بسم الله الرحمن الرحيم

القيادة العامة للقوات المسلحة الأردنية – الجيش العربي
مديرية الخدمات الطبية الملكية
شعبة المشتريات المركزية

الرقم : ش/٧٠٠/١/٢٠٢٥ / ٩٥٥٥
التاريخ : ١١ / محرم / ١٤٤٧
: ٦. / تموز / ٢٠٢٥

السادة : المناقصون
الموضوع : تعديل المواصفات

تحية وبعد ،،،

الاشارة : العطاء رقم ش/٧٠٠/١/٢٠٢٥
شراء (احتياج وحدة ترميم الجمجمة والوجه )

- مرفق طيه التعديل والتوضيح لبعض نقاط المواصفات المطلوبة للعطاء أعلاه وحسب ما وردنا من الجهة المعنية .

واقبلوا فائق الاحترام،،،،

ع / العميد الطبيب
مدير عام الخدمات الطبية الملكية
عقيد صيدلاني طارق محمد الجبوري

جهاز رقم (١)

| التعديل | اسم المادة | رقم المادة |
|---|---|---|
| Supercut, serrated, 18 cm length | Kaye scissors | 4 |
| Standard 17.5 cm length | Mitchells trimmer | 7 |
| 14 cm length, 30° angle | Dandy artery forceps lateral curved | 8 |
| Size:<br>10mm, 15mm, 20mm, 25mm,<br>30 mm width and 20 cm length | Malleable brain re-tractor set (all sizes) | 9 |
| Length 18 cm. | Microsurgery set | 12 |

JORDAN ARMED FORCES

The DIRCETROATE OF ROYAL MEDICAL SERVICES

THE INSTITUE OF BIOMEDICAL TECHNOLOGY

| Item 1 | Syringe pump, PCA | Qty. (3) |
|---|---|---|

**IMPORTANT NOTE:**

*Vendors are required to fill out the product details, compliance information, and brochure reference page number in the columns below*

**TECHNICAL SPECIFICATIONS:**

The unit must meet or exceed the requirements listed in the table below.

| **Product Details** | |
|---|---|
| Name of Manufacturer | |
| Model/ catalogue number | |
| Country of Origin for the offered model | |
| Country where the manufacturer is based | |
| Delivery time | |
| Full warranty period | |
| FDA clearance OR CE Mark | |

| | **Minimum Requirements** | **Compliance (Y/N), Notes** | *Brochure Page No.* |
|---|---|---|---|
| **1.** | **Durable, compact, heavy duty construction, reliable ergonomic design and easy to use: Sample must be submitted with the offer** | | |
| **2.** | It should be stated clearly in the catalogue that the unit is dedicated to PCA (patient controlled analgesia) | | |
| **3.** | Syringe protective mechanism that prevents accidental or unauthorized removal of syringe | | |
| **4.** | The unit must prevent unauthorized change in parameters | | |
| **5.** | Protection against accidental or unauthorized overdose | | |
| **6.** | Any required accessories and consumables that are necessary to operate the unit should be quoted | | |
| **7.** | **Performance data:** | | |
| | a- Flow rate setting: 0.1-999 ml/hr or better | | |
| | b- Volume (VTBI) setting 0.1-999 ml or better | | |
| | c- Drug dose rate calculation function | | |

| | Minimum Requirements | Compliance (Y/N), Notes | *Brochure Page No.* |
|---|---|---|---|
| | d- Automatic and manual bolus administration that is adjustable in rate and volume | | |
| | e- KVO (keep vein open) function | | |
| | f - Prime (purge) function | | |
| | g - Accuracy ± 2% (or better) of displayed flow rate and volume | | |
| | H - Adjustable limit for occlusion pressure. | | |
| | i - Automatic pressure should be reduced when occlusion detected | | |
| | J - To accept syringe sizes from 10 ml up to 50 ml (or better) from several manufacturers (major suppliers) while maintaining accuracy (open system). | | |
| 8. | **Pump Mechanism**: Syringe driver | | |
| 9. | Microprocessor controlled unit with start-up self-test capability with an error code system | | |
| 10. | **Medication database:** pre-programmed drug protocols | | |
| 11. | **Clear and easy to read LCD** display of the following: | | |
| | a- Flow rate | | |
| | b- Volume infused and volume to be infused | | |
| | c- Time remaining | | |
| | d- Battery capacity | | |
| | e- Pressure level | | |
| | f- Bolus delivery | | |
| | g- Medication name and settings | | |
| | h- Alarms and error codes | | |
| 12. | Hygienic, flat, soft touch keys control panel with visible and clearly identified controls | | |

| | Minimum Requirements | Compliance (Y/N), Notes | *Brochure Page No.* |
|---|---|---|---|
| 13. | Battery and mains AC power supply indicators | | |
| 14. | Built in maintenance free rechargeable battery that is easy accessible with capacity not less than 6 hours operation at 5 ml/h flow rate when fully charged. | | |
| 15. | Visual and audible alarms with adjustable alarm volume control, The following alarms should be included: | | |
| | a- High pressure occlusion. | | |
| | b- Syringe unlocked | | |
| | c- Malfunction alarms. | | |
| | d- Low battery alarm. | | |
| | e- Empty and near empty syringe | | |
| 16. | Easy and secured clamping mechanism for pole fixation | | |
| 17. | Light weight (≤ 3 kg) | | |
| 18. | The unit shall meet at least the following standards: | | |
| | **a- Defibrillation proof type CF** | | |
| | **b- IEC 60601-2-24:** safety and essential performance of infusion pumps | | |
| | **c- IPX 2 (or higher):** Fluid ingress protection | | |
| 19. | To work on either internal or external power supply | | |
| 20. | A list of standard accessories for the offered model. | | |
| 21. | Please quote for any other optional accessories; IV pole, docking station, etc... | | |

