Date: - 27th November, 2019

Corrigendum-2
for
4K Laparoscopy Imaging System with ICG compatibility for the Department of Surgical Gastroenterology

NIT Issue Date	**:**	**21st August, 2019**
NIT No.	**:**	**Admn/Tender/39/2019-AIIMS.JDH**
Pre-Bid Meeting	**:**	**02nd September, 2019 at 03:00 PM**
Earlier Last Date of Submission	**:**	**28th November, 2019 at 03:00 PM**
Extended Last Date of Submission	**:**	**16th December, 2019 at 03:00 PM**
Bid opening	**:**	**17th December, 2019 at 03:15 P.M**

The following revised and additional specification will be added:-

1. **In Corrigendum, Page no. 2, Point no. 2 , Point a)**

For

A Powerful 300 Watt Xenon Lamp or equivalent LED light source.

Read as:

Two powerful 300 Watt or more Xenon or equivalent LED light source to provide 4K and ICG output for independent working.

2. **In Corrigendum, Page no. 2, Point no. 2 , Point e)**

For

300W Xenon light source with spare lamp (10 extra bulbs) having " NBI technology" / ICG technology

Read as:

300W Xenon light source with 5 spare bulbs having "NBI technology" / ICG technology

3. **In Corrigendum, Page no. 5, Point no. 8 , Point (b), "Image and Video recording and data archiving management system":**

For

Should be a 4K UHD / full HD medical grade video and still images recorder

Read as:

Should be able to record 4K High definition medical grade video and still images recorder

4. **In Technical specification, Page No. 12, Point No. 8, sub-point No. (f)"**

For

Should have dual channel simultaneous recording for 4K UHD /full HD

Read as:

Should have dual channel simultaneous recording for 4K High Definition images and videos.

5. In Corrigendum, Page no. 5, Point no. 8 , Sub-Point (O), "Image and Video recording and data archiving management system":

For

Should have support for physician print, media, annotations, patients field, procedure Settings and individual surgeon profile & preferences.

Read as: Optional

6. In Corrigendum, Page no. 5, Point no. 8 , Sub-Point (r), "Image and Video recording and data archiving management system":

For

Following accessories should be supplied: Cordless mouse and cordless silicone keypad and foot pedal control

Read as: Optional; provide if available.

7. **In Corrigendum, Page no. 6, Point no. 9:**

For

All products should be same manufacturer & should be US FDA or European CE approved (with four digit notified body number)

Read As

All products should be from same manufacturer & must be US FDA or European CE approved (with four digit notified body number).