All Bidders

Amendment -II

Date: 12.07.2019

Project: Supply, Installation, Testing & Commissioning of Medical Gas Manifold System for Hospital Block at All India Institute of Medical Sciences (AIIMS), Nagpur

IFB No.HSCC/SES/MGMS/AIIMS/Nagpur/2019 Date:13.06.2019

This has reference to above IFB.

The following Amendment may be noted which shall be treated as part of the tender document and to be submitted duly signed & stamp along with tender.

Sr. No.	Bidders' Queries	Reply
1	Vol 4 - Technical Specifications, Page 2, Para 1.2 Oxygen Manifold Supply System (without Cylinders): Oxygen Manifold should consist of 2 row/s of respective numbers of class D-type bulk oxygen cylinders.	Oxygen Manifold should consist of 2/1 row/s of respective numbers of class D-type bulk oxygen cylinders.
	The Standard configuration of Manifolds available with most of the leading manufacturers are of Single (1) Row. Therefore we request you to amend the sentence as; Oxygen Manifold should consist of 2/1 row/s of respective numbers of class D-type bulk oxygen cylinders. In addition to this, and for enhancing safety of the manifold system, we suggest that you should demand that the manifold system should not contain any Halogenated Polymer materials.	Manifold system should not contain any Halogenated Polymer materials
2	Vol 4 - Technical Specifications, Page 2, Para 1.2 Oxygen Manifold Supply System (without Cylinders): Header bar/s assembly shall be provided with a highpressure shut off valve. It may be noted that HTM or ISO Standards doesnot permit the use of Shut Off Valves on the Headers. Moreover, if any Shut Off Valve are provided on Manifold Header, thenthe same will not be compliant / satisfying the requirement of HTM /ISO Standards. Hence, we request you to delete the sentence:	Header bar/s assembly shall be as per standards mentioned in the technical specification of tender.

	Header bar/s assemblyshall be provided with a	
	high pressure shut off valve.	
3	Vol 4 - Technical Specifications, Page 2, Para 1.2 Oxygen Manifold Supply System (without Cylinders): The manifold should be hydraulically tested to 3500psig or as per guideline of standard to be followed. In India, the general practise is that the medical	Manifolds are designed and tested with at leastinlet pressure of 3000 psig or as per standards mentioned in the technical specification of tender.
	gas manifolds are tested at a minimum pressure of 3500 psig. This is inherent and can be verified in many of the earlier tenders floated by HSCC as well. Accepting of any manifold which are tested	
	below 3500 psig isactually diluting the safety standards and requirement, and therebyincreasing the risk of untoward incidents happening. We therefore request you to maintain the requirement of manifold hydraulically tested to	
	minimum 3500 psig, and delete the wordings or as per guideline of standard to be followed.	
4	Vol 4 - Technical Specifications, Page 3, Para 1.3	Header bar/s assembly shall be as per standards mentioned in the technical
	Emergency Oxygen Manifold (without Cylinders): Header bar/s assembly shall be provided with a highpressure shut off valve.	specification of tender.
	It may be noted that HTM or ISO Standards doesnot permit the use of Shut Off Valves on the Headers.	
	Moreover, if any Shut Off Valve are provided on Manifold Header, thenthe same will not be compliant / satisfying the requirement of HTM / ISO Standards.	
	Hence, we request you to delete the sentence: Header bar/s assemblyshall be provided with a high pressure shut off valve.	
5	Vol 4 - Technical Specifications, Page 3, Para 1.3	Manifolds are designed and tested with at leastinlet pressure of 3000 psig or as
	Emergency Oxygen Manifold (without Cylinders): The manifold should be hydraulically tested to 3500psig or as per standard to be followed.	per standards mentioned in the technical specification of tender.
	In India, the general practise is that the medical gas manifolds are tested at a minimum pressure of 3500 psig. This is inherent and can be	

	Verified in many of the earlier tenders floated by HSCC as well.	
	Accepting of any manifold which are tested	
	below 3500 psig is	
	actually diluting the safety standards and	
	requirement, and thereby	
	increasing the risk of untoward incidents	
	happening.	
	We therefore request you to maintain the	
	requirement of manifold hydraulically tested to minimum 3500 psig, and delete the wordingsor as	
	per guideline of standard to be followed.	
6	Vol 4 - Technical Specifications,	Tender terms and condition
	Page 3, Para 1.3	prevails
	Emergency Oxygen Manifold (without	
	Cylinders):	
	Please confirm if any Pr Regulating arrangement	
	is required to beoffered along with the	
	Emergency Oxygen Manifold?	
	If YES, what is the flow capacity & other	
	technical requirement?	
7	Vol 4 - Technical Specifications,	BIS/US FDA/European CE
	Page 3, Para 1.5	Certified with 4 digit
	Overson Flour motor with Humidifion Pottle.	notified body number or American ETL/ UL listed
	Oxygen Flow meter with Humidifier Bottle: I) should be BIS/CE certified/ UL Listed	American ETL/ OL fisted
	should be bis/ell certified/ ell bisted	
	It is observed that elsewhere in the tender	
	technical specs, it is mentioned as: It should be	
	US FDA/European CE Certified with 4	
	digitnotified body number or American ETL/ UL	
	listed.	
	In this case the 4 Digit notified body number is missing?	
	Even for a less critical item like Ward Vacuum	
	Unit, you have asked for	
	European CE with 4 Digit notified body number.	
	Considering that the Oxygen Flowmeter is a	
	critical item and workingunder positive pressure,	
	we request that this also be classified as: It	
	should be US FDA/European CE Certified with 4 digit notified bodynumber or American ETL/ UL	
	listed.	
	Out of all the items sought in tender, only for this	
	item BIS Standard isaccepted. Why? It should be	
	ideally sought for all items or for none.We	

	Vol 4 - Technical Specifications,	Specification of Fully
	Page 1, Para 1.1	Automatic Oxygen Control
		Panel should be as per
	Fully Automatic Oxygen Control Panel	standards mentioned in the technical specification.
	This specification is tilted towards NFPA	or and a second
	Standard.	
	It is not possible to meet word by word	
	requirement in toto of Technical Specs, as this is	
	make and standard specific. We request you to	
	permit us to offer as per should fully meet and	
	complies with ISO 7396-1/ HTM0201 /	
	NFPA99C standards, and as per manufacturers	
	own design for a capacity of 2000 LPM at 50 / 60 PSI.	
8	Vol 4 - Technical Specifications,	Specification ofFully
	Page 9, Para 2.1	Automatic Nitrous Oxide
	Fully Automatic Nitrous Oxide Control Panel	Control Panel should be as
	This specification is tilted towards NFPA	per standards mentioned in
	Standard.	the technical specification.
	It is not possible to meet word by word	
	requirement in toto of Technical Specs, as this is	
	make and standard specific. We request you to	
	permit us to offer as per should fully meet and complies with ISO7396-1/ HTM0201 /	
	complies with ISO7396-1/ HTM0201 / NFPA99C standards, and as per manufacturers	
	own design for a capacity of 2000 LPM at 50 / 60	
	PSI.	
9	Vol 4 - Technical Specifications,	Header bar/s assembly shall
	Page 10, Para 2.2	be as per standards
	Nitrous Oxide Manifold (Without Cylinders)	mentioned in the technical
	Header bar/s assembly shall be provided with a	specification of tender.
	high pressure shut off valve.	
	It may be noted that HTM or ISO Standards does	
	not permit the use of Shut Off Valves on the Headers.	
	Moreover, if any Shut Off Valve are provided on	
	Manifold Header, then the same will not be	
	compliant / satisfying the requirement of HTM	
	/ISO Standards. Hence, we request you to delete	
	the sentence: Header bar/s assembly shall be	
	provided with a high pressure shut off valve.	
10	Vol 4 - Technical Specifications,	Manifolds are designed and
	Page 10, Para 2.2	tested with at least inlet
	Nitrous Oxide Manifold (Without Cylinders)	pressure of 3000 psig or as
	The manifold should be hydraulically tested to	per standards mentioned in
	3500 psig or as per guideline of standard.	the technical specification of
	In India, the general practise is that the medical	tender.
	gas manifolds are tested at a minimum pressure	
	of 3500 psig. This is inherent and can be verified	1

	in many of the earlier tenders floated by HSCC as well.	
	Accepting of any manifold which are tested	
	below 3500 psig is actually diluting the safety	
	standards and requirement, and thereby	
	increasing the risk of untoward incidents	
	happening.	
	We therefore request you to maintain the	
	requirement of manifold hydraulically tested to	
	minimum 3500 psig, and delete the wordings or	
1.1	as per guideline of standard to be followed.	T. 1 11 1
11	Vol 4 - Technical Specifications,	It shall be as per standards
	Page 10, Para 2.2	mentioned in the technical
	Emergency N2O Manifold (Without Cylinders)	specification of tender.
	Each header bar assembly shall be provided with	
	a high pressure shut off valve. It may be noted	
	that HTM or ISO Standards does not permit the	
	use of Shut Off Valves on the Headers.	
	Moreover, if any Shut Off Valve are provided on	
	Manifold Header, then the same will not be	
	compliant / satisfying the requirement of HTM /	
	ISO Standards.	
	Hence, we request you to delete the sentence:	
	Header bar/s assembly shall be provided with a	
	high pressure shut off valve.	
12		Tender terms & conditions
12	Vol 4 - Technical Specifications,	Tender terms & conditions prevail.
12	Vol 4 - Technical Specifications, Page 10, Para 3	Tender terms & conditions prevail.
12	Vol 4 - Technical Specifications, Page 10, Para 3 MEDICAL AND SURGICAL AIR SYSTEM:	
12	Vol 4 - Technical Specifications, Page 10, Para 3 MEDICAL AND SURGICAL AIR SYSTEM: The medical air plant shall fully comply with the	
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12	Vol 4 - Technical Specifications, Page 10, Para 3 MEDICAL AND SURGICAL AIR SYSTEM: The medical air plant shall fully comply with the requirements of the HTM 02-01/ NFPA 99C/EN/DIN/ISO 7396-1. It should be US FDA/European CE certified with 4 digit notified	
	Vol 4 - Technical Specifications, Page 10, Para 3 MEDICAL AND SURGICAL AIR SYSTEM: The medical air plant shall fully comply with the requirements of the HTM 02-01/ NFPA 99C/EN/DIN/ISO 7396-1. It should be US FDA/European CE certified with 4 digit notified body number or American ETL/ UL listed	prevail.
12	Vol 4 - Technical Specifications, Page 10, Para 3 MEDICAL AND SURGICAL AIR SYSTEM: The medical air plant shall fully comply with the requirements of the HTM 02-01/ NFPA 99C/EN/DIN/ISO 7396-1. It should be US FDA/European CE certified with 4 digit notified body number or American ETL/ UL listed It may be noted that NFPA99C doesnot permit	prevail. Combination or de-
	Vol 4 - Technical Specifications, Page 10, Para 3 MEDICAL AND SURGICAL AIR SYSTEM: The medical air plant shall fully comply with the requirements of the HTM 02-01/NFPA 99C/EN/DIN/ISO 7396-1. It should be US FDA/European CE certified with 4 digit notified body number or American ETL/ UL listed It may be noted that NFPA99C doesnot permit for use of a Combined Medical & Surgical Air	Combination or decombination of Medical Air
	Vol 4 - Technical Specifications, Page 10, Para 3 MEDICAL AND SURGICAL AIR SYSTEM: The medical air plant shall fully comply with the requirements of the HTM 02-01/NFPA 99C/EN/DIN/ISO 7396-1. It should be US FDA/European CE certified with 4 digit notified body number or American ETL/UL listed It may be noted that NFPA99C doesnot permit for use of a Combined Medical & Surgical Air System.	Combination or decombination of Medical Air & Surgical Air system
	Vol 4 - Technical Specifications, Page 10, Para 3 MEDICAL AND SURGICAL AIR SYSTEM: The medical air plant shall fully comply with the requirements of the HTM 02-01/NFPA 99C/EN/DIN/ISO 7396-1. It should be US FDA/European CE certified with 4 digit notified body number or American ETL/ UL listed It may be noted that NFPA99C doesnot permit for use of a Combined Medical & Surgical Air System. Medical Air and Surgical Air Plants are required	Combination or decombination of Medical Air & Surgical Air system should be followed as per
	Vol 4 - Technical Specifications, Page 10, Para 3 MEDICAL AND SURGICAL AIR SYSTEM: The medical air plant shall fully comply with the requirements of the HTM 02-01/NFPA 99C/EN/DIN/ISO 7396-1. It should be US FDA/European CE certified with 4 digit notified body number or American ETL/ UL listed It may be noted that NFPA99C doesnot permit for use of a Combined Medical & Surgical Air System. Medical Air and Surgical Air Plants are required to be separate as per NFPA99C.	Combination or decombination of Medical Air & Surgical Air system should be followed as per standards mentioned in the
	Vol 4 - Technical Specifications, Page 10, Para 3 MEDICAL AND SURGICAL AIR SYSTEM: The medical air plant shall fully comply with the requirements of the HTM 02-01/NFPA 99C/EN/DIN/ISO 7396-1. It should be US FDA/European CE certified with 4 digit notified body number or American ETL/UL listed It may be noted that NFPA99C doesnot permit for use of a Combined Medical & Surgical Air System. Medical Air and Surgical Air Plants are required to be separate as per NFPA99C. If any bidder is offering a combined Medical &	Combination or decombination of Medical Air & Surgical Air system should be followed as per standards mentioned in the technical specification of
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13	Vol 4 - Technical Specifications, Page 10, Para 3 MEDICAL AND SURGICAL AIR SYSTEM: The medical air plant shall fully comply with the requirements of the HTM 02-01/NFPA 99C/EN/DIN/ISO 7396-1. It should be US FDA/European CE certified with 4 digit notified body number or American ETL/UL listed It may be noted that NFPA99C doesnot permit for use of a Combined Medical & Surgical Air System. Medical Air and Surgical Air Plants are required to be separate as per NFPA99C. If any bidder is offering a combined Medical & Surgical Air plant and confirming that it is as per NFPA, then it is a complete violation of NFPA99C standards. This may be kindly noted and taken into consideration. Vol 4 - Technical Specifications, Page 11, Para 3.1	Combination or decombination of Medical Air & Surgical Air system should be followed as per standards mentioned in the technical specification of tender.
13	Vol 4 - Technical Specifications, Page 10, Para 3 MEDICAL AND SURGICAL AIR SYSTEM: The medical air plant shall fully comply with the requirements of the HTM 02-01/NFPA 99C/EN/DIN/ISO 7396-1. It should be US FDA/European CE certified with 4 digit notified body number or American ETL/UL listed It may be noted that NFPA99C doesnot permit for use of a Combined Medical & Surgical Air System. Medical Air and Surgical Air Plants are required to be separate as per NFPA99C. If any bidder is offering a combined Medical & Surgical Air plant and confirming that it is as per NFPA, then it is a complete violation of NFPA99C standards. This may be kindly noted and taken into consideration. Vol 4 - Technical Specifications, Page 11, Para 3.1 MEDICAL AND SURGICAL AIR SYSTEM:	Combination or decombination of Medical Air & Surgical Air system should be followed as per standards mentioned in the technical specification of tender. Tender terms & conditions
13	Vol 4 - Technical Specifications, Page 10, Para 3 MEDICAL AND SURGICAL AIR SYSTEM: The medical air plant shall fully comply with the requirements of the HTM 02-01/ NFPA 99C/EN/DIN/ISO 7396-1. It should be US FDA/European CE certified with 4 digit notified body number or American ETL/ UL listed It may be noted that NFPA99C doesnot permit for use of a Combined Medical & Surgical Air System. Medical Air and Surgical Air Plants are required to be separate as per NFPA99C. If any bidder is offering a combined Medical & Surgical Air plant and confirming that it is as per NFPA, then it is a complete violation of NFPA99C standards. This may be kindly noted and taken into consideration. Vol 4 - Technical Specifications, Page 11, Para 3.1 MEDICAL AND SURGICAL AIR SYSTEM: Air Compressor Modules:	Combination or decombination of Medical Air & Surgical Air system should be followed as per standards mentioned in the technical specification of tender. Tender terms & conditions
13	Vol 4 - Technical Specifications, Page 10, Para 3 MEDICAL AND SURGICAL AIR SYSTEM: The medical air plant shall fully comply with the requirements of the HTM 02-01/NFPA 99C/EN/DIN/ISO 7396-1. It should be US FDA/European CE certified with 4 digit notified body number or American ETL/UL listed It may be noted that NFPA99C doesnot permit for use of a Combined Medical & Surgical Air System. Medical Air and Surgical Air Plants are required to be separate as per NFPA99C. If any bidder is offering a combined Medical & Surgical Air plant and confirming that it is as per NFPA, then it is a complete violation of NFPA99C standards. This may be kindly noted and taken into consideration. Vol 4 - Technical Specifications, Page 11, Para 3.1 MEDICAL AND SURGICAL AIR SYSTEM: Air Compressor Modules: It should be Oil-Less Screw Compressors /Scroll	Combination or decombination of Medical Air & Surgical Air system should be followed as per standards mentioned in the technical specification of tender. Tender terms & conditions
13	Vol 4 - Technical Specifications, Page 10, Para 3 MEDICAL AND SURGICAL AIR SYSTEM: The medical air plant shall fully comply with the requirements of the HTM 02-01/ NFPA 99C/EN/DIN/ISO 7396-1. It should be US FDA/European CE certified with 4 digit notified body number or American ETL/ UL listed It may be noted that NFPA99C doesnot permit for use of a Combined Medical & Surgical Air System. Medical Air and Surgical Air Plants are required to be separate as per NFPA99C. If any bidder is offering a combined Medical & Surgical Air plant and confirming that it is as per NFPA, then it is a complete violation of NFPA99C standards. This may be kindly noted and taken into consideration. Vol 4 - Technical Specifications, Page 11, Para 3.1 MEDICAL AND SURGICAL AIR SYSTEM: Air Compressor Modules:	Combination or decombination of Medical Air & Surgical Air system should be followed as per standards mentioned in the technical specification of tender. Tender terms & conditions

	<u></u>	<u></u>
	standby. HTM & ISO Standards permit use of Oil injected	
	Air Screw Compressors for Medical & Surgical Air.	
	It may be noted that Lubricated Screw	
	Compressors are very economical in terms of	
	initial investment and also maintenance and	
	lifetime cost compared to oilless scroll / screw.	
	This type of Air Compressors are installed all	
	over the world in major hospitals.	
	Also, it is to be noted that a Scroll technology is	
	not a very good choice for 10 Bar plant	
	requirement.	
	Hence, we request you to include oil injected	
1.5	screw compressors in the specifications.	
15	Vol 4 - Technical Specifications,	Padlocks available to allow
	Page 11, Para 3.1	locking of the valves in both
	Pressure Reducing Station:	open and closed positions
	Padlocks available to allow locking of the valves	and must have easy to read
	in both open and closed positions and must have	pressure gauges or as per
	easy to read pressure gauges or as per guideline	guideline of standard to be
	of standard to be followed.	followed.
		Base plate mounted and
	We fail to understand why padlocks are required	supplied with copper stub
	to be provided in the Pressure reducing Station?	pipes for ease of installation
	Please note that this appears to be a particular	using inert joining
	make specific. Hence, we request you to delete	procedures
16	the requirement of padlocks for the valves.	The air receiver / vacuum
10	Vol 4 - Technical Specifications, Page 12, Para 3.2	reservoir capacity should be
	1 age 12, 1 ara 3.2	as per standard mentioned in
	Vertical Air Receiver:	the technical specification of
	Total air receiver capacity shall be at least 50%	tender.
	$(\pm 5\%)$ of the primary plant capacity mentioned	
	in the BOQ) in 1 minute in terms of free air	
	delivered at normal working pressure or as per	
	guideline of standard to be followed.	
	Our submission is that, for fair evaluation and	
	comparison of bidders of various standards and	
	for parity purposes, the minimum capacity of the	
	Air Receiver should be specified, which should	
	be complied by bidders for all standards.	
	If an open ended statement like as per guideline	
	of standard to be	
	followed is mentioned, then this would be a huge	
	disadvantage for HTM	
	/ ISO eqpt suppliers.	
	Contrarily, it may be noted that, NFPA does not	

	have any guideline on sizing the capacity of Air	
	Receivers. It has been left to the discretion of	
	manufacturers.	
	It has been observed that Medical Air Systems of	
	NFPA Standards are usually supplied with a	
	small capacity Air Receivers.	
	Hence, to avoid this ambiguity, we suggest that	
	minimum capacity of Air Receiver is clearly	
	specified in the amended tender specs, and which	
	should be followed by all bidders.	
17	Vol 4 - Technical Specifications,	System controls be offered
	Page 12, Para 3.4	as per the requirement of the
	System Controls:	standard.
	The cabinet shall have status display to include	
	system pressure, dew point pump operation,	
	accumulated time, maintenance interval, fault	
	conditions, and silence button, lighted Hand-Off-	
	Automatic selector switches	
	It appears that the specification is of a particular	
	make and standard.	
	We request you to delete the words silence	
	button, lighted Hand-Off-Automatic selector	
	switches.	
	Further, it is suggested that this be amended as:	
	System controls be offered as per the requirement	
	of the standard.	
18	Vol 4 - Technical Specifications,	The air receiver / vacuum
10	Page 13, Para 4.2	reservoir capacity should be
	1 age 13, 1 ata 4.2	as per standard mentioned in
	Vacuum Receiver:	the technical specification of
	Vacuum reservoir shall have total volume of at	tender.
		tender.
	least 100 % of Primary plant output(± 5%)	
	(Capacity as mentioned in the BOQ)	
	Our submission is that for fair avaluation and	
	Our submission is that, for fair evaluation and	
	comparison of bidders of various standards and	
	for parity purposes, the minimum capacity of the	
	Vacuum Receiver should be specified, which	
	should be complied by bidders for all standards.	
	If an open ended statement like as per guideline	
	of standard to be followed is mentioned, then this	
	would be a huge disadvantage for HTM/ISO eqpt	
	<u> </u>	
1	suppliers. Contrarily, it may be noted that, NFPA	
	suppliers. Contrarily, it may be noted that, NFPA does not have any guideline on sizing the	
	suppliers. Contrarily, it may be noted that, NFPA does not have any guideline on sizing the capacity of Vacuum Receivers. It has been left to	
	suppliers. Contrarily, it may be noted that, NFPA does not have any guideline on sizing the capacity of Vacuum Receivers. It has been left to the discretion of manufacturers. It has been	
	suppliers. Contrarily, it may be noted that, NFPA does not have any guideline on sizing the capacity of Vacuum Receivers. It has been left to the discretion of manufacturers. It has been observed that Medical Vacuum Systems of NFPA	
	suppliers. Contrarily, it may be noted that, NFPA does not have any guideline on sizing the capacity of Vacuum Receivers. It has been left to the discretion of manufacturers. It has been observed that Medical Vacuum Systems of NFPA Standards are usually supplied with a small	
	suppliers. Contrarily, it may be noted that, NFPA does not have any guideline on sizing the capacity of Vacuum Receivers. It has been left to the discretion of manufacturers. It has been observed that Medical Vacuum Systems of NFPA	

	minimum capacity of Vacuum Receiver is clearly	
	specified in the amended tender specs, and which should be followed by all bidders.	
19	Vol 4 - Technical Specifications,	System controls be offered
	Page 14, Para 4.3	as per the requirement of the standard.
	System Controls:	
	The cabinet shall have status display to include	
	system pressure, dew point pump operation, accumulated time, maintenance interval, fault	
	conditions, and silence button, lighted Hand-Off-	
	Automatic selector switches	
	It appears that the specification is of a particular make and standard.	
	We request you to delete the words silence	
	button, lighted Hand-Off-Automatic selector	
	switches. Further, it is suggested that this be amended as:	
	System controls be offered as per the requirement	
	of the standard.	
20	Vol 4 - Technical Specifications,	Tender terms and conditions
	Page 14, Para 4.4	prevail
	Bacterial Filters:	
	The dryer should be particulate filter dryer with	
	ability to remove particles as small as 1 micron.	
	This appears to be a typo error.	
21	Please clarify and issue necessary amendments.	The modes 1 - 11
21	Vol 4 - Technical Specifications, Page 14, Para 8	The package should consist of two dry rotary
	1 1,1 1111 0	vane/Claw type vacuum
	AGSS (Anesthetic Gas Scavenging System)	pumps (Dry/Oiless) or
	Plant: The package should consist of two dry rotary	Blower pump as per guideline of standard
	vane/Claw type vacuum pumps (Dry/Oiless) or as	guideinie di Standald
	per guideline of standard to be followed,	
	As per HTM & ISO Standards, the Blower type	
	Pumps are normally used in the AGSS System.	
22	Please confirm if this is acceptable.	Quantity mantioned in the
22	Vol 4 - Technical Specifications, Page 14, Para 8	Quantity mentioned in the attached revised BOQ
	AGSS (Anaesthetic Gas Scavenging System)	
	Plant: Connecting hose suitable to fit with anaesthesia	
	Workstation should be provided.	

	Discourse its the Oter C.C. ti	
	Please provide the Qtys of Connecting Hoses	
	required for the Anaesthesia Workstations and	
	their Technical Specs.	
	Ideally, the connecting hoses are supplied by the	
	respective Anaesthesia Workstation Suppliers,	
	and therefore can be deleted from the tender	
	specs.	
23	Vol 4 - Technical Specifications,	The Area Valve Service
	Page 17, Para 11	Unit should incorporate pre-
	1 450 17,1 414 11	fitted ball valve in a box
	AREA VALVE SERVICE UNIT:	
		with emergency access.
	The Area Valve Service Unit should incorporate	
	a ball valve with NIST/else connectors either side	77.1
	mounted in a lockable box with emergency	Valve sizes and quantities
	access or as per guideline of standard to be	are mentioned in the BOQ.
	followed. Please confirm if the AVSU is required	
	to be supplied along with ball Valves? If yes,	
	specify the valve sizes configuration required in	
	each of the AVSU models 2 gas - 6 gas services.	
24	Vol 4 - Technical Specifications,	Shall be as per standards
	Page 17, Para 11	mentioned in the technical
		specification of tender
	AREA VALVE SERVICE UNIT:	specification of tender
	The Area Valve Service Unit should incorporate	
	a ball valve with NIST/else connectors either side	
	mounted in a lockable box with emergency	
	access or as per guideline of standard to be	
	followed.	
	Our submission is that, for fair evaluation and	
	comparison of bidders of various standards and	
	for parity purposes, and also for the Safety and	
	backup arrangement and continuity of gas	
	supplies of the MGPS, the NIST is a very critical	
	requirement for AVSU, which should be	
	complied by bidders for all standards.	
	If an open ended statement like as per guideline	
	of standard to be followed is mentioned, then this	
	would be a huge disadvantage for HTM/ ISO	
	eqpt suppliers.	
	11 11	
	Contrarily, it may be noted that, NFPA doesnot	
	categorically mention about the requirement of	
	NIST. However, the Isolation Valves as per	
	NFPA Mandates to have provisions for inlet ports	
	on the pipe extensions, wherein the NIST	
	connectors can be fixed. Hence, to avoid this	
	ambiguity, we suggest that NIST Connectors are	
	mandatory is clearly specified in the amended	
	tender specs, and which should be followed by all	
	bidders.	
25	Vol 4 - Technical Specifications,	Extruded Aluminium/MS

Page 17, Para 11	powder coated
1 450 17,1 414 11	powder coaled
AREA VALVE SERVICE UNIT: The box shall be made from extruded aluminium to prevent corrosion or as per guideline of standard to be followed. It may be noted that the AVSU Box	
manufactured by all major suppliers is made of	
Steel and powder coated etc to prevent corrosion.	
Hence, we request you to kindly delete the wordings extruded aluminium from the tender	
	I .

specs.

26	Vol. 4 Tachnical Specifications	Didder shall be responsible for all
26	Vol 4 - Technical Specifications, Page 18, Para 12.1	Bidder shall be responsible for all cabling from local alarm panels
		(OTs, ICUs)to Master alarm
	Master Alarm :	panel.
	The master alarms should be capable to monitor	
	minimum 30-40 Point.	
	Bidder shall be responsible for all cabling from	
	local alarm panels to master alarm panel.	
	Please confirm if all the Area Alarms are	
	required to be connected with the Master Alarm	
	Panel by providing cabling? If Yes, then the Size of	
	Master Alarm Panel is not sufficient, as it has	
	provision to connect only 30-40 points.	
	Whereas, the total number of AVSU is 93 Nos.	
27	Vol 4 - Technical Specifications,	The medical gas central alarms
_,	Page 19, Para 12.2	should be capable of monitoring
	1 450 17,1 414 12.2	up to 6 medical gas services (As
	Medical Gas Area Alarm:	specified in BOQ)
	The medical gas central alarms should be	specified in 200)
	capable of monitoring up to 5 medical gas	
	services (As specified in BOQ)	
	It appears the central mentioned in the tender	
	specs is a typo error.	
	Please clarify.	
	Also, the 5 medical gas services mentioned in	
	the specs is not correct, as the BOQ mentions	
	upto 6 medical gas services. Please clarify.	
28	Vol 4 - Technical Specifications,	Tender terms and conditions
	Page 22, Para 13b	prevail
	Manufacturer Authorization:	
	This is a limiting clause and appears to be	
	favouring few companies only and which may	
	restrict competition to a few bidders only. You	
	are requested to delete the requirement of	
	Manufacturers AuthorisationCertificates. By	
	getting committed to a particular manufacturer	
	while quoting, the bidders lose their ability to	
	negotiate better prices and terms at the time of	
	supply and also quite often are restricted to the	
	products of the particular manufacturer despite	
	better products becoming available from other	
	manufacturers while the tender is still under	
	consideration of the tenderer. This requirement	
	is inviting cartelisation because local / Indian	
	representatives of particular foreign	
	manufacturers have got their specification	
	incorporated in the major tender and the leading	
	bidder will have their supporting bidders.	

	Moreover, it may be noted that the requirement of Manufacturers				
	Authorisation Certificate is against the CCI Act 2012.				
29	Vol 4 - Technical Specifications, Page 24,	Tender prevail	terms	and	conditions
	Responsibility of bidder: 5. Rota meters for measurement of consumption of Oxygen and Compressed air To enable fair evaluation and parity among all bidders, Please provide Technical Specs for the				
	Rotameters. Also, request you to include this in the line item of the BOQ, as this is a major item.				
30	Vol 4 - Technical Specifications, Page 24, Para 15	Tender prevail	terms	and	conditions
	The following systems/items must be from the same principal company/Manufacturer: a. Control Panels & Manifold for O2, N2O & CO2				
	b. Medical air plant c. Medical Vacuum Plant d. AGSS Plant				
	e. Area & Master Alarm f. All types Outlets g. AVSU				
	h. Line Isolation valves i. High Pressure tubes				
	We request you to kindly delete the sentence The following systems/items must be from the same principal company/Manufacturer: , and allow the bidders to select the best possible supplier option, as long as all the MGPS Products are from the same single MGPS Standard selected for which the bid is being submitted.				
	It may be kindly noted that there are only a few and limited suppliers worldwide who manufacture (or outsource) all the MGPS products.				
	On the other hand, there are certain manufacturers who are having exceptionally good experience and reputation in their respective domain like supplying MGPS Plant Source Equipments (Medical Air				
	Plant, Vacuum & AGSS Plants), and there are other's who can offer exceptionally good quality				

	MODO Divilai o v E i v 1	T
	MGPS Distribution System Equipment and	
	similarly for Architectural Systems (Bed Head	
	panels etc), SOT	
	Products (Oxygen Flowmeter, Ward Vacuum	
	Units, Theatre Vacuum	
	Units etc).	
	Linde shall take the single point responsibilty for	
	all the equipment	
	supplied by the respective OEM Suppliers, and	
	shall be fully	
	responsible for the complete installation and	
	during warranty and CMC	
	Periods.	
21		Tandan tanna and acaditions
31	Vol 4 - Technical Specifications,	Tender terms and conditions
	Page 26, Para 23	prevail
	Inter connection to Manifolds with LMO tank	
	with necessary automatic switchover panels	
	between LMO & Manifold will be up to	
	responsibility of bidder up to 100m distance.	
	We request you to provide drawings of the site	
	to identify the location of proposed LMO system	
	and the location of Manifold Room, to enable	
	ascertain the distance, routing and cost	
	estimation.	
32.	Vol 3 - SCC, Page 35	The word 'Operation' is deleted
32	Vol 3 - SCC, Page 35,	The word 'Operation' is deleted.
32		The word 'Operation' is deleted.
32	After provisional taking over, the contractor	The word 'Operation' is deleted.
32	After provisional taking over, the contractor shall provide operation and maintenance	The word 'Operation' is deleted.
32	After provisional taking over, the contractor shall provide operation and maintenance services for the complete MEDICAL GAS	The word 'Operation' is deleted.
32	After provisional taking over, the contractor shall provide operation and maintenance services for the complete MEDICAL GAS MANIFOLD SYSTEM EQUIPMENT till the	The word 'Operation' is deleted.
32	After provisional taking over, the contractor shall provide operation and maintenance services for the complete MEDICAL GAS MANIFOLD SYSTEM EQUIPMENT till the successful completion of Defect Liability	The word 'Operation' is deleted.
32	After provisional taking over, the contractor shall provide operation and maintenance services for the complete MEDICAL GAS MANIFOLD SYSTEM EQUIPMENT till the successful completion of Defect Liability Period.	The word 'Operation' is deleted.
32	After provisional taking over, the contractor shall provide operation and maintenance services for the complete MEDICAL GAS MANIFOLD SYSTEM EQUIPMENT till the successful completion of Defect Liability Period. Please clarify the meaning of Operation. Will	The word 'Operation' is deleted.
32	After provisional taking over, the contractor shall provide operation and maintenance services for the complete MEDICAL GAS MANIFOLD SYSTEM EQUIPMENT till the successful completion of Defect Liability Period. Please clarify the meaning of Operation. Will the bidder have to provide manpower at site? If	The word 'Operation' is deleted.
32	After provisional taking over, the contractor shall provide operation and maintenance services for the complete MEDICAL GAS MANIFOLD SYSTEM EQUIPMENT till the successful completion of Defect Liability Period. Please clarify the meaning of Operation. Will the bidder have to provide manpower at site? If Yes, how many in each shift etc?	The word 'Operation' is deleted.
32	After provisional taking over, the contractor shall provide operation and maintenance services for the complete MEDICAL GAS MANIFOLD SYSTEM EQUIPMENT till the successful completion of Defect Liability Period. Please clarify the meaning of Operation. Will the bidder have to provide manpower at site? If Yes, how many in each shift etc? The BOQ file doesnot have relevant areas to	The word 'Operation' is deleted.
32	After provisional taking over, the contractor shall provide operation and maintenance services for the complete MEDICAL GAS MANIFOLD SYSTEM EQUIPMENT till the successful completion of Defect Liability Period. Please clarify the meaning of Operation. Will the bidder have to provide manpower at site? If Yes, how many in each shift etc? The BOQ file doesnot have relevant areas to provide the Operation Cost. Please amend the	The word 'Operation' is deleted.
32	After provisional taking over, the contractor shall provide operation and maintenance services for the complete MEDICAL GAS MANIFOLD SYSTEM EQUIPMENT till the successful completion of Defect Liability Period. Please clarify the meaning of Operation. Will the bidder have to provide manpower at site? If Yes, how many in each shift etc? The BOQ file doesnot have relevant areas to provide the Operation Cost. Please amend the BOQ file, if this scope of work is required to be	The word 'Operation' is deleted.
32	After provisional taking over, the contractor shall provide operation and maintenance services for the complete MEDICAL GAS MANIFOLD SYSTEM EQUIPMENT till the successful completion of Defect Liability Period. Please clarify the meaning of Operation. Will the bidder have to provide manpower at site? If Yes, how many in each shift etc? The BOQ file doesnot have relevant areas to provide the Operation Cost. Please amend the BOQ file, if this scope of work is required to be provided.	The word 'Operation' is deleted.
33	After provisional taking over, the contractor shall provide operation and maintenance services for the complete MEDICAL GAS MANIFOLD SYSTEM EQUIPMENT till the successful completion of Defect Liability Period. Please clarify the meaning of Operation. Will the bidder have to provide manpower at site? If Yes, how many in each shift etc? The BOQ file doesnot have relevant areas to provide the Operation Cost. Please amend the BOQ file, if this scope of work is required to be	The word 'Operation' is deleted. Tender terms and conditions
	After provisional taking over, the contractor shall provide operation and maintenance services for the complete MEDICAL GAS MANIFOLD SYSTEM EQUIPMENT till the successful completion of Defect Liability Period. Please clarify the meaning of Operation. Will the bidder have to provide manpower at site? If Yes, how many in each shift etc? The BOQ file doesnot have relevant areas to provide the Operation Cost. Please amend the BOQ file, if this scope of work is required to be provided.	
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	After provisional taking over, the contractor shall provide operation and maintenance services for the complete MEDICAL GAS MANIFOLD SYSTEM EQUIPMENT till the successful completion of Defect Liability Period. Please clarify the meaning of Operation. Will the bidder have to provide manpower at site? If Yes, how many in each shift etc? The BOQ file doesnot have relevant areas to provide the Operation Cost. Please amend the BOQ file, if this scope of work is required to be provided. Vol 3 - SCC, Page 49,	Tender terms and conditions
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	After provisional taking over, the contractor shall provide operation and maintenance services for the complete MEDICAL GAS MANIFOLD SYSTEM EQUIPMENT till the successful completion of Defect Liability Period. Please clarify the meaning of Operation. Will the bidder have to provide manpower at site? If Yes, how many in each shift etc? The BOQ file doesnot have relevant areas to provide the Operation Cost. Please amend the BOQ file, if this scope of work is required to be provided. Vol 3 - SCC, Page 49, MANUFACTURER'S AUTHORIZATION FORM: Dear Sir,	Tender terms and conditions
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	After provisional taking over, the contractor shall provide operation and maintenance services for the complete MEDICAL GAS MANIFOLD SYSTEM EQUIPMENT till the successful completion of Defect Liability Period. Please clarify the meaning of Operation. Will the bidder have to provide manpower at site? If Yes, how many in each shift etc? The BOQ file doesnot have relevant areas to provide the Operation Cost. Please amend the BOQ file, if this scope of work is required to be provided. Vol 3 - SCC, Page 49, MANUFACTURER'S AUTHORIZATION FORM: Dear Sir, Tender No	Tender terms and conditions

	agent) which has been our dealer/distributor	
	since, to submit a bid, and sign the contract with	
	you for the goods manufactured by us against	
	the above tender.	
	No company or firm or individual other than	
	M/s are authorized to bid and conclude the	
	contract for goods manufactured by us against	
	this specific tender.	
	We hereby extend our full guarantee and defect	
	Liability period as per the clause of Condition of	
	Contract and Additional Specific Conditions of	
	Contract of above tender for goods and services	
	offered for supply by our authorized firm.	
	To enable a Principal Supplier to follow, we	
	request you to amend the	
	format as:	
	MANUFACTURER'S / PRINCIPAL	
	SUPPLIER'S AUTHORIZATION FORM:	
	Dear Sir,	
	Tender No	
	We who are established and reputed	
	manufacturer / Supplier of (name & description	
	of goods offered) having factories at (address of	
	factory) do hereby authorize M/s (Name &	
	address of agent) which has been our	
	dealer/distributor since, to submit a bid, and sign	
	the contract with you for the goods	
	manufactured / supplied by us against the above	
	tender.	
	No company or firm or individual other than	
	M/s are authorized to bid and conclude the	
	contract for goods manufactured /supplied by us	
	against this specific tender.	
	We hereby extend our full guarantee and defect	
	liability period as per the clause of Condition of	
	Contract and Additional Specific Conditions of	
	Contract of above tender for goods and services	
34	offered for supply by our authorized firm.	Davied of work completion 7
34	Vol 1- Page 3 Period of work completion 6 Months	Period of work completion – 7 Months from the date of letter of
	Period of work completion – 6 Months Considering the huge quantum of work and	commencement
	imported equipment which needs to be arranged,	
	we request you to kindly amend the work completion time to minimum 9 Months	
35	Vol-III, Special Conditions of	Tender terms and conditions
33	Contract, Page No. SCC-, Clause	Tender terms and conditions prevail
	39.2.4, Page 25	pievan
	Water Supply & Power Supply:	
	The contractor will provide water & electricity	
	The confractor will provide water & electricity	

	to the Consultant's office free of cost for the required quantity by the Consultant's site office. MGPS Works require very less electrical power and water for construction purposes. We request to delete the sentence: The contractor will provide water & electricity to the Consultant's office free of cost for the required quantity by the Consultant's site office.	
36	Vol-II, General Conditions of Contract, Clause 47.1 & Annexure- B (Appendix To Tender), Page 38 & 71 Liquidated Damages for Delay: Amount of Liquidated damages: 1% (one percent) of contract price per calendar week of delay Limit of liquidated damages: 10% (Ten percent) of contract price. Please consider the aggregate maximum of liquidated damages payable under clause No. 47.1 shall not exceed 0.5% of contract value per week of delay and shall be subjected to maximum amount of 5% on overall contract price.	Tender terms and conditions prevail
37	General Point Release of Final 10% Payment After completion of installation works, if the Commissioning / Trial Run of the MGMS system is delayed for more than 3 months due to reasons not attributable to Contractor, then the final 10% payment should be released without any further delay against submission of Bank Guarantee. Please confirm acceptance of this.	Release of final 10% payment of BOQ contract rates after final acceptance of system by the client
38	General Point DLP Period Start Date After completion of installation works, if the Commissioning/ Trial run of the MGMS system is delayed for more than 3 months due to reasons not attributable to Contractor, then DLP period start date would be considered from that date. Please confirm acceptance of this.	Tender terms and conditions prevail
39	General Point Customs Duty Please confirm customs duty is under customer or bidders scope. Also confirm the applicable rate of customs duty for the job. Also confirm if CDEC (Customs Duty Exemption Certificate)	Tender terms and conditions prevail

	would be issued by customer. If Yes, what	
	would be the rate of Customs Duty exemption?	
40	General Point	Tender terms and conditions
	Statutory Duties	prevail
	In case of any statutory variation in duties like	
	GST, Customs Duty, IGST etc within the	
	contractual delivery date shall be borne by	
	client.	
	Kindly confirm this.	
41	For safety of personnel & patients at hospital premises, it is highly recommended that all cylinders are fitted with Valve Guards to prevent incidents during cylinder handling / tripping. Hence, we request you to include the following in the Technical Specs for Cylinders:	Valve Guard should be fitted on all cylinders for the protection of Cylinder Valve
	14. Supply of O2 Cylinders – Class D Type Should be as per BIS/IS/ASME Standard. Valve Guard should be fitted on all cylinders for the protection of Cylinder Valve, to prevent incidents happening during cylinder handling / tripping.	
	15. Supply of N2O Cylinders – Class D Type Should be as per BIS/IS/ASME Standard Valve Guard should be fitted on all cylinders for the protection of Cylinder Valve, to prevent incidents happening during cylinder handling / tripping.	
	16. Supply of CO2 Cylinders – Class D Type Should be as per BIS/IS/ASME Standard Valve Guard should be fitted on all cylinders for the protection of Cylinder Valve, to prevent incidents happening during cylinder handling / tripping.	
42	Oxygen Flow meter with Humidifier Bottle We understand that you missed out to mention USFDA/European CE Certified with 4 digit notified body no or American UL /ETL Listed as you have mentioned the same for Ward Vacuum Unit and Theatre Vacuum unit. You are requested to kindly amend and mention European CE Marked with 4 Digit Notified Body No /USFDA Certified /UL Listed /ETL Listed.	BIS/US FDA/European CE Certified with 4 digit notified body number or American ETL/ UL listed.

43	Humidifier Bottle asked in the tender is of polycarbonate /polysulfone material. We recommend adding polypropylene material for humidifier bottle, which is better quality material autoclavable at 134°C Medical and Surgical Air System You missed out to mention the Variation of +1-10% for Vacuum System as mentioned in past tenders published by HSCC. You are requested to kindly amend the same and also mention the same in BOQof tender. • In the BOQ, it is mentioned that vendor may offer two plants. Since the total Flow requirement for is 15000 LPM, may we offer 7500 LPM working and 7500 LPM as standby or 9000 LPM as working or 6000 LPM as standby. Please clarify and confirm the same Vacuum System – Youmissed out to mention the Variation of +1-10% for Vacuum System as mentioned in past tenders published by HSCC. You are requested to kindly amend the same and also mention the same in BOQof tender	+/-10 % to flow capacity of plant is permitted Tender terms and conditions prevail +/-10 % to flow capacity of plant is permitted
45	 Master Alarm / Medical Gas Area Alarm In HTM Standard, the Area / Master Alarms are LED Type and not Digital / Touch Screen type. Kindly amend the same accordingly. In Bill of Quantity, for Medical Gas Area Alarm for 2 Services, there is a typographical 	 Digital or as per standard mentioned in the technical specification of tender Read as Oxygen and
	error in mentioning the gases for 2 Service Alarm, It should be Oxygen and Vacuum instead of MA4Bar. You are requested to kindly correct the same. • Further the specifications mentioned in the tender are more of NFPA Standard, you are requested to kindly add technical specifications as per HTM 02-01 standard for the following items as published by HSCC in their past tenders of MGPS, For your reference, copy of technical specifications enclosed: Master Alarm and Area AlarmGas Outlet	Vacuum instead of MA4 bar. • Tender terms and conditions prevail
46	Horizontal / Vertical Bed Head Panel- You are requested to kindly amend the same to 3 Channel instead of 2 tier / 2 Channel, because electrical and Gas Outlets are not possible to mount of 2 channel Bed Head Panel.	3 Channel / 2 tier / 2 Channel/3 Partition rows

47	Responsibility of bidder - Being an accessory item, High Pressure tubing must not be a part of this clause. You are therefore requested to kindly delete the same from clause 15 of responsibility of bidders.	Tender terms and conditions prevail
48	Point to be Clarified - Please confirm who will supply 200 KW DGSet and when?	Tender terms and conditions prevail
49	1.2 of INSTRUCTION TO APPLICANTS 5 of Volume – I COMPLETION PERIOD: 6 months from the date of order of commencement. We request you to kindly change the COMPLETION PERIOD to 12 months instead of 6 months from the date of order of commencement.	Period of work completion – 7 Months from the date of letter of commencement
50	2.2 (ii) of INSTRUCTION TO APPLICANTS 5 of Volume – I The Applicant should meet the following minimum criteria for Pre-Qualification: Experience of having successfully completed similar work during last 7 years ending last day of month previous to the one in which tenders are invited should be either of the following: Three similar* completed works costing not less than the amount equal to 40% of the estimated cost. or Two similar* completed works costing not less than the amount equal to 50% of the estimated cost. or One similar* completed work costing not less than the amount equal to 80% of the estimated cost. We wish to inform you that, we fulfill all your eligibility criteria to great extent. However, we have observed that the Pre- Qualification Criteria is restricted & biased to favour some proven cartel companies to participate. In view of this, we request you to kindly amend the Pre-Qualification Criteria in line with 2 Options requested below. This will ensure wider participation from various bidders including us and there will be a price advantage to HSCC in view of huge participation, price comparison,	Tender terms and conditions prevail

and no encouragement to cartelization.

OPTION 1: Please go through attached HITES Tender Eligibility criteria for this size of MGPS work and same may be amended accordingly. We are surprised to note that HSCC and HITES both are under Ministry of Health & Family Welfare but asking different Eligibility Criteria. PQ Criteria - Request for Amendment:

"Minimum work of similar Nature: Eligible bidders should have successfully executed globally in last Seven years from the date of tender opening, similar turnkey project of value, equivalent to or exceeding 50% of the estimated schedule / tender value. Out of total 50% value, at least one single order for similar work of minimum 10% of the estimated schedule / tender value should have been executed globally". Please refer to Annexure - II for similar condition in one of the tenders of HITES issued on 14.02.2019.

OPTION 2:

(i) As can be seen we are very eager to participate in this tender and in this regard, we request you to kindly allow the bidders to submit the "Experience of having successfully completed similar work during last 10 years ending last day of month previous to the one in which tenders are invited instead of 7 years:

51 Standards/ Guideline

1 of Volume – IV

Standards/Guideline

The design & selection of all imported items should be of international standard like NFPA 99(latest version) standard and UL listed or ISO-7396-1/DIN/ EN (latest version) and UL listed/European CE or HTM 02 01 (latest version) guideline and European CE. This supersedes single/multiple standards mentioned at any other places in the tender specification involving item/system/capacity etc. The imported products should be of one standard only. All indigenous items should be of high quality and to be compatible to the main system.

We have observed that most of the technical

Design following one of International standards mentioned in the technical specification of tender will be accepted.

	specifications are written around one standard only i.e. NFPA. This may please be amended to Generic specifications in nature and OEM's should allowed to use their own design and follow any of the international standards. In case you do not want to change the present technical specifications, you may need to mention somewhere in the tender that "The tender technical specification are general in nature. However, manufacturer own design following one of the given international standard in the tender will be accepted, without compromising the flow rate".	
52	10.1	The Performance security shall be
		5% of the estimated cost.
	GCC - 10 of	
	VOLUME – II	
	Performance Security: The Contractor shall	
	provide security for his proper performance of	
	the Contract to the Employer within 15 days	
	after the receipt of the Letter of Acceptance. The	
	performance security shall be in the form of	
	bank guarantee. The amount of the bank	
	guarantee shall be 10 percent of the Contract	
	Price. It shall be issued by a Nationalised bank of India. When providing such security to the	
	of India. When providing such security to the Employer, the Contractor shall notify the	
	Engineer of so doing.	
	Engineer or so doing.	
	Performance BG should not be more than 5% of	
	order value and submission must be minimum	
	30 days from the date of your firm order. In	
	place of Nationalised Bank you should mention	
	Scheduled banks.	
53	12.1 & 12.2	Tender terms and conditions
		prevail
	SCC-12 of	r
	Volume – III	
	Bid Security: The Bidder shall furnish, as part of	
	his Bid, a Bid	
	Security of the amount of Rs. 25, 00, 000/-	
	(Rupees Twenty Five	
	Lakhs Only) for Medical Gas Manifold System,	
	Hospital Block, AIIMS,	
	Nagpur. No deviation shall be permitted from	
	this.	
	The Bid Security shall be in the form of a	
	Demand Draft/Pay	
	· · · · · · · · · · · · · · · · · · ·	•

Order/Bank Guarantee in favour of HSCC (India) Ltd. Payable at New

Delhi/NOIDA from any Nationalised/Scheduled bank.

Bid Security may please be reduced to 1% of estimated value only which means Rs. 15 lac in place of Rs. 25 lac,

you can understand that this blocks bidders finances as you know government takes long time in deciding the

tender and then returning the EMD and with this huge amount bidders cannot bid other on going and upcoming

tenders even in your case you have floated three big tenders on almost on same dates. This is unfair and there is

no such CVC guidelines for this huge EMD amount which some of you were referring in pre-bid meeting.

54 21.0 & 21.1

SCC-38 of Volume – III

Terms of Payment: For purposes of estimating the contract value

of works executed for certificate of payment, the following norms

shall be followed:

- 70 % of the BOQ contract rates on delivery of equipment at site
- after inspection ,passing and issue of dispatch clearance on pro-data

basis.

- 20% of BOQ contract rates on satisfactory take over certificate by
- client after erection and installation, testing and commissioning of

equipments on pro-data basis.

- 10 % of BOQ contract rates after successful completion of trial run of
- 30 days from the date of handover to the client
- (1) Please amend the payment terms as: 85% against delivery of material on or before 30 days from the date of supply the material. 10% against installation on pro-rata basis within 30

- 1) 75% of the BOQ contract rates on delivery of equipment at site after Inspection and Despatch Clearance Report on pro-data basis.
- 2) 15% of BOQ contract rates on Installation (Installation certificate to be provided with bill) of MGPS.
- 3) 10% of BOQ contract rates after final acceptance of system by the client

days of submission of bills and balance 5% against testing and commissioning within 30 days of submission of bills, in case site is not ready or electrical supply, gas supplies are not provided, contractor payment will be released within 30 days of testing and commissioning against undertaking that as and when electricity and gases will be provided contractor will do the needful without any extra charges. (2) Since, your technical specifications asking for International standards which can only be met with foreign manufacturer in totality which clearly means that you are looking for imported components. In such case you should give an option for quoting in foreign currency and you need to pay these foreign suppliers by means of letter of credit, with this you will be 100% assured that you are getting 100% imported material directly from manufacturers. This will avoid foreign makes stamping on local products, which is happening very commonly. Standards/Guideline Tender conditions and terms prevail 1 of Volume – IV Standards/Guideline The design & selection of all imported items should be of international standard like NFPA 99(latest version) standard and UL listed or ISO-7396-1/DIN/ EN (latest version) and UL listed/European CE or HTM 02 01 (latest version) guideline and European CE. This supersedes single/multiple standards mentioned at any other places in the tender specification involving item/system/capacity etc. The imported products should be of one standard only. All indigenous items should be of high quality and to be compatible to the main system. We have observed that most of the technical specifications are written around one standard only i.e. NFPA and make specific for example Item 1.1, Page No. 1

& 2, Item 2.1, Page No. 9 & 10, Item 3, Page

Item 4, Page No. 13 & 14, Item 10, Page No. 17, Item 12, Page No. 18 & 19, Item 17 on Page No.

No. 10, 11, 12 & 13,

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19 and Item 20, Page No. 21. This may please be amended to Generic specifications in nature and OEM's should be allowed to use their own design as per any of the international standards. In case you do not want to change the present technical specifications, you may need to mention somewhere in the tender that "The tender technical specification are general in nature. However, manufacturer own design following one of the given international standard in the tender will be accepted, without compromising the flow rate". For Example: In case of Air Plant please remove the word oil-less compressors because as per HTM which is mentioned in your tender does not require oil-less compressors because all the UK manufacturers who are producing Air plants as HTM are using oil-floaded compressors (because their efficiencies are many times higher than oil-less compressors) and all UK hospitals and even in Europe as per ISO:7396 and Middle east hospitals are using oilflooded air compressors. Once you are demanding HTM standard / recommendations then it is the responsibility of manufacturer to design and manufacture the air plants with medical air quality by using any types of air compressors but by using this oil-less word you only inviting **NFPA** standard recommendations and mentioning of HTM is an eye. We strongly object this approach to favour only proven cartel companies. 1.5 conditions Tender terms and prevail 3 of Volume – IV LIQUID MEDICAL OXYGEN STORAGE TANK The double walled Vacuum Insulated

Evaporator shall be constructed of stainless steel inner vessel contained within a carbon steel outer vessel. The annular space between the vessels shall be filled with non-inflammable perlite insulation material to insulate under vacuum. The VIE should be self-pressurizing type by partial evaporation of liquid oxygen

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through a pressure building coil by a non-ferrous imported pressure regulator. The vessel shall be supplied as a functional whole with all materials of construction & the cleaning regime suitable for medical grade liquid oxygen.

Please remove this item "LIQUID TANK" because 99% of the government and private hospitals go for this directly through oxygen manufacturers on rental basis. It is consumables hence hospital is supposed to buy on regular basis, we do not understand why this item is being purchased and that through by MGPS installation companies, who will have no control on what kind of gas will be supplied, when someone else is liquid tank and someone elseis oxygen gas with this hospital will have to deal with two parties one for maintenance and one for gas and tanksupplier will say problem occurred due to gas supply decantation etc and gas supplier will say gas pressure holding is an issue to due tank and over and above MGPS bidder will be depending upon third party like INOXCVA who is tank manufacturer. By removing this you will not only come out of these problems but save lot of money but also avoid re-occurrence of Gorakhpur incident. It is our duty to make you aware well in advance. Theanotheradvantage is that Estimated value will come down and more and more bidders will be able to participate with lower eligibility criteria which is dependent on Estimated Value.

14, 15 & 16

19 of Volume -IV Supply of Oxygen, N2O and CO2 Cylinders.

Please remove this items "Oxygen, N2O and CO2 Cylinders" because 99% of the government and private hospitals go for this directly through these gases manufacturers on rental basis, It is consumables hence hospital is supposed to buy on regular basis, we do not understand why this item is being purchased and that through by MGPS installation companies, who will have no control on what kind of gas will be supplied, when someone else is liquid tank and someone else is oxygen gas with this hospital will have to deal with two parties one for maintenance and one for gas and tank supplier will say problem

Tender terms and conditions prevail

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occurred due to gas supply decantation etc and gas supplier will say gas pressure holding is an issue to due tank and over and above MGPS bidder will be depending upon third party like Rama or Everest Kanto etc.Cylinders who are Cylinders manufacturers. By removing this you will not only come out of these problems but save lot of money but also avoid re-occurrence of Gorakhpur incident. It is our duty to make you aware well in advance. The another advantage is that Estimated value will come down and more and more bidders will be able to participate with lower eligibility criteria which is dependent on Estimated Value.

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Manufacturer Authorization: Eligible bidders should submit amandatory letter of authority from the Foreign Principal /Manufacturer, mentioning country of origin with name of manufacturing company for major products quoted by them.

For the following major items, Manufacturer's Authorization as per format Volume-II SCC of tender document should be submitted:

- 1. Fully Automatic Oxygen Control Panel
- 2. Oxygen Flow meter
- 3. Fully Automatic Nitrous Oxide Control Panel
- 4. Fully Automatic Control panel for CO2 System
- 5. VACUUM SYSTEMS
- 6. MEDICAL AND SURGICAL AIR SYSTEM
- 7. ALARM SYSTEM
- 8. AREA VALVE SERVICE UNIT
- 9. BED HEAD PANELS
- 10. GAS OUTLETS
- 11. AGSS (Anesthetic Gas Scavenging System)

This is purely a favourable clause to favour companies who have done exclusive tie-ups for Indian market with three to four American companies, who are also not manufacturing these Air Plants, Vacuum Plants and AGSS/WAGD plants themselves but most of them buy from Powerex USA only and they simply print their catalogues with powerex plants specifications and certifications, which has unfortunately never asked or checked by HSCC ever. Ideally there should be an enquiry on this subject. This is an another example of

Tender terms and conditions prevail

restricted conditions to buy only NFPA standard recommendations products and other international standards mentioned in the tender are just eye wash. Under the circumstances this highly objectionable clause may please be removed from this tender and we expect justice at this stage from your end only. We are surprised you being Engineering organization and consultants is not having this information and you did not bother to check online as well before publishing such tenders. Worldwide plants manufacturers are different as they manufacture not only for Medical Industry but for many other industrial requirements too and MGPS manufacturers worldwide do manufacturer these plants except one who are also dealing with Medical and Industrial. You should also not ask single standard for all theitems because it has no relevance, we expect you the explain to us for our knowledge what advantage you get with this and what disadvantage you have without this when the aim of all the standards manufacture products for medical applications and full fill your flow rate, pressure and quality requirements. These are nothing but to favour companies of your choice probably or due to lack of knowledge which both the cases are not expected from government consultants like you.

In view of the facts presented above, we request you to remove this requirement of submission of Manufacturer Authorization Letter from Foreign Principal / Manufacturer, so that bidders should have freedom to buy from any of the international manufacturer and make them to compete with each other which will ultimately benefit you in the price which is ultimate aim of Govt. of India by means of inviting open tender and for your information most of the Govt. organizations like PWD, CPWD do not ask for such manufacturer authorization letter (If you want we can submit the copy of PWD tender in this regards). Under circumstance we should come out of foreign manufacturers' slavery.

Tender and conditions terms prevail

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> The following systems/items must be from the same principal

company/Manufacturer:

- a. Control Panels & Manifold for O2, N2O & CO2
- b. Medical air plant
- c. Medical Vacuum Plant
- d. AGSS Plant
- e. Area & Master Alarm
- f. All types Outlets
- g. AVSU
- h. Line Isolation valves
- i. High Pressure tubes

This is purely a favourable clause to favour companies who have done exclusive tie-ups for Indian market with three to four American companies, who are also not manufacturing these Air Plants. Vacuum Plants AGSS/WAGD plants themselves but most of them buy from Powerex USA only and they simply print their catalogues with powerex plants specifications and certifications, which has unfortunately never asked or checked by HSCC ever. Ideally there should be an enquiry on this subject. This is an another example of restricted conditions to buy only NFPA standard recommendations products and international standards mentioned in the tender are just eye wash. Under the circumstances this highly objectionable clause may please be removed from this tender and we expect justice at this stage from your end only. We are surprised you being Engineering organization and consultants is not having this information and you did not bother to check online as well before publishing such tenders. Worldwide plants manufacturers are different as they manufacture not only for Medical Industry but for many other industrial requirements too and manufacturers worldwide do manufacturer these plants except one who are also dealing with Medical and Industrial. You should also not ask single standard for all the items because it has no relevance, we expect you the explain to us for our knowledge what advantage you get with this and what disadvantage you have without this when the aim of all the standards manufacture products for medical applications and full fill your flow rate, pressure and quality requirements. These are nothing but to favour companies of your

	choice probably or due to lack of knowledge which both the cases are not expected from	
	government consultants like you. Page No.24 of Volume – IV	
60	Point No.6	Bidder shall execute following
60	Bidder shall execute all required civil, electrical, plumbing, lighting, fire safety, exhaust systems and other works as maybe required for complete installation and trouble-free functioning as a part of the 'turnkey work'. Please elaborate all required civil, electrical, plumbing, lighting, fire safety, exhaust systems and other works as maybe required for complete installation. Please specify the areas where these work has to be executed. Please mention quantity of all additional work required to be carried out by MGPS Vendor in tender BOQ.	turnkey works in addition to the works mentioned in the technical specification of tender: -Providing and fixing of Exhaust fan with IS marked Motor and louver for ventilation of MGPS Plant room and Manifold roomOnly Electrical Power supply will be provided at one location inside the Plant room by client. - Air-conditioning (Ductable with exhaust) to run 24x7 inside the Plant room and Manifold room. -Providing and fixing of cable from local alram panels (OTs & ICUs) - Construction of Overhead/Under Ground trench size approx 1.5mx1m as standard for interconnection between buildings/plant/manifold/etc block. - SITC of 3.5 core 185 sq.mm XLPE Cable as per IS: 7098 inside the gas manifold and plant room including Electrical Distribution Panel for plant & Manifold rooms. -SITC of Electrical Distribution Panel for Plant & Manifold rooms. - Providing of dedicated chemical earthing for MGPS Plant room as per IS: 3043 - Wiring for light point/fan point/exhaust fan point/call bell point with 1.5 sqmm FRLS PVC insulated copper conductor single core cable in surface/recessed medium plate, suitable GI box and

		-	3- Point 15	= 2 an	Point = 30, ad Group-C-
61	Point No.6 Bidder will be responsible for trenching or other associated work related to installation and commissioning of complete MGPS system. Since this work is costly, you are requested to kindly mention the same in tender BOQ as extra works.	Tender prevail	terms	and	conditions
62	Point No.19 Bidder should be responsible for dedicated earthing (Chemical type) for MGPS Plant (if required)	Tender prevail	terms	and	conditions
	In the responsibility of bidder, you have mentioned that dedicated chemical earthing for MGPS Plant room is to be provided by MGPS bidder, while the same is not considered in tender BOQ. Since this item includes cost also, we therefore request you to kindly add the same in tender BOQ.				
63	Bidder should submit the MGPS plant and Manifold equipment loading design with footprint of all components as per their offered plant along with bid within the area of 200 sq.m bidder may keep tanks inside. Bidder has to consider proper sitting space for technicians, cylinder storage space for filled and empty including Plant room equipment.	Tender prevail	terms	and	conditions
	Any drawings related to MGPS work will be designed only after issuance of NOA against the Autocad Drawing / Floor Plan of Hospital provided by HSCC / Consignee. Hence it is not possible to submit the same at the time of submission of bid. You are requested to kindly delete the lines stating "Bidder should submit the MGPS Plant and Manifold equipment loading design with foot print of all component as per their offered plant along with bid within the area of 200 sq m. bidder may keep the tanks inside, only when their offered plant and				

	manifold are coming within the 200 sq m area	
	along with proper sitting space for technicians,	
	cylinder storage space for filled and empty	
	including Plant Room Equipment".	
64	Item No.1.1	Fully Automatic Oxygen Control Panel should have digital display
	Fully Automatic Oxygen Control Panel	
	The Manifold Control Panel should be Digital / Analogue	
	Since Analogue is outdated technology and have no comparison with Digital Technology, we recommend amending it to "The Manifold control panel should have 10" LCD Digital Display". Kindly amend the same accordingly.	
	Digital is only specified in all other AIIMS & PMSSY tenders published by Hites& NBCC India Ltd AIIMS tenders.	
	If the requirement is more than flow capacity requirement automatic control panel the bidders has to supply 02 numbers of Automatic Control Panel and design the system in such a way to meet the flow requirement of respective institute.	
	Since the Primary Source for Oxygen is Liquid Medical Oxygen and Secondary source is Automatic Control Panel. Can you please confirm, why there is a requirement of an additional Automatic control panel.	
	You are requested to kindly delete invalid line	
65	from tender specifications. Item No.1.4	BIS/US FDA/European CE
0.5	1001111001.7	Certified with 4 digit notified
	OXYGEN FLOW METER WITH HUMIDIFIER BOTTLE	body number or American ETL/ UL listed.
	Point No. I) – Should be BIS / CE Certified / UL Listed	
	Kindly amend it By mistakenly you have missed to add European CE Marked with 4 Digit Notified Body No / USFDA Certified / UL Listed / ETL Listed	
	Please note same is specified in all other AIIMS	

	& PMSSY tenders published by HITES & NBCC India Ltd AIIMS tenders.	
66	Item No.1.5 LIQUID MEDICAL OXYGEN Mov. Working prossure : 17 Per G	Max. working Pressureis 17-17.6 Bar G
	Max. Working pressure: 17 Bar G Max. Working Pressure is 17.6 Bar G instead of 17 Bar G. Please amend the same.	bai G
	Hydraulic test pressure: 26 bar G Hydraulic test pressure should be as per EN 13458 code instead of 26 bar G. please amend	Hydraulic test pressure should be 26 bar G or as per EN 13458
	the same. Joint Efficiency: 100%	Joint Efficiency 100% or as per code EN 13458
	Joint Efficiency: As per code EN 13458	Tender terms and conditions prevail
	Inspection: By 3rd party (SGS/LLOYDS/TUV) Kindly also add 3 rd party BVIS	
	Page 8, Requirement of the Cryogenic Vessel should be: Inner vessel maximum allowable working pressure: 17 kg/cm2	Maximum allowable working pressure: 17-17.6Bar G
	Max. Working Pressure is 17.6 Bar G instead of 17 Bar G. Please amend the same.	Inner Vessel hydrostatic test pressure: greater than 26 kg/cm2 or as per EN13458
	Inner Vessel hydrostatic test pressure: greater than 26 kg/cm2 Hydraulic test pressure should be as per EN 13458 code instead of 26 bar G. please amend the same.	
	Vaporiser Coil	Tender terms and conditions prevail
	Duty Cycle: Continuous duty	
	Duty Cycle: 6-8 hrs. Kindly amend the same.	
	Safety Fitting Vessel Low Liquid level alarm	Deleted
	In Safety Fitting, Vessel Low Liquid level alarm is not required since low pressure alarm will be provided. Kindly delete Vessel Low Liquid Level Alarm from tender technical specifications.	

67	Item No.2.3	Manifolds are designed and tested
	EMERGENCY N2O MANIFOLD	with aatleast inlet pressure of 3000 psig or as per standards mentioned in the technical
	The Manifold should be hydraulically tested at 3500 psig.	specification of tender.
	Please note as per NFPA standard, Manifolds are designed with a maximum inlet pressure of 3000 psig instead 3500 psig. You are requested to kindly amend the same and mention "The Manifold should be hydraulically tested at 3000 psig".	
68	Item No.3 MEDICAL AND SURGICAL AIR SYSTEM	Shall be as per standard mentioned in the technical specification of tender
	System Control – The control include individual self-protected combination motor controls with short circuit protection, single phase and thermal overload protection, individual control circuit transformers with fuseless primary and secondary protection.	
	As per NFPA Standard, transformers used are fused instead of fuseless. You are therefore requested to kindly amend the same accordingly.	
69	Item No.3 MEDICAL AND SURGICAL AIR SYSTEM	Tender terms and conditions prevail
	Tender Required Total Capacity of Medical & Surgical Air System (combined medical air plant) is 15000 LPM	
	Please allow options:	
	1 complete medical air plant = Total Capacity of 15000 LPM Or 2 medical air plant = Total Capacity of 15000	
	LPM Or 3 medical air plant = Total Capacity of 15000 LPM	

	The total output of 1 or 2 or 3 medical Air plant should meet with tender specifications & Total Capacity requirement.	
70	Item No.4 VACUUM SYSTEM	Shall be as per standards mentioned in the technical specification of tender
	System Control — The control include individual self-protected combination motor controls with short circuit protection, single phase and thermal overload protection, individual control circuit transformers with fuseless primary and secondary protection.	
	As per NFPA Standard, transformers used are fused instead of fuseless. You are therefore requested to kindly amend the same accordingly.	
71	Item No.4	Tender terms and conditions prevail
	VACUUM SYSTEM	
	Tender Required Total Capacity of 28000 LPM	
	Please allow options 1 complete medical vacuum plant = Total Capacity of 28000 LPM Or	
	2 medical vacuum plant = Total Capacity of 28000 LPM	
	Or 3 medical vacuum plant = Total Capacity of 28000 LPM	
	The total output of 1 or 2 or 3 medical vacuum plant should meet with tender specifications& Total Capacity requirement.	
72	Item No.6	
	LOW FLOW WARD VACUUM UNIT Technical Specifications of Low Flow Unit is given in tender, while the same is not considered in BOQ for pricing. You are requested to kindly add Low flow ward vacuum unit in BOQ of	Mentioned in the BOQ attached herewith.
	tender.	Tender terms and conditions

	Please amend the capacity 0-250m/bar for low	prevail			
	flow vacuum regulator Please amend the capacity of suction jar 600- 1000ml (1000cc)	Tender prevail	terms	and	conditions
	The complete low flow vacuum unit with regulator and jar should be European CE Marked with 4 Digit Notified Body No / USFDA Certified / UL Listed / ETL Listed	Tender prevail	terms	and	conditions
	Please note same is specified in all other AIIMS & PMSSY tenders published by HITES & NBCC India Ltd AIIMS tenders.				
73	Item No.				
	WARD VACUUM UNIT				
	Please amend the capacity 0-1000m/bar for low flow vacuum regulator	Tender prevail	terms	and	conditions
	Please amend the capacity of suction jar 600-1000ml (1000cc)	Tender prevail	terms	and	conditions
	The complete ward vacuum unit with regulator and jar should be European CE Marked with 4 Digit Notified Body No / USFDA Certified / UL Listed / ETL Listed Please note same is specified in all other AIIMS & PMSSY tenders published by HITES & NBCC India Ltd AIIMS tenders.	Tender prevail	terms	and	conditions
74	Item No.12.2	Amende	d as Oxy	ygen a	nd Vacuum
	MEDICAL GAS AREA ALARM				
	In Bill of Quantity, for Medical Gas Area Alarm for 2 Services, there is a typographical error in mentioning the gases for 2 Service Alarm, It should be Oxygen and Vacuum instead of MA4 Bar. You are requested to kindly correct the same.				
75	Item No.13	Tender prevail	terms	and	conditions
	LINE ISOLATION VALVES	pievan			
	Kindly add Lines stating:				
	• "Line Isolation Valves should be 3" Piece ball				

	type with Bronze body Lockable with Stuffed Pipes. Each Valve should be separately packed as per the standard.	
	• Valves should be Single Port for 12mm to 76 mm and for 108mm, it should be dual port.	
	This is safety feature and must be added for quality assurance.	
76	Regarding Manufacturer Authorization Ref. Bed Head Panel, you had asked for specific	3 Channel / 2 tier / 2 Channel/3 Partition rows
	authorization for this item, you are requested to kindly clarify whether you need BHP of indigenous or imported make.	
	Bed Head Panels should be 3 channel/3 partition rows 1 channel /1 partition row for gas outlets (pre piped)	
	1 channel /1 partition row for electricals sockets (pre wired)	
	1 channel /1 partition row for Data sockets, nurse call, telephone (not pre wired) This is mandatory feature as per the standards.	
	Please note same is specified in all other AIIMS & PMSSY tenders published by Hites& NBCC India Ltd AIIMS tenders.	
77	Volume-I, Page no. 3,	Period of work completion – 7 Months from the date of letter of
	Period of Completion: 6 Months	commencement
	We request the period of completion should be 9 months after approval of drawings. You would appreciate that quantum of this Project is large and arranging quantity of material takes time and resources. This is not mere supply of equipments items like Touch Screen Control Panel, Surgeon & Anaesthesia Pendant, OT Light with Camera & Monitor etc are imported items for which procurement only starts after approval of final drawing which is a time consuming process.	
70	We hereby request to kindly increase the completion schedule to 9 months.	The empire Configurate C
78	Volume-I, Pre-Qualification Criteria; Page no. 5	The experience Certificate for

& 6, Clause no. 2.2 (ii) completed similar works during last 7 years should be ending last date previous to the date of submission of tender. (ii) Experience of having successfully completed similar work during last 7 years ending last day of month previous to the one in which tenders are invited should be either of the following: We request the successfully completed experience certificate of similar work should from the last date of receipt of application for tender instead of last day of the month previous to the one in which tenders are invited. The reason is this is a prestigious and high value tender. The Experience should be relaxed enough, enabling bidders to arrange & place experience certificate of this magnitude. Therefore it is requested to kindly amend it to "Experience of having successfully completed similar work during last 7 years ending last day of receipt of tender". 79 Volume-I, Page no. 6, Tender conditions and terms 2.2 (iii) prevail Solvency Certificate Considering the estimated cost of the tender, we request M/s HSCC to be liberal & relaxed in terms of value of Solvency Certificate. We request 1 more option should be allowed i.e. Net Worth Certificate from Chartered Accountant. Sir, this qualification criteria is in practice in M/s HLL Infra Tech Services Ltd. Tenders. A copy is enclosed herewith for your ready reference at Page 1 to 3. The criteria should be: Average Net Worth: Eligible bidders should have an Average Net Worth (i.e. Assets minus Liabilities) for the last five years (i.e. from 2013-14 to 2017-18) of not less than 10% of the cumulative estimated value of work to qualify in tender. Please appreciate in this way the bidder has the option to either submit Solvency Certificate or Net Worth Certificate by Chartered Accountant.

> Considering the estimated cost of the tender the bidder should be allowed with an option of Net

	Worth Certificate from Chartered Accountant or Solvency Certificate of 40% estimated cost. Needless to emphasise by this criteria more and more bidders will participate in the tender.	
80	Volume-III, SCC, Page no. 12, 12.4 Bid Security,	Tender terms and conditions prevail
	Document Fee and EMD are exempted for NSIC registered Firm.	
	As you are aware, to promote Micro Small & Medium Enterprise, Government of India had given the facility of NSIC Certification to the manufacturing firm. This may please be noted that this clause is only applicable for manufacturing of goods in India and not for procurement of Imported goods. If we go through the tender Specifications, there are imported items such OT Light, Pendant etc	
	We request this clause should be suitably amended so that NO Bidder could take undue advantage of NSIC Certification and all the bidders should be treated on single platform. This has been done in earlier HSCC Sangrur, Shimla & Raebareli Tenders.	
81	Volume-III, Page no. SCC-38, Clause no. 21.0 Terms of Payment	1) 75% of the BOQ contract rates on delivery of equipment at site after Inspection and Despatch
	For purposes of estimating the contract value of works executed for certificate of payment, the following norms shall be followed:	Clearance Report on pro-data basis. 2) 15% of BOQ contract rates on Installation of MCPS
	1) 70% of the BOQ contract rates on delivery of equipments at site after inspection and passing on pro-data basis.	Installation of MGPS. 3) 10% of BOQ contract rates after final acceptance of system by the client
	2) 20% of BOQ contract rates on satisfactory take over certificate by client after erection and installation, testing and commissioning of equipments on pro-data basis.	
	3) 10 % of BOQ contract rates after successful completion of trial run of 30 days from the date of handover to the client on pro-data basis.	
	We request, the payment terms should be amended as:	

- 1) 75% of the BOQ contract rates on delivery of equipment at site after inspection and passing on pro-data basis.
- 2) 20% of BOQ contract rates on satisfactory take over certificate by M/s HSCC after Installation.
- 3) 5% of BOQ contract rates after successful completion of trial run of 30 days from the date of handover to the client on pro-data basis.

Please appreciate, as soon as the work order is received, the contractor has to initiate necessary actions for successful execution of the work order. Among the very important, Contractor need to place order to the respective suppliers along with 100% payment because supplier will not wait till the completion of the project. Here it is worthwhile to say that contractor do not get 75% payment as 5% amount gets hold for Security from Running Bill; 1% towards Labour 10% towards Performance Guarantee, 12% towards GST & in case of delay in supply liquidated damages. After going through all this in the netshell a contractor gets approximately 50% amount at the initial stage. It is just a eye wash that 70% payment will be released on pro-rata basis. Initial payment is the lifeline to the contractors, which gives relief up to some extent.

With regard to 20% payment, please be noted that commissioning and handing over has no difference. After commissioning by our experience, most of the sites are not ready for handover such as civil work is not complete, hospital staff is not available etc etc. Without any fault of contractor, he need to wait for the payment till the handing over takes place. Therefore this payment should be at the time of erection, installation.

You will appreciate GST tax regime has been implemented since 1st July 2017 by Govt. of India. With the implementation of this system vis-a-vis in current payment structure, most of the projects gets delayed or handing over not taken by the Hospital/Institute/Department; because of this the balance payment gets stuck for longer duration. This way the liquidity get

	11 1 1 1 1 1 1 1				
00	blocked and input credit is lost.				
82	No Deduction from Running Bills We understand, a common ideology & practice has been adopted as that of M/s HLL Lifecare Ltd. in the present tender. While implementing all rules, we request there should not be any deductions with regard to retention of security amount etc.	Tender t prevail	erms an	d cond	itions
83	Part Completion Certificate We request Part Completion Certification should be accepted for Bidder's Evaluation in Tender. As it is project and sometimes due to unavailability of manpower or handover not taken by user it results in delay in commissioning for executed project. For instance, in one of the M/s HLL's Project of J&K Modular Operation Theatre, the complete Installation is done but commissioning is not done from last Four years as the Site is not Clear for Commissioning. Because of this the project remains incomplete.	Tender prevail	terms	and	conditions
	We therefore request Part Completion Certificate should be Accepted in Evaluation.				
84	Performance of the Company We request, to kindly ensure that the bidder who has executed the similar nature of work who has worked in Government Hospital through Government Agencies such as M/s HSCC (India) Ltd., M/s HLL Lifecare Ltd, UPRNN, NBCC etc., the performance of the company should be satisfactory.	Tender prevail	terms	and	conditions
85	We understand, Imported means the place from where the goods are mined, cultivated, grown, manufactured, produced or processed and Outside India. Accordingly the certification of the product applies i.e. if it is UL Listed certification same shall be applicable to American Manufacturer only and European CE Certification same shall be applicable to European Manufacturer only. We request his criteria should be strictly applied and maintained.	Tender prevail	terms	and	conditions

	It is further requested that the European CE Certified/UL listed Criteria for NFPA-99 STANDARD SHOULD BE "CERTIFICATION, WHEREVER APPLICABLE FOR ALL THE ITEMS of MEDICAL GAS".	
86	Page no. 1 & 2 1.1 Fully Automatic Oxygen Control Panel (Imported): Control panel should have Alarm reset switch/Mute /acknowledgement switch to control and monitor the alarm indications by the operator.	Control panel should have Alarm reset/Mute /acknowledgement switches to control and monitor the alarm indications by the operator" is deleted.
	Fully Automatic Oxygen Control Panel is defined as Automatic whereas at the bottom para it is mentioned "reset switch". Both the statements are contradicting to each other. Once the system is Automatic the reset word does not apply. The system will be automatic. All the features are available in Master Alarm this is duplicacy.	
	We request, "Control panel should have Alarm reset/Mute /acknowledgement switch to control and monitor the alarm indications by the operator" should be deleted.	
87	Page no. 2 1.2 Oxygen Manifold Supply System (without Cylinder) The Manifold should be hydraulically tested at 3500 psig.	Manifolds are designed and tested with atleast inlet pressure of 3000 psig or as per standards mentioned in the technical specification of tender.
	Please note as per NFPA standard, Manifolds are designed with a maximum inlet pressure of 3000 psig instead 3500 psig. You are requested to kindly amend the same and mention "The Manifold should be hydraulically tested at 3000 psig". The working pressure is 2000 LPM, we are giving 1.1/2 time more pressure.	
88	Page no. 3 1.3 Emergency Manifold Supply System (without Cylinder) The Manifold should be hydraulically tested at 3500 psig.	Manifolds are designed and tested with at least inlet pressure of 3000 psig or as per standards mentioned in the technical specification of tender.

89	Please note as per NFPA standard, Manifolds are designed with a maximum inlet pressure of 3000 psig instead 3500 psig. You are requested to kindly amend the same and mention "The Manifold should be hydraulically tested at 3000 psig". The working pressure is 2000 LPM, we are giving 1.1/2 time more pressure.	BIS/US FDA/European CE
09	Page no. 3 1.4 Oxygen Flow meter with Humidifier Bottle	BIS/US FDA/European CE Certified with 4 digit notified body number or American ETL/ UL listed
	We request, Oxygen Flow Meter with Humidifier Bottle should be European CE with 4 digit notified no. /UL Listed/USFDA for better quality product.	
90	Page no. 9 & 10 2.1 Fully Automatic Nitrous Oxide Control Panel (Imported): Control panel should have Alarm reset switch/Mute /acknowledgement switch to control and monitor the alarm indications by the operator.	Control panel should have Alarm reset/Mute /acknowledgement switches to control and monitor the alarm indications by the operator" is deleted.
	Fully Automatic Oxygen Control Panel is defined as Automatic whereas at the bottom para it is mentioned "reset switch". Both the statements are contradicting to each other. Once the system is Automatic the reset word does not apply. The system will be automatic. All the features are available in Master Alarm this is duplicacy.	
	We request, "Control panel should have Alarm reset/Mute /acknowledgement switch to control and monitor the alarm indications by the operator" should be deleted.	
91	Page no. 10 2.2 Nitrous Oxide Manifold (without Cylinder) The Manifold should be hydraulically tested at 3500 psig.	Manifolds are designed and tested with at least inlet pressure of 3000 psig or as per standards mentioned in the technical specification of tender.
	Please note as per NFPA standard, Manifolds are designed with a maximum inlet pressure of 3000 psig instead 3500 psig. You are requested to kindly amend the same and mention "The Manifold should be hydraulically tested at 3000 psig".	

92	Page no. 10	Manifolds are designed and tested with at least inlet pressure
	2.3 Emergency Nitrous Oxide Supply System (without Cylinder) The Manifold should be hydraulically tested at 3500 psig.	of 3000 psig or as per standards mentioned in the technical specification of tender.
	Please note as per NFPA standard, Manifolds are designed with a maximum inlet pressure of 3000 psig instead 3500 psig. You are requested to kindly amend the same and mention "The Manifold should be hydraulically tested at 3000 psig".	
93	Page no. 10 & 11 3. Medical and Surgical Air System (Package Unit - Imported)	+/-10 % to flow capacity of plant is permitted
	- Variation of + 10% is missing	
	Please appreciate throughout the worldwide, the Models and the Capacity (LPM) of Air System are Pre-Defined by Manufacturers. Air System is not manufactured as per the requirement. Based on the Pre-Defined Air System, the Models are selected as per the requirement. Like in M/s HSCC (India) Ltd. Tender no. HSCC/SES/MGMS/2018 (IIT Kharagpur) Amendment no. IV dated 02.02.2018; Tender no. HSCC/SES/MGMS/PGI/SANGRUR/2019 Dated 09.01.2019 (PGI Sangrur); Tender no. HSCC/SES/MGMS /SSB/Shimla/2019 Dated: 31.01.2019; Tender no. HSCC/SES/MGMS /AIIMS/Raebareli/2019 Dated: 29.03.2019 variation of +/- 10% is given & like wise and M/s HLL Tenders such as SIX AIIMS for MGPS etc, + 10% variation is given. This +/- 5% / 10% variation is mentioned for ease in procurement.	
	We therefore request, the Air Compressor plant capacity should be defined with variation of +/- 10% and same should be as per Standard.	
94	Page no. 11 Stage 3: Bacteria filter for removing particles down to 0.01 micron.Purity should betested as per the American Pharmacopeia / European Pharmacopeia standard	Testing should be American Pharmacopeia/ European Pharmacopeia/Third Party likeSGS/Lloyd/TUV/ Bureau Veritas

	We request the testing should be American Pharmacopeia/ European Pharmacopeia/Third Party such as TUV etc, which will be more appropriate for bidders.	
95	Page no. 11, 12	The air receiver / vacuum
	3. Medical & Surgical Air System (Package Unit - Imported)	reservoir capacity should be as per standard mentioned in the technical specification of tender.
	Total air receiver capacity shall be atleast 50% (+/- 5%).	
	You have asked 50% standby capacity of air receiver. This should be as per standard. Please amend it to corresponding standards quoted by the bidder'- The capacity should be as per NFPA 99/HTM 02 01 standards as done in Tender no. HSCC/SES/MGMS /AIIMS/Raebareli/ 2019 Dated: 29.03.2019; Tender Enquiry No. TC-1404/GT/Manifold/19-20/FSC [AIIMS Jai Prakash Narayan Apex Trauma Centre, New Delhi] Copy Enclosed at Page no. 4 to 5	
96	Page no. 11, 12 3. Medical & Surgical Air System (Package Unit - Imported)	Air Compressor should be factory fitted, factory tested, packed, prewired & pre-piped.
	We request the Air Compressor (Imported) should be factory fitted, factory tested, packed, pre-wired & pre-piped and tank mounted. As the plants are expensive items and that too imported the genuiness& authenticity of the product should be utmost priority.	
97	Page no. 13, 14	+/-10 % to flow capacity of plant
	4. 4. VACUUM SYSTEMS (Package unit – imported)	is permitted
	- Variation of + 10% is missing	
	Please appreciate throughout the worldwide, the Models and the Capacity (LPM) of Vacuum System are Pre-Defined by Manufacturers. Vacuum System is not manufactured as per the requirement. Based on the Pre-Defined Vacuum System, the Models are selected as per the requirement. Like in M/s HSCC (India) Ltd. Tender no. HSCC/SES/MGMS/2018 Amendment no. IV dated 02.02.2018 (IIT	

	Kharagpur); Tender no. HSCC/SES/MGMS	
	/SSB/Shimla/2019 Dated: 31.01.2019; Tender no. HSCC/SES/MGMS /AIIMS/Raebareli/ 2019	
	Dated: 29.03.2019 variation of +/- 10% is given	
	and M/s HLL such as SIX AIIMS Tender no.	
	HLL/PCD/ PMSSY/AIIMS-II/14-RT-01/15-16	
	dated 31.12.2015 for MGPS etc, +/- 10%	
	variation is given. This +/- 10% variation is mentioned for ease in procurement.	
	mentioned for ease in procurement.	
	We therefore request, the Vacuum plant capacity	
	should be defined with variation of + 10% and	
00	same should be as per Standard.	
98	Page no. 13, 14	Shall be as per standard
	4. VACUUM SYSTEMS (Package unit –	mentioned in the technical specification of tender.
	imported)	specification of tender.
	4.4 Bacterial Filters	
	Bacteria Filters does not come in NFPA-99 Standard The Posteria Filter is in built in the	
	Standard. The Bacteria Filter is in built in the vacuum system. Bacteria Filters comes in HTM	
	Standard. We therefore request to kindly take a	
	note of it and issue necessary amendment.	
99	Page no. 13, 14	The air receiver / vacuum
	4 WACHINA GWOTENIG (D. 1	reservoir capacity should be as per
	4. VACUUM SYSTEMS (Package unit – imported)	standard mentioned in the technical specification of tender.
	Vacuum reservoir shall have total volume of at	technical specification of tender.
	least 100% of Primary plant output (+/- 5%).	
	You have asked 100% (+/-5%) standby capacity	
	of air receiver. This should be as per standard.	
	Please amend it to corresponding standards	
	quoted by the bidder'- The capacity should be as	
	per NFPA 99/HTM 02 01/ DIN standards as	
	done in Tender no. HSCC/SES/MGMS /AIIMS/Raebareli/ 2019 Dated : 29.03.2019;	
	/AIIMS/Raebareli/ 2019 Dated : 29.03.2019; Tender Enquiry No. TC-1404/GT/Manifold/19-	
	20/FSC [AIIMS Jai Prakash Narayan Apex	
	Trauma Centre, New Delhi] Copy Enclosed at	
	Page no. 4 to 5.	
100	Page no. 13, 14	Vacuum System should be factory
	4. VACUUM SYSTEMS (Package unit –	fitted, factory tested, packed, pre- wired & pre-piped
	imported)	whea & pre-piped
	Standby Plant Capacity	
	We request the Vacuum (Suction) System	
1	(Imported), should be factory fitted, factory	

	tested, packed, pre-wired & pre-piped and tank mounted. As the plants are expensive items and that too imported, the genuiness& authenticity of the product should be utmost priority.	
101	Page no. 15 8.0 AGSS Anesthesia Gas Scavenging System (Imported):-	Anaesthesia Gas Scavenging System should be factory fitted, factory tested, packed, pre-wired & pre-piped.
	We request the Anesthesia Gas Scavenging System (Imported), should be factory fitted, factory tested, packed, pre-wired & pre-piped and tank mounted. As the plants are expensive items and that too imported, the genuiness& authenticity of the product should be utmost priority.	
102	Page no. 16 9. DISTRIBUTION PIPING 8.1 Piping specifications	Tender terms and conditions prevail
	We request the Medical Grade Copper Pipe should be Kite Mark. Here, Lloyd is 3rd party Inspection Agency whereas Kite Mark product and service quality certification mark which is owned and operated by the British Standards (BSI Group). It is a voluntary mark of manufacturers and service industries use to demonstrate safety and reliability. The product has been proven to meet the agreed high standard. We therefore request Copper Pipe should be kite marked.	
103	Page no. 17 11. AREA VALVE SERVICE UNIT: The Area Valve Service Unit should incorporate a ball valve with NIST/else connectors either side mounted in a lockable box with emergency access.	Shall be as per standard mentioned in the technical specification of tender.
	The NIST Connection as mention is as per HTM Standard. We work on the Principals of NFPA-99 Standard, where no NIST connection is applicable. We therefore request NIST Connection should be deleted.	
104	Page no. 18, 19 12.1 Master Alarm System The emissions from alarms should conform	Shall be as per standard mentioned in the technical specification of tender.

	with EMC standard	
	with EMC standard	
	We work on the principals of NFPA-99 standard where EMC Standard is not applicable. This standard may be applicable to HTM Standard. Therefore we request you to please delete.	
105	Page no. 19 & 20	All Bed Head Panels shall be Horizontal
	17. Horizontal/ Vertical Bed Head Panel	
	Kindly clarify the no. of Horizontal and no. of Vertical Bed Head Panel required as there is costing involved in it.	
106	Page no. 18, 19 12. Alarm System	Alarms shall be digitalor as per standards mentioned in the technical specification of tender
	12.2 Medical Gas Area Alarm	
	We request the Alarm System should be Touch Type Alarm Technology – Please refer to the Tender no. HITES/PCD/AIIMS-	
	IV/14/MGPS/18-19 dated 14.02.2019 for New AIIMS Gorakhpur and Bhatinda under PMSSY	
	Phase-IV & V and Tender no. HITES/PCD/PMSSY-IV/02/MGPS/18-19 dated 14.02.2019	
	for 7 Medical Colleges /Institutions getting upgraded under PMSSY Phase-IV; where in it is clearly defined DIGITAL. [Tender Papers enclosed for your ready reference].	
107	Page no. 24 13. The Medical Gas Pipe Line System must follow Single Standard any one only from:	The Medical Gas Pipe Line System except accessories like Ward Vacuum Unit, Theatre Vacuum Unit, Flow meter with
	NFPA 99c/HTM02-01/ ISO 7396-1/DIN/EN except Copper pipe.	Humidifier bottle must follow Single Standard any one from: NFPA 99c/HTM02-01/ ISO 7396-
	As mentioned the products should of one standard only; kindly note rest of the medical gas items are Engineering Products. Items such as Ward Vacuum Unit, Theatre Vacuum Unit,	1/DIN/EN including Copper pipe.
	Flowmeter with Humidifier bottle are Accessories and are not part of Medical Gas Pipeline System. The Standard applies from Pipe Distribution to Gas Outlets Worldwide and all the Companies works on this methodology.	
108	Page no. 24 Bidder shall execute all required civil, electrical,	Bidder shall execute following turnkey works in addition to the works mentioned in the technical
	plumbing, lighting, fire safety, exhaust systems,	specification of tender:

false ceiling trap door/ cutout and repair(if any) and other works as maybe required for complete installation and trouble-free functioning as a part of the 'Civil Modification'.