JORDAN ARMED FORCES

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| Item 2 | May-field horseshoe U shape 3point headrest for operation table | Qty. (1) |
|---|---|---|

**IMPORTANT NOTE:**

*Vendors are required to fill out the product details, compliance information, and brochure reference page number in the columns below*

**TECHNICAL SPECIFICATIONS:**

The unit must meet or exceed the requirements listed in the table below.

| Product Details | |
|---|---|
| Name of Manufacturer | |
| Model/ catalogue number | |
| Country of Origin for the offered model | |
| Country where the manufacturer is based | |
| Delivery time | |
| Full warranty period | |
| FDA clearance OR CE Mark | |

| | Minimum Requirements | Compliance (Y/N), Notes | *Brochure Page No.* |
|---|---|---|---|
| | Compatible with the available surgical tables in OR department in royal rehabilitation center (Eschmann model T20 A+ or Trumpf model Mars ) preferable both table models. | | |
| 1 | U-shaped head support | | |
| 2 | The head bracket can move forward & backward and left and right horizontally, and up and down vertically | | |
| 3 | Adjust at various angle | | |
| 4 | The locking mechanism is convenient | | |
| 5 | Three-point type fixed | | |
| 6 | Simple and convenient installation and operation. | | |
| 7 | Cross-bar and tighten arm are all made of stainless steel, economy and durable | | |
| 8 | All additional required supports and clamps should be included | | |

## *SPECIAL TERMS*

( for item 1 + 2 )

- *Offers not complying with any of the special terms or the technical specifications shall be considered non-conforming with tender requirements.*
- *Any vendor providing FORGED documents shall be disqualified from the current tender and banned from participating in any future RMS tenders.*
- *Wherever term "based" is mentioned it refers to the country where the manufacturing company is founded & established.*

1. *All equipment must be the most recently released model/version which is equal to or higher than the range of the specifications of the required system (low, mid or high) and equal to or higher than the level of technology and required options mentioned in the technical specifications.*

2. *Required certificates (must be submitted with the technical offer):*

   2.1 *FDA clearance (510K) for equipment of USA based companies.*

   2.2 *MDR (EU) 2017/745 certificate for equipment of EU (European Union) based companies.*

   2.3 *UKCA certificate For Equipment of Great Britain based companies (England, Scotland, Northern Ireland, and Wales).*

   2.4 *ARTG (Australian Register of Therapeutic Goods) certification or approval for Australian and New Zealand based companies.*

   2.5 *Establishment License with the relevant Device License issued by the Therapeutic Products Directorate for Equipment of Canadian based companies.*

   2.6 *PMDA (Pharmaceuticals and Medical Devices Agency) certification or approval for equipment of Japanese based companies.*

   2.7 *Swissmedic (Swiss agency for therapeutic products) certification or approval for equipment of Swiss based companies.*

   2.8 *Norwegian Medicines Agency certification or approval for equipment of Norwegian based companies.*

   2.9 *Only for class I medical equipment manufactured by companies based in one of the countries mentioned above, submission of either one of the certificates mentioned above or a free sale certificate in any of these countries shall be accepted.*

2.10 *For the following equipment:*

- *All X-ray equipment, MRI, ultrasound, and nuclear medicine systems (regardless where the manufacturing companies are based).*
- *Where the manufacturing companies are based in other origins than the mentioned in terms 2.1 - 2.8.*

*The following are required:*

*a- At least two of the certificates mentioned above, one of which has to be FDA clearance (510K) (Only for class I medical equipment submission of certificates mentioned in 2.9 shall be accepted).*

*b- Evaluation certificate from the Royal Medical Services for the same offered model with at least 80% passing grade.*

*If the evaluation is not applicable (based on purchasing committee perspective) bidder should submit a list of installation basis of the same offered model and/ or previous models in at least two of the following hospitals (King Hussein Cancer Center, National Center for Diabetes Endocrinology and Genetic Diseases, Jordan University Hospital or King Abdullah University Hospital) with at least three years of operation, list should include: Name of hospital, Model installed, Quantity, and date of installation.*

*The purchasing committee has the right to officially contact any of these hospitals and disqualify any offer where the feedback is negative in operation, after sales service or local agent performance.*

2.11 *The vendor is responsible to ensure through official documents that classified medical devices are manufactured in conformity with applicable quality system standards (ISO,*

*IEC); (the international quality systems standard for medical device manufacturing ISO 13485) or ISO9001.*

2.12 *With each offer, bidders must provide a formally endorsed document issued by the manufacturer stating that the bidder is the sole certified agent for the offered item.*

2.13 *In all of the above cases (except 2.12) certificates must be formally endorsed by JFDA.*

2.14 *Any vendor not submitting all required certificates will be eliminated.*

3. *Official website of the manufacturer must be provided. The website must clearly demonstrate the history profile and manufacturing features of the company including the same offered model and its brochure.*

   *In case of inconsistency between the information on the website with the provided information of the local agent regarding any offered product, the committee has the right to eliminate that product from the awarding process.*

4. *Offered items should be from reputable well known manufacturers and excellent experience in the field and shall have multiple installations of the same offered model and/ or previous models in RMS main hospitals with at least two years of operation and excellent experience in operation, after sales service & local agent performance; otherwise the purchasing committee has the right to request <u>any</u> of the following:*

   a) *An evaluation certificate as mentioned in term 2.10.b.*

   b) *A sample of the same offered model at any time during the purchasing process, the sample unit shall be delivered within three weeks from the date of a written notification; any vendor rejects to deliver a sