


Terms & Condition:

- 1. The bidders who unable to submit the complete requisite documents (as per Annexure – I) considered as technically disqualified.**
2. Firm to mention Make/Brand name in their quotation.
3. Taxes, if any (Kindly mention in above table) should be clearly mentioned in the offer.
4. Document relating to registration of firm i.e. GST and relevant document should be submitted along with quotation.
5. Products are certified from ISI/ISO/CE/GMP/BIS as applicable, the Certificate to this effect should be attached.
6. Supply should be done within 15 days after Placement of PO.
7. Price should be FOR Destination basis (i.e. concerned department).
8. 100% Payment will be released after certification from concerned department.
- 9. Quotation Name/No. and due date of opening must be mentioned on top of envelops.**
10. LD @ 0.5% of delayed supply per week or part of week for delay of supply of material subject to maximum up to 10% of delayed supply to be deducted.
11. AIIMS Raipur reserves the right to place order for full or part quantity to one or more firms. The AIIMS, Raipur reserves the right to increase/decrease the number of required quantity.
12. All other terms & condition as per GFR 2017.
13. Material to be delivered at **Obstetrics and Gynecology Department and B-Block OT, AIIMS Raipur.**
14. **Validity of the quotation should be 90 days from the date of opening.**
15. **Bidders should submit their email id, contact details with GST registration.**
16. **AIIMS Raipur reserves the right to ask the tenderers for arranging demonstration of their samples for feel & amp; finish for which rates have been quoted, to the concerned committee, if required.**


Officer In-charge

Procurement (DMC)

AIIMS, Raipur (C.G.)

Dr. Yogendra N. Keche
(Officer In-charge, Procurement of
Drugs, Medicine & Consumable)
AIIMS, Raipur (C.G.)

Annexure - I

Technical Specification

- Pouch Rolls having inbuilt chemical indicator intended to be used to enclosed medical devices that are to be terminally sterilized in Sterrad NX 50L/ 100L Sterilizer.
- Pouch Roll material must be validated by appropriate authority for use in Sterrad NX 50L/ 100L sterilizer for terminal sterilization already installed in our institute.
- Validation certificate for demonstrated Seal strength test, microbial barrier properties, burst test, peel open test after processing in the sterad NX 50L/ 100L sterilizer must be produced from appropriate authority.
- Validation certificate for demonstrated Sterilant penetration and package integrity after processing in the sterad NX 50L/ 100L sterilizer must be produced from appropriate authority.
- Validation certificate for demonstrated maintenance of sterility for 12 months post processing in the sterad NX 50L/ 100L sterilizer must be produced from appropriate authority.
- As matter of terminal surgical instrument sterilization is very serious and involves with patient safety; all of the above claims by manufacturer must be validated from appropriate authority like CDSCO-India or equivalent Indian Authority/ US-FDA/ European CE for the intended use in Sterrad NX 50L/ 100L Sterilization system installed in our institute.
- At-least one year of shelf life of the delivered material from the date of delivery.