It is requested to Please elaborate all required civil, electrical, plumbing, lighting, fire safety, exhaust systems, false ceiling trap door/ cutout and repair (if any) and other works as maybe required for complete installation. Please specify the areas where these work has to be executed. Please mention quantity of all additional work required to be carried out by MGPS Vendor in tender BOQ

- -Providing and fixing of Exhaust fan with IS marked Motor and louver for ventilation of MGPS Plant room and Manifold roomOnly Electrical Power supply will be provided at one location inside the Plant room by client.
- Air-conditioning (Ductable with exhaust) to run 24x7 inside the Plant room and Manifold room.
- -Providing and fixing of cable from local alram panels (OTs & ICUs)
- Construction of Overhead/Under Ground trench size approx 1.5mx1m as standard for interconnection between buildings/plant/manifold/etc block.
- SITC of 3.5 core 185 sq.mm XLPE Cable as per IS: 7098 inside the gas manifold and plant room including Electrical Distribution Panel for plant & Manifold rooms.
- -SITC of Electrical Distribution Panel for Plant & Manifold rooms.
- Providing of dedicated chemical earthing for MGPS Plant room as per IS: 3043
- -Wiring for light point/fan point/exhaust fan point/call bell point with 1.5 sqmm FRLS PVC insulated copper conductor single core cable in surface/recessed medium plate, suitable GI box and earthing the point with 1.5 sq.mm FRLS PVC insulated copper conductor single core cable etc as required- Group-A - Point = 30, Group-B- Point = 2 and Group-C-Point = 15Revised BOQ attached

In volume V of your tender document (BOQ)
, We request that the stand –by asked for the
Air plant in the BOQ should be as per
International standards(like

Standby in Air Plant should be as per the standardsmentioned in the technical specification of tender.

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Bio int pro Air	pa99,htm0201,din) followed by the dder/Company because different ternational standards have different otocals when it comes to the standby for the r Plants.	The air receiver / vacuum reservoir capacity should be as per standard mentioned in the technical specification of tender.
als dif wh	milarly in Air Receivers the capacity should so be as per the standard followed by the as fferent standards have different protocols nen it comes to the Air receiver capacity for r plants.	+/-10 % to flow capacity of plant is permitted
be con cap	so a range of +/- of 10%, instead of +/- 5% given for plant capacities as different mpanies may have different plant pacities.	
do pla sta	milarly in Volume V (BOQ) of your tender cument, the standby asked for the Vacuum ant is 100% whereas different International andards have different protocols for back	Standby in Vacuum plants should be as per the standardsmentioned in the technical specification of tender.
asl sta	, so our submission is that the standby ked in Vacuum plants should be as per the andards followed by the bidder/Company in se of Vacuum Plants.	The air receiver / vacuum reservoir capacity should be as per standards mentioned in the technical specification of tender.
she con pre	milarly in Vacuum Receivers the Volume ould be as per the standard followed by the mpany as different standards have otocols for the volume of Vacuum ceivers.	+/-10 % to flow capacity of plant is permitted
ins dif caj	so kindly give the range of +/- of 10% stead of +/-5% for the plant capacities as ferent companies have different plant pacities and to reach the capacity asked in a tender documents.	
cla Wi	n page no17 of TS, point no 11, please arify whether the AVSU are with Valves or ithout Valves, otherwise it will have effect the Quantities of the Valves as in the BOQ.	AVSU are with valves
	n page no 22, you have asked Manufacturer athorisation Form certificate for only 11, ms	Tender terms and conditions prevail

	Our Submission is that kindly include Ward Vacuum Unit,Low flow ward Vacuum unit and Theatre Vacuum unit should also be included in the list so that you get quality products of a good standard and there are no compatibility issues among different products.				
113	On page 25 of TS(Technical Specification), point no 15, where it is written that The following systems/items must be from the same principal company/Manufacturer Our submission is that Ward Vacuum Unit, Theatre suction Unit,Low flow Ward Vacuum unit should also be included in the list and they should be from the same manufacture so that there is no compromise on the Quality of the product and also there is no issue with the compatibility among the products.	Tender prevail	terms	and	conditions
114	In Pre Qualification Criteria on page no 5, point no (ii), you have asked that Experience of having successfully completed similar work during last 7years ending last day of month and Three similar* completed works costing not less than the amount equal to 40% of the estimated cost. Or Two similar* completed works costing not less than the amount equal to 50% of the estimated cost. Or One similar* completed work costing not less than the amount equal to 80% of the estimated cost Our submission is that keeping the above criteria only one company will be fulfilling the above Qualification criteria, so to have more companies that will be eligible to bid the tender, the Pre Qualification criteria should be amended to	Tender prevail	terms	and	conditions

	Experience of having successfully completed similar work during last 10 years ending last day of month and Three similar* completed works costing not less than the amount equal to 30% of the estimated cost. or Two similar* completed works costing not less than the amount equal to 40% of the estimated cost. or One similar* completed work costing not less than the amount equal to 50% of the estimated cost.	
115	The time period to complete the Entire work given in the tender documents is 6 months Our submission is that it should be amended to 8-10 months as it is a big work and cannot be completed in 6 months timeline.	Period of work completion – 7 Months from the date of letter of commencement
116	In vol II, clause no 10.1 where in of your tender document, you have asked for 10% of the contract price as Performance Security Our submission is that kindly amend it to 5% of the contract Price as 10% of contract price is a very big amount keeping in view of the Budget for the tenders.	The Performance security shall be 5% of the estimated cost.
117	We also request to kindly amend the payment terms clause 21.0 of volume III of your tender document where it is written that the payment will be released as per 70/20/10 ratio Our submission is to kindly amend the payment terms and it should be released inthe ratio of 80/10/10(80% on supply(pro rata basis),10 % on installation and remaining 10% on handover/Trial run). Also request to extend the dates of the tender atleast by 15-20 days days from the day of uploading of the amendments. Hope our request will be taken into account and the amendments will be uploaded so that	1) 75% of the BOQ contract rates on delivery of equipment at site after Inspection and Despatch Clearance Report on pro-data basis. 2) 15% of BOQ contract rates on Installation (Installation certificate to be provided with bill) of MGPS. 3) 10% of BOQ contract rates after final acceptance of system by the client
	every company has a chance to	

	bid/Participate in the tender process of your prestigious Institute.	
118	Technical specification : Page No. 15 Point no. 8 in Volume IV	Tender terms and conditions prevail
	Anesthesia Gas Scavenging System (Imported)	
	There is 2 different technologies i.e. Rotary Vane / Claw type mentioned in NIT specs which meets only NFPA99 guidelnes. Where as HTM/ISO 7396 standard allows "Blower" technology which is Oil Free produce better output with less energy consumption. Request you to please add "Oil Free Blower Technology" as per ISO 7396 standard. ISO 7396 standard is adopted by latest Indian Medical Gas Pipeline Standard	
119	Technical specification: Page No. 3 Point no.	Tender terms and conditions prevail
	1.4 in Volume IV	pievan
	Flow meter with Humidifier C. The flow meter body should be made of brass chrome plated materials.	
	Request you to please add "Brass Nickle Chrome plated" along with "Brass Chrome plated.	
120	Technical specification: Page No. 19 Point no. 12.2 in Volume IV	Area Alarm should have digital display or as per standards mentioned in the technical specification
	The area alarm should have a digital display of pressures	
	LED display of pressure is allowed under HTM / ISO standard. The main purpose of the alarms is to have a audio visual display in case pressure goes out of the prescribed limits So, LED display can be visualized from disptance	

	and helpful for the nursing staff. So, kindly allow it.				
121	Technical specification : Page No. 11 Point no. 3.1 in Volume IV	Tender prevail	terms	and	conditions
	It should be Oil-Less Screw Compressors / Scroll Compressors				
	Oil less screw compressors are high in capacity which is not suitable & design for the the hospital application. Oil less screw compressors are meant for the industrial application. The Medical Air Plant with oil injected screw compressors are permitted as per ISO 7396 / HTM guidelines and are suitable & design for the hospital application. ISO 7396 standard is adopted in the latest Indian Medical Gas Pipeline standard. So, kindly oil injected screw compressors as per ISO 7396 / Indian Medical Gas Pipeline Standard.				
122	Pre-Qualification Criteria: Page 5, Point 2.2 (ii)	Tender prevail	terms	and	conditions
	Three similar* completed works costing not less than the amount equal to 40% of the estimated cost. or Two similar* completed works costing not less than the amount equal to 50% of the estimated cost. or One similar* completed work costing not less than the amount equal to 80% of the estimated cost.				
	The estimated cost of the project is very high and therefore, the present PQ condition will restrict bidder to qualify for this tender. The financial capability of the bidder can be checked thru other parameter like average turnover clause, a solvency certificate and networth of the company and these parameters are alreday listed in the tender. Many prestigious hospital like AIIMS, Delhi & tendering authorities like HITES has relaxed PQ condition for the other new AIIMS tenders to attract bidders So, we request you to relax it to 10% single order of estimated value and total cumulative 50% of the estimated value				

	project work to be completed during last 7 years	
123	5	Period of work completion – 7 Months from the date of letter of
	1.2 COMPLETION PERIOD	commencement
	6 months from the date of order of commencement	
	Going by the outlets disposition chart of MGPS tender, the OT's are in three different floorsground floor (02 nos), first floor (23 nos) and third floor (02 nos). Since AIIMS projects are done in phases, we request more clarity on readiness of the site to carry out the works within the stipulated time period and accordingly accept PBG to commence work.	
124	Pg. no. SCC 14,	Tender terms and conditions
	Clause 15 (c)	prevail
	The contractor must fill up price in Indian Rupees against each item of BOQ (Volume V) online both in words and figures in the blanks spaces provided in the respective columns.	
	Request that imported items be allowed to be quoted in Euro/US dollar and payment through Letter of Credit be allowed. This will greatly reduce our financial stress.	
125	Pg. No. GCC Page No. 10	Performance security should be 5% of the estimated cost
	10.1 Performance Security	
	The amount of the bank guarantee shall be 10 percent of the Contract Price	
	Request that the performance guarantee be lowered to 5 percent of the contract price since the estimated cost is higher.	
126	Vol. IV Technical Specification Pg. No. 10	+/-10 % to flow capacity is
	3. MEICAL AND SURGICAL SYSTEM	permitted
	Kindly provide tolerance 10 +/-10 % to flow	

	capacity.				
127	Pg. 11 3.1 It should be Oil Less Screw Compressors/Scroll Compressors to produce the plant output of {minimum Liters Per Minutes (LPM)} as mentioned in BOQ of respective institute as primary and same capacity as standby. Kindly include oil free screw compressor /oil free tooth compressor.	Combination or de-combination of Medical Air & Surgical Air system should be followed as pe standards mentioned in the technical specification of tender.			
	According to NFPA99 guidelines, medical air and surgical air must be from two independent sources. Hence please add "If the bidder is quoting as per NFPA 99 then surgical air system and medical air system must be from completely separate sources /compressor systems". This point is necessary since the tender requires third party				
128	Pg. 13 4 VACUUM SYSTEM Primary plant output (+/- 5%) capacity	+/-10 % to flow capacity is permitted			
	Kindly increase tolerance to +/-10%				
129	Commercial criteria: 1. Since most of the items belongs to US/UK/EUROPE origin. Request you to please consider products which belongs to US/UK/EUROPE origin those must be consider in their respective currency.	Tender terms and conditions prevail			
	2. Major products like Air Compressors & Vacuum Systems is a customized product according to client requirement. And these systems dispatches from ''Ship Cargo'' due to highly in larger size. So, we request you to kindly accept 240 days instead of 180 days.				
130	Pre-Qualification criteria: 1. According to NIT, only Indian Experience have been asked. Whereas other India government organizations like HLL/HITES also accept Global Experience.	Tender terms and conditions prevail			

131 Minimum Works of Similar Nature: Eligible Tender conditions terms and bidders should have successfully executed prevail globally in last Seven years from the date of tender opening, similar turnkey project of value, equivalent to or exceeding 50% of estimated schedule/tender value. Out of total 50% value, at least one single order for similar work of minimum 10% of the estimated schedule/tender value should have been executed globally. The details of requirement of cumulative schedule value of MWSN (minimum work of similar nature) are mentioned in Eligibility Table. The value of the executed works shall be brought to current costing level by enhancing the actual value of work at simple rate of 7% per annum, calculated from the date of completion to last date of receipt of tenders. Example/Clarification: Similar projects means that Medical Gas Pipeline System meeting major technical parameters irrespective of material of construction. 132 1. "Air Compressor Modules" As per NIT, Oil free Rotary Screw/Scroll a) Tender terms and conditions compressors only accept. prevail a) Atlas Copco also manufacturers "OIL FREE ROTARY TOOTH" compressors which is smaller version of Oil Free screw. Oil Free Screw requires for larger requirement & Oil Free Tooth designed for lesser demand especially for the Medical Application as compare to Oil Free Screw Tooth. The only difference between both the technologies is Element Size. And this technology is globally accepted. Our sincere request you to kindly refer the attached technical brochure which gives you more glimpse & advantages of the technology. b) Standby Capacity- As per NIT, there is 'standby capacity' is mentioned. b) Standby in Compressed Air specific system should be as per the Whereas every standard i.e. HTM/NFPA99/ISO 7396 has its own calculation for system stand by standardsmentioned in the capacity. Request you to please accept "system technical specification of tender. standby capacity" as per relevant standards. - As per NFPA99 compliance- Secondary should be 25% of primary flow. - As per HTM 0201/ISO 7396-1 compliance- If there are 6 Vacuum Pumps, 2 Pumps should

maintain as secondary.

c) Air Receiver Capacity- As per NIT, this is also specified for one standard. Air Receiver capacity and size must be design as per the relevant standards HTM/NFPA99/ISO 7396. For ex: HTM/ISO says Air Receiver capacity should be minimum of 50% flow whereas NFPA99 says i.e. 5.1.3.5.6 Medical Air Receivers be of a capacity sufficient to prevent the compressors from short-cycling.

The above mentioned tender seeks a declaration from the manufacturer, (in case NFPA system is being offered) that the quoted system conforms to NFPA99 guidelines. The technical specifications of tender demands the Medical Air and Surgical Air be a combined package unit whereas NFPA99 doesn't allow common source for Medical Air and Surgical Air.

In case a combined system is installed, it will fail to comply third party audit to conform to NFPA99.

NFPA 99 2005 edition does not contemplate the use of combined air systems in which Medical Air and Instrument air derive from a single source. This is a violation of a basic principle of the standard which seeks to ensure that no one failure can deny supply, and in this case, it could deny supply to two systems.

- 1. According to NFPA99 edition 2005 para **no. 3.3.80** states that
- 3.3.80 Instrument Air. For the purposes of this standard, instrument air is intended for the powering of medical devices unrelated to human respiration (e.g., surgical tools, ceiling arms). Medical air and instrument air are distinct systems for mutually exclusive applications. Instrument air is a medical support gas that falls under general requirements for medical gas. (PIP)
 - 2. Medical Air and Instrument are different sources as mentioned in **Para no. 5.1.3.3.1.3**
- 5.1.3.3.1.3 Any of the following system shall be permitted to be located together in the same room:

c) The air receiver / vacuum reservoir capacity should be as per standard mentioned in the technical specification of tender.

Combination or de-combination of Medical Air & Surgical Air system should be followed as per standards mentioned in the technical specification of tender.

- 1) Medical air compressor supply sources (see 5.1.3.5.3.)
- 2) Medical-surgical vacuum sources (see 5.1.3.6)
- 3) Waste anaesthetic gas disposal (WAGD) sources (See 5.1.3.7)
- 4) Instrument air compressor sources (See 5.1.3.8)
- 5) Any other compressor, vacuum pump, or electrically powered machinery
 - 3. There are different set of quality air levels as per NFPA99 for Medical Air and Surgical Air Quality.

4. System configuration

The NFPA 99 guidelines defines different system configuration for medical and instrument air system and the guideline states different dew point levels for medical air system and surgical air system. It is self-explanatory that the medical and instrument/surgical air system must be from mutually exclusive and different sources.

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2. "Vacuum Pump Module"

- a) NIT says only one technology that is "Rotary Vane Oil Lubricated". Whereas other technologies are also available which is more energy efficient i.e. "Rotary Screw Oil Lubricated and Dry Vane" Request you to kindly allow these technologies. These kind of pumps delivers higher output in with less motor capacity as compare to Rotary Vane Oil Lubricated. I have enclosed product catalogue explanatory about which is self these technologies.
- b) None of the standards says that Primary system capacity is same as Secondary system capacity. Because as per your NIT Primary Plant output and secondary output is same which is bias the standard norms & regulations.
- As per NFPA99 compliance- Secondary should be 25% of primary flow.

- a) Tender terms and conditions prevail
- b) Standby in Compressed Air system should be as per the standardsmentioned in the technical specification of tender.

	- As per HTM 0201/ISO 7396-1 compliance- If there are 6 Vacuum Pumps, 2 Pumps should maintain as secondary. Vacuum Receiver Capacity- As per NIT, this	The air receiver / vacuum reservoir capacity should be as per standard mentioned in the
	is also specified for one standard. Vacuum Receiver capacity and size must be design as per the relevant standards HTM/NFPA99/ISO 7396.	technical specification of tender.
	For ex: HTM/ISO says Air Receiver capacity should be minimum of 50% flow whereas NFPA99 says i.e. 5.1.3.6.3 Vacuum Receivers	
	be of a capacity based on the technology of the pumps.	
134	pumps	Alarm should be digital or as per
	Master Alarm and Area Alarm Panel According to NIT- only area alarm should have digital display/Touch screen type of pressures. We would request you to kindly accept ''touch screen panel'' only for both Master and Area Alarm Panel.	standards mentioned in the technical specification of tender
135	Page no. 3, Volume-I, Period of Completion: 6 Months	Period of work completion – 7 Months from the date of letter of
	Our Suggestion: The period of completion should be 10 months. The important items of Modular	commencement
	Operation Theatres are imported and takes time to finally arrive in India and being big Project in the history	
	of MOT the period of completion should be atleast 10 months. One should also consider the vicinity of the	
	project. Therefore please extend the completion period to 10 months.	
136	Page no. 5 & 6, Clause no. 2.2 (ii) Volume-I, Pre-Qualification Criteria; (ii) Experience of having	The experience Certificate for completed similar works during last 7 years should be ending last
	successfully completed similar work during last 7 years ending last day of month previous to the one in	date previous to the date of submission of tender.
	which tenders are invited should be either of	
	the following: Our Suggestion: The completed Experience	
	Certificate of similar work i.e. Modular	
	Operation Theatre should be last date of receipt of application for tender. It should not be lest	
	of application for tender. It should not be last day of month previous to the one in which	
	tenders are invited. This is big project in the	
	history of MOT therefore the qualification	
	criteria should be relaxed, so that bidders like us could take part in the tender. Therefore amend	
	experience tolast date of receipt of application	

Page no. SCC-38, Clause no. 21.0, Volume-III, Terms of Payment Our Suggestion: The terms of payment should be: A. 80% of the BOQ contract rates on delivery of equipment at site after inspection and passing on pro rata basis. B. 10% of BOQ contract rates on satisfactory take over certificate by client after erection. C. 10% of BOQ contract rates after successful completion of trial run of 30 days from the date 1) 75% of the BOQ on delivery of equipment after Inspection after I	ipment at site and Despatch on pro-data ontract rates on ation certificate with bill) of contract rates
Volume-III, Terms of Payment Our Suggestion: The terms of payment should be: A. 80% of the BOQ contract rates on delivery of equipment at site after inspection and passing on pro rata basis. B. 10% of BOQ contract rates on satisfactory take over certificate by client after erection. C. 10% of BOQ contract rates after successful on delivery of equipment after Inspection after	and Despatch on pro-data ontract rates on ation certificate with bill) of contract rates
Our Suggestion: The terms of payment should be: A. 80% of the BOQ contract rates on delivery of equipment at site after inspection and passing on pro rata basis. B. 10% of BOQ contract rates on satisfactory take over certificate by client after erection. C. 10% of BOQ contract rates after successful after final acceptance Report basis. 2) 15% of BOQ contract lates on satisfactory to be provided MGPS. 3) 10% of BOQ after final acceptance Report basis. 2) 15% of BOQ contract lates on satisfactory after final acceptance Report basis. 3) 10% of BOQ after Inspection and Clearance Report basis. 4) 15% of BOQ contract lates on satisfactory after final acceptance Report basis. 5) 10% of BOQ after Inspection and Clearance Report basis. 6) 10% of BOQ contract rates on delivery of equipment at site after inspection and passing on provided to be provided after final acceptance Report basis. 6) 10% of BOQ contract rates on satisfactory after final acceptance Report basis. 7) 10% of BOQ contract rates on satisfactory after final acceptance Report basis. 8) 10% of BOQ contract rates on satisfactory after final acceptance Report basis. 9) 10% of BOQ contract rates on satisfactory after final acceptance Report basis. 10% of BOQ contract rates on satisfactory after final acceptance Report basis. 10% of BOQ contract rates on satisfactory after final acceptance Report basis.	on pro-data ontract rates on ation certificate with bill) of contract rates
be: A. 80% of the BOQ contract rates on delivery of equipment at site after inspection and passing on pro rata basis. B. 10% of BOQ contract rates on satisfactory take over certificate by client after erection. C. 10% of BOQ contract rates after successful Clearance Report basis. 2) 15% of BOQ contract Installation (Installation to be provided MGPS. 3) 10% of BOQ after final acceptance for the provided basis. Clearance Report basis. 3) 10% of BOQ and after final acceptance for the provided basis.	entract rates on ation certificate with bill) of contract rates
A. 80% of the BOQ contract rates on delivery of equipment at site after inspection and passing on pro rata basis. B. 10% of BOQ contract rates on satisfactory take over certificate by client after erection. C. 10% of BOQ contract rates after successful 2) 15% of BOQ contract installation (Installation to be provided MGPS. 3) 10% of BOQ after final acceptant	ation certificate with bill) of contract rates
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take over certificate by client after erection. C. 10% of BOQ contract rates after successful 3) 10% of BOQ after final acceptant	
C. 10% of BOQ contract rates after successful after final acceptant	
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of handover to the client on pro-data basis.	
The payment is important part of contract. The	
department should understand, to secure "A"	
payment, a contractor has to incur various	
expenses, such as payment to	
supplier/manufacturer, transportation cost,	
contract expenses such as PBG, Labour Cess,	
Security Charges, GST etc etc. After so much of	
struggle contractor gets payments. Therefore	
change in Terms of Payment is utmost important	
and should be looked upon.	
•	nd conditions
Certificate prevail	id conditions
Our Suggestion: We participate in	
Government/Private Tenders, floated by various	
Departments such as MES,	
PWD etc for similar nature of work. One such	
Department M/s HLL Infra tech service	
ltd'squalification criteria	
is:	
"Average Net Worth: Eligible bidders should	
have an Average Net Worth (i.e. Assets minus	
Liabilities) for the last five years (i.e. from	
2013-14 to 2017-18) of not less than 10% of the	
cumulative estimated value of work to qualify in	
tender".	
This is big project in the history of MOT	
therefore the qualification criteria should be	
relaxed. The Solvency Certificate should be 10%	
of the estimated cost of tender or bidder should	
be allowed to submit Average Net worth	
Certificate for the last five years of not less than	
10% of the estimated cost from Chartered	
Accountant.	
Therefore please amend the criteria for more	
participation.	
Page 2- 1.2 Oxygen Manifold Supply System Manifolds are des	igned with at

	(without Cylinder) - The Manifold should be	least inlet pressure of 3000 psig or
	hydraulically tested at 3500 psig.	shall be as per standard.
	Page 3- 1.3 Emergency Manifold Supply	
	System (without Cylinder) - The Manifold	
	should be hydraulically tested at 3500 psig.	
	Page 10 - 2.2 Nitrous Oxide Manifold	
	(without Cylinder) - The Manifold should be	
	hydraulically tested at	
	3500 psig.	
	Page 10 - 2.3 Emergency Nitrous Oxide	
	Supply System (without Cylinder) - The	
	Manifold should be	
	hydraulically tested at 3500 psig.	
	Our Suggestion: It is requested to please note as per	
	NFPA standard, Manifolds are designed with a	
	maximum inlet pressure of 3000 psig instead 3500	
	psig. Please amend "The Manifold should be	
	hydraulically tested at	
	3000 psig".	
140	Page no. 10 & 11 - 3. Medical and Surgical	+/-10 % to flow capacity is
	Air System (Package Unit - Imported)	permitted
	Page no. 13, 14 - 4. 4. VACUUM SYSTEMS	
	(Package unit – imported)	
	Our Suggestion : We request +/- 10% variation	
	should be provide as in all Government Tenders	
	variation is	
	allowed.	
141	Page no. 11, 12 - 3. Medical & Surgical Air	The air receiver / vacuum
	System (Package Unit - Imported) - Total air	reservoir capacity should be as per
	receiver capacity	standard mentioned in the
	shall be atleast 50% (+/- 5%). Page no. 13, 14 - 4. Vacuum Systems (Package	technical specification of tender.
	unit – imported) - Vacuum reservoir shall	
	have total	
	volume of atleast 100% of Primary plant	
	output (+/- 5%).	
	Tangan () Tangan	
	Our Suggestion: The air receiver / vacuum	
	reservoir capacity should be as per standard	
	HTM 02 01/NFPA-99	
	Standard.	
142	Page no. 18, 19 - 12. Alarm System 12.2	Alarm system shall be digitalor as
	Medical Gas Area Alarm	per standards mentioned in the
		technical specifications of tender
	Our Suggestion: We suggest the alarm system	document.
	should be digital as this is the latest	
1.40	technology.	Digata ED / 7
143	Page no. 3 - 1.4 Oxygen Flow meter with	BIS/US FDA/European CE

	Humidifier Bottle - Our Suggestion: The certificates of the product	Certified with 4 digit notified body number or American ETL/UL listed
	should be European CE Certified with notified 4 digit body no. / UL Listed.	
144	Quality standards-US FDA/European CE with 4 digit notified body no./ETL/UL etc for quoted model	Quality standards-BIS/US FDA/European CE with 4 digit notified body no./ETL/UL etc for quoted model
145	Operationand Maintenance of MGPS	The Estimated cost of MGPS Project is excluding of Cost of Operation and CMC Agreement for Operation and CMC shall be made between AIIMS and the contractor. AIIMS will award for Operation and CMC and make payment accordingly for the same. In addition to technical specification, details of manpower planning and management of Operation and Maintenance are mentioned in the attached sheet. Payment for Operation shall be half yearly basis after satisfactory completion of operation of the said period Revised BOQ attached.

All other terms & conditions remain unchanged.

Sr. Chief General Manager -I , HSCC (I) Ltd. For & on behalf of Director, AIIMS, Nagpur

Added part under Technical Specification

Operation and Maintenance of Medical Gas Pipeline system at AIIMS

Scope of the work:

The bidder should ensure safe and reliable MGPS and their efficient Operation and use as per standards. Bidder will be responsible for operation and maintenance of MGPS as following:

- Medical oxygen System -Liquid oxygen system , Manifold and Control panels
- Nitrous oxide System-Manifold and Control Panel
- Medical and Surgical Air System-Compressor systems , Control panel, Dryers, Reservior, Filters etc
- Medical Vacuum System-Vacuum pumps, Control panel, Reservoir, Filters etc.
- Waste anaesthetics gas scavenging systems (AGSS)
- Carbon dioxide manifold system
- Copper pipelines
- Area Valve Service Units
- Isolation Valves
- Area Alarm panels and Master alarm panels
- Gas Outlets
- Bed Head Panels
- Pendants (If any)

Staff responsible for plant operation should be aware of the activities necessary to ensure the continuous safe operation of the system and action necessary to be taken in an emergency. The authorised person of MGPS provider in particular should take a lead in explaining to users the function of the system and should be adequately trained and informed about the system. Operator shall be responsible for safe cylinder handling, storage and transportation. Any work involving alterations, extensions or maintenance work on the system should be subject to the permit-to-work procedure as per standards.

Operation of Medical Gas Pipeline System

The contractor should provide manpower to operate the plant 24 x 7 and 365 days in a year. The duty of the worker in each shift should be limited to 8 hours per day.

Sl.	Designation	Shift-	Shift-	Shift-	General	Leave	Total
No.		A	В	C	Shift	substitute	
1	Supervisor (Diploma in Mechanical/Electrical)With 5 years' Experience in installation maintenance & operation of MGPS				1		1
2.	Medical Gas Technicians (ITI)With 2year Experience in installation maintenance & operation of MGPS	1	1	1		1	3
3.	Helpers (8 th Standard or more with minimum 5 years' experience in installation, maintenance & operation of MGPS	2	2	2			6

The operators should ensure a trouble free seamless supply at the outlets at the required pressure. They should Monitor the consumption of O2& N2O on hourly basis and submit a consolidated report weekly, Timely intimation of cylinders refill due date, Timely intimation of oxygen plant refill due date based on consumption, and other service maintenance has to be done by the operator.

The contractors and operator should be fully aware of the safety regulation applied to Medical gas System. It is the responsibility of the contractor to conduct training sessions of adequate level to the workforce periodically to keep them fit for handling the plants and associated systems. All tests to be conducted by authorized persons, competent persons, quality controller etc have to be arranged by the contractor additionally as required. The contractor may refer to relevant part of standards for details.

Routine Activity

1. Oxygen plant

Checking oxygen pressure and liquid level
Entering details in the log book
Checking for leaks
Checking the change over
Intimating the preventive Maintenance one week ahead of the schedule
Supervising Maintenance jobs and checking reports

2 Manifold (Oxygen and Nitrous oxide)

Inspection of following
Checking for leakage
Checking inlet and outlet pressure
Checking the change over
Loading the cylinder as required
Replacement of defective parts
Notifying breakdown
Logging details

3 Compressed Air

Checking change over
Checking pressure
Checking the dryer and change over
Checking the receiver
Checking the filter

4 Medical Vacuum system

Checking vacuum pump
Checking vacuum level
Checking controls
Checking change over
Checking for drop in Vacuum level
Checking the filters

5 Medical Gas Lines

Checking for leakage Checking the isolation valves Checking the vacuum lines for block Checking alarms Replacing leaking lines

6 Bedhead Panels, Gas outlets, Pendants

Checking for leakage Checking for defective valves Replacement of defective parts