / bidder shall					
		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	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
		Amplification Test 2. Real- Time PCR					b. Terms of
		technology integrating sample preparation, amplification and					payment at least 30 days from receipt
		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,					of Sales invoice

7				
	HIV viral load, HPV			
	Genotype, CT/NG, BCR-			
	ABL, MRSA, Sars CoV-			
	2, Flu, etc.			Ŋ.
	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			
	1			
	8. Reaction chamber			
	thermistors calibrated			
	using National Institute of			
	Standards and			
	Technology (NIST)			
	traceable standards			
	9. With			
	Desktop/Computer			
	System System			
			j'	
	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			
	1 4			
	valid certificate of			
	authorized or exclusive			
	distributorship.			
	2. The bidder must			
	provide the current and			
	valid certificate of			
	Manufacturer's		1	
	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	V		
	I'			
	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	
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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11. Warranty	$\perp$
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.	

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		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.  17. Certification that the supplier.  18. With 24/7 Technical Support.  19. Delivery period — 30					
8	CL- 08- 2024	calendar days.  Supply, Delivery, Installation, Testing and Commissioning of a	1	Clinical Laboratory Section	NCMH Modular Hospital II	National Center for Mental	a. One-time delivery within 30
		Brand-New Incubator, Microbiological/General			(Temporary Lab	Health	calendar
		Incubator			Building)		days upon
							receipt of
		SPECIFICATIONS:					Notice to Deliver
		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					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					li
Regulator					
8					
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				1	
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.		_			
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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	(		
1	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.			1
	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	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
		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					b. Terms of payment at least 30 days from receipt of Sales invoice
		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					

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	
l .	
Regulator	
STANDARD	
REQUIREMENTS:	1 1
1. The bidder must	
provide the current and	1
valid certificate of	
authorized or exclusive	
distributorship.	l,
2. The bidder must	
provide the current and	
Manufacturer's	
compliance with ISO	
13485.	
3. The bidder/supplier	
shall secure a certificate	1
from the manufacturer	
stating that the equipment	
is brand new, unused and	
not a discontinued model	
or was listed in the market	
recall.	
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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	
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			l <sup>1</sup>	
delivery.				
9. Certification that the				
supplier has the capability				
or authority for corrective				
and preventive				
maintenance of the unit.		8		
The certificate must be				
issued by the				
manufacturer to the				
bidder.				
10. Certification from the				
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manufacturer				
authenticated by the				
Philippine Consulate from				
the country of origin of				1
the unit that the warranty				
should not be affected				
with a change of				
distributor.				
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Certification:				
Certification that the				
bidder/supplier shall				
provide a three (3) year				
warranty for parts and				
services that includes				
corrective maintenance,				
preventive maintenance,				
				1
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			1	
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	9			
manual in English		1		
Language upon delivery		i i		
of the equipment.				