Request for offer 5/2026

Requirements:

Source of Medicines:

- All medicines and their manufacturers should be registered in the National Medicines & Poisons Board (NMPB). \ Medicines with narrow therapeutic index, high toxicity, biological products, serums, vaccines must be from their Originators, Manufacturers holding WHO Prequalification status for specific products or from countries with Stringent Regulatory Authorities (SRAs) determined by NMPB.
- The NMSF may, where necessary, procure unregistered medicines or pharmaceutical products under the following conditions and subject to prior written approval of the National Medicines and Poisons Board (NMPB):
- The products must originate from countries with Stringent Regulatory Authorities (SRAs) as determined by the NMPB. International institutions with globally recognized competence in medical supply.
- Manufacturers holding WHO Prequalification status for specific products.
- For consumables, bidders must submit a valid certificate of registration for the manufacturer and for each item. In cases where the items are not registered, an official No Objection Letter for importation issued by the NMPB must be provided. Furthermore, all relevant supporting certificates must be submitted, including but not limited to: Free Sale Certificate (FSC), ISO 13485:2016, EC Certificate (as applicable to the device classification), USFDA approval, and/or a valid registration certificate issued in one of the following countries: the European Union, Japan, Australia, Republic of Korea, Canada, the Kingdom of Saudi Arabia, or Brazil
- For consumables, three non–returnable samples, including different sizes for the single item must be provided, during or before the closing date of the Tender. Each sample must be labeled with item code, serial number and the name of the local agent
- A colored, clear and readable artwork or clear photo from all sides must be submitted, and goods to be supplied must be strictly in accordance with the original artwork.
- All labeling requirements that have been stated by NMPB policy for registration of pharmaceutical products.

- Commercial invoice signed and stamped in full details, items names must be written in generic names with full specifications, expiry date, manufacturer name, country of origin, storage condition.
- Packing list must be submitted with commercial invoice, with full information and with details of quantities of each batch, number of cartons, pallets, batch numbers, manufacture and expiry dates.
- A copy of the commercial invoice is submitted to NMSF Madani & Khartoum, and the original must be submitted Khartoum for payment.

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