AMENDMENT - II

Dated:- 10.1.2014

Subject: Amendment to the tender Enquiry Document.

Ref: Tender Enquiry No.: HSCC/PUR/AIIA/HOSPITAL EQUIPMENT/2013

The pre-bid meeting for the referred tender enquiry was held on 10.12.2013 & 11.12.2013. The following amendments are being incorporated in the referred tender enquiry document.

For:

Package - 60

1	Digital Mobile X-Ray Unit		1	300,000	
2	500 mA	Digital	Fluro	1	
	Radiograph	y System			

Read As:

Package - 60 (A)

1	Digital Mobile X-Ray Unit	1	
			1,60,000

Package - 60 (B)

1	500 mA High Frequency X-	1	1,40,000
	Ray unit with Image		
	Intensifier		

PACKAGE NO. 37 SINGLE PUNCTURE LAPARASCOPE

Point No.	Existing Tender Specification	Amendment requested by Bidder	Amended by AIIA / HSCC
3.3	Full High Definition(HD) Endoscopic camera with T.V. medical grade monitor and printer A.2)Endoscopic High Definition Camera (Digital) 1. 3X1/3 CCD image sensor. 2. Should have progressive scanning and should support 16:9 format 3. Should have option of controlling the compatible endoscopic units in hands of surgeon/ touchscreen 4. Should be compatible with 23- 26 inch monitor 16:9 HD format 5. Upgradeable 6. Resolution should be 1900 x 1080p or more 7. Light weight camera head with programmable function key 8. PAL system/ multimedia as existing in this country 9 Automatic white balancing 10. Freely programmable camera head buttons 11. Cable should have buckling protection 12. Facilities for fine focus for smooth	Bidder	Point 3.3 stands Deleted .
	function. Microprocessor controlled. 13. Built in antifogging device.		

1	. Camera head should be compatible	
W	th telescope of any make and light of	
a	y make.	
1	. Integrated universal power supply	
1	. Compatible with medical grade	
n	onitor with multimedia projection	
a	ailable in this country.	
1	. Should have specific built in facility	
fo	camera functionality automatically	
0	timizing all settings	
1	. Camera should be ready to use as	
S	on as it is connected to camera	
C	ntrol unit.	
1	Universal coupler	
2	. Inbuilt electronic Fibre optic filters	
В	2) Camera Control Unit	
1	Should have microprocessor control	
2	The Camera CCU should be capable of	
e	her down-converting HD signals to SD	
	up-converting SD signals to HD.	
3	It should have provision of working /	
	mpatible with lower models of	
	mera heads.	
	. It should allow images from one	
	rmat to be viewed, on displays in	
	ferent format ie it is the HD system is	
	mpatible with both SD and HD.	
	Should have multiple video input and	
	t puts – BNC,RGB,Y/C, DVI-D	
	cket,digitalSDI signal, DV for digital	
	cording etc	
6	Should have all necessary connecting	

cables between camera	ead and video	
monitor	icaa ana viaco	
monitor		
C. 2 HD MEDICAL GRADE	MONITOR flat	
screen,LCD/LED/ TFT MC		
1.Desktop or wall mount		
2. Multinorm/PAL system		
for different color system		
the country.	3	
3. Compatible with endo	ision camera	
of any makes		
4. Screen size diagonal 23	/24/26" Ultra	
high resolution, more the		
5. Aspect ratio 16:10		
6. Should preferably hav	advanced	
technology feature to pe	form interlace	
to progressive conversio		
7. Number of colors show	ld be	
approximately 16.8 million		
8. Viewing angle should	e wide	
9. Monitor menu display		
capabilities and operation		
keys, user defined caption	ns, easy to use	
and highly dependable.		
10. Should be composite	•	
video input and out puts	– BNC, RGB,	
Y/C, SDI, DVI etc		
11. Power supply of 200-	240 VAC. 50	
/60Hz		
12.Should have facilities	•	
the data on computer /d	gital Video	
recorders/CD		
13. On screen menu for	nonitor setting	

, Compact and light weight ,Drip water	
protected dust proof, all connecting	
cables to be supplied	
14. Brightness 400cd/m2, contrast ratio	
1000:1	
15. Antireflection quoted front glass.	
16. Should have consistent illumination	
level.	
17. Should preferably have facility for	
upgradation and should be compatible	
with lower models.	
18. Should be supplied with power	
supply, monitor stand and mains cord	
19. Camera, CCU, & Monitor should be	
compatible with each other and	
preferably should be of same make.	
D) Documentation system for storage	
and transfer of digital data	
1- Digital storage of still HD images and	
video/ audio files. It should have the	
facility editing/cutting of recorded data.	
2 Auto detection of the connected	
camera system on HD_SD/ SD-SDI input	
5 Archiving on DVD CD- ROM or USB	
stick, Multi- Session and Multi –Patient	
6 Network saving	
7 Automatic generation of standard	
reports Approved use of computers and	
monitors in the or environment as per	
60601- 1	

PACKAGE NO. 37(2 Operative Gynaecological Laparoscope Set)

Point No.	Existing Tender Specification	Amendment requested by Bidder	Amended by AIIA / HSCC
Point -3.4	Full High Definition (HD) Endoscopic camera with TV medical grade monitor and printer	Apart of existing features, kindly add following important point also i. Three Chip Camera head should produce at head itself Pure Digital signal with high definition video (1920*1080P) with aspect ratio of CCD chip and video format of 16:9 or 16:10 ii. Should have integrated Digital/optical zoom lens 15-32mm + 10%zoom range, to increase and decrease the size of image which should remain in focusing zone, without readjusting the	Amendment requested by Bidder stands DELETED .

focus and fully soakable. iii. Should have port for direct recording of still & video sequences On external hard drive
of 1TB and 2 Nos. of this external hard drive to be supplied along with camera. iv. All camera
functions to be controlled from camera head buttons or through key board at Camera
control unit to make it controllable from both sterile and non sterile zone. The camera should also
comply with the following standards: According to :IEC 60601-2- 18, UL2601.1 CSA22.2 NO.601.1-M90:
- Type of protection against electrical shocks: Protection class

- Degree of protection against electrical shocks: Applied part of type CF defibrillator proof According to Medical Device Directive (MDD) the camera should belong to Class I and bear the CE mark in accordance with MDD 93/42/EEC
Should have part to connect to communication data BUS.

PACKAGE NO. 60 (ITEM 1 DIGITAL MOBILE X RAY)

Point No.	Existing Tender Specification	Amendment requested by Bidder	Amended by AIIA / HSCC
2. Generator	vi. Shortest Exposure time: Should be 1ms or less	Shortest exposure time should be less than 4ms or less.	Should be 4ms or less
II of 2	Generator It should have digital display of mAS and kV and digital timer	It should have digital display of mAS and kV	No Change
III of 3	X-ray Tube It should have dual focus. Large focus :1.3mm and small focus 0.6mm or better.	It should have single/dual focus. Having 0.8 or better in case of single focus. Large focus :1.3mm and small focus 0.6mm or better in case of dual	No Change

		focus.	
4.Flat Panel Detector System	I. Minimum size of detector must be 14" x 17"	flat Panel Detector of size at least 16" x 16" size.	No Change
	IV. Pixel size / pitch should be 160µm or less	Pixels size should less than 200µm	No Change
IV of 5	The battery should be able to charge from a normal 15A 220-240 V single phase socket in less than 6hrs, preferably.	When fully discharged, the battery should be able to charge from a normal 15A 220-240 V single phase socket in less than 12hrs, preferably.	The battery should be able to charge from a normal 15A 220-240 V single phase socket not more than 12 hrs, preferably.
Point No.4- Flat Panel detector		Request you to kindly specify the detector scintilator whether it should be CSI or GOS Also as you are looking for 63% DQE so this is only possible with CSI and since DQE cannot be measured locally. So request you to kindly specify the detector as CSI only.	detector scintilator should be CSI
	A grid of ratio 10:1 of appropriate size preferably 17" x 17" should be supplied	Size of Grid should get changed to 14" x 17" (As detector of size 14" x 17" has been asked therefore grid of same size will be provided.	14" x 17"
	Vendor will get approval for the site plan from AERB for installation of the equipment	Should get DELETED NOT REQUIRED FOR MOBILE UNIT.	Site plan approval not requried

V of 2	It should be capable of delivering	It should be capable of	It should be capable of
	upto 300mAS in different steps.	delivering upto 200mAS in	delivering upto 200mAS in
		different steps.	different steps.
14.	Company/ supplier should have	Quoted model should have US	Quoted model should have US
	CE/FDA approval certificate and	FDA & CE certified. And quoted	FDA or European CE certified.
	quoted model should have	model should have AERB type	And quoted model should have
	AERB type approval.	approval	AERB type approval
Point No.14	- Company / Supplier should have	The quoted model must have	The quoted model must have
	CE/FDA approval certificate and	CE, USFDA and AERB type	European CE or US FDA and
	quoted model should have AERB	approval- this will lead to best	AERB type approval
	type approval.	technical specification for the	
		required product.	

PACKAGE NO. 60, Item No.2 (500mA Digital Fluro Radiography System).

Point No.	Existing Tender Specification	Amendment requested by Bidder	Amended by AIIA / HSCC
		Kindly add that x-ray tube should be USA FDA approved and manufactured in same country where main equipment is manufactured	Not required
7	Dual Focus X-ray Tube with Large focus 1.0mm and small focus 0.6 mm or smaller.	should get changed to 1.2mm	Dual Focus X-ray Tube with Large focus 1.2mm or less and Mention size of small & large Focal Spot.

		0.6 / 1.2mm.	
20	Under table 12 inch image	PLS Delete 12 inch Image	12" or more
	intensifier system with high		
	resolution CCD camera.	Under table 16 inch Triple field	-
		image intensifier systems with	
		high resolution 1 k CCD	
		Camera.	

PACKAGE NO61. CT SCAN

Point No.	Existing Tender Specification	Amendment requested by Bidder	Amended by AIIA / HSCC
X-ray Section.2:	The System X-ray power should be 70 kw and above	The System X-ray power should be 100 kw and above.	The System X-ray power should be 100 kw and above.
			Point to be added: Dual Energy Applications Application (Should be FDA approved) 1) Calculi Characterization 2) Gout Characterization 3) Monoenergetic imaging
		Oncology Application	Point to be deleted

PACKAGE No. 68: ANESTHESIA WORKSTATION

For:

Should be US- FDA approved product

Read As:

Should be US FDA or CE approved.

All other terms & conditions of bid document shall remain unchanged.

Director (AIIA)