BBQR OF MEDICINES

- 1. Quoted manufacturer should have WHO/GMP Certificate.
- 2. Should have valid Drug Manufacturing Licence No
- 3. Should have Marketing Manufacturing Certificate (MMC) if drug manufactured in India.
- 4. The sellers are submit notarized undertaking as per format attached.
- 5. Annual Turnover certificate more than Rs. 20.00 Crore per year (for the last 3 Year) of the manufacturer/firm quoted, certified by CA or VAT Form 21/22.
- 6. **OEM CERTIFICATE**:- A scanned copy of valid authorised letter of OEM or authorised marketing agent (if any) on letter head addressed to Commandant, Command Hospital (WC) Chandimandir uploaded in PDF. Authorised signatory should mention name, contact number & E-mail.
- 7. Certificate of acceptance of terms & conditions of GeM on firm's letterhead. Undertaking that firm has not been blacklisted and undertaking for recovery of cost of entire quantity ordered ad acceptance of punitive action as per existing DGQA policy in case of item is declared toxic/defective
- 8. Compliance statement including item wise manufacturing company quoted for and if authorization for the same attached, (as per appx B). The compliance statement should be in proper format stating all the requisite details, or else the entire bid will be rejected.
- Valid PAN/TIN/GST/VAT registration certificate.

10. Primary package - tablets to be packed in strip/blister as applicable.

Dated: ♥ Aug 2024

(Bishnu Raj)

Lt Col

OIC MSPC

COUNTERSIGNED

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COMPLIANCE CERTIFICATE

Details	and the committee of th	entervador (tedpetiro, escuestrator anti-se vivalmentratification dis-se	and the second section of the second		
Name of the firm and the address with telephone number: (As given in Drug License)			de description of a section of the section of		
Drug License No. No.	in form 25 & 28 (OR Import License		general and the state of the st	ggint to the second state of the second seco
Manufacturer of the item offered	Compliance to RFP specification. Whether Yes/No	compliance deviation from	ons IP/BP/US		Is the manufacturer as per DPM 2009? i.e Has the the Complied with the Compliance sheet as per Appx 'G'
	Name of the firm number: (As given in Drug I Drug License No. No. Manufacturer of	Name of the firm and the address number: (As given in Drug License) Drug License No. in form 25 & 28 (No. Manufacturer of the item offered RFP specification.	Name of the firm and the address with telephone number: (As given in Drug License) Drug License No. in form 25 & 28 OR Import License No. Manufacturer of the item offered Compliance to RFP specification. Whether Yes/No RFP to be specified in unambiguous	Name of the firm and the address with telephone number: (As given in Drug License) Drug License No. in form 25 & 28 OR Import License No. Manufacturer of the item offered Compliance to RFP specification. Whether Yes/No RFP to be specified in unambiguous	Name of the firm and the address with telephone number: (As given in Drug License) Drug License No. in form 25 & 28 OR Import License No. Manufacturer of the item offered Compliance to RFP specification. Whether Yes/No Whether Yes/No No. In case of non-compliance deviation from RFP to be specified in unambiguous terms FDA/CE Whether Yes/No Whether Yes/No Whether Yes/No No. Whether Yes/No Whether Yes/No No. Whether Yes/No No. In case of non-compliance deviation from RFP to be specified in unambiguous terms No. Whether Yes/No No. No. No. No. No. No. No.

I, the u	inderstand do solemnly affirm: -
(a)	That I am the proprietor/partner/agent/of M/S
(b) firm to	That the rates quoted for the items are either at per or lower than that as quoted by the any other Hospital and Institution.
(c)	Rates are lower than MRP of items.
(d)	There will be been no failure in delivery, compliance or quality of Drug (s)supplied.
Place:	Authorized Signatory
Date:	Name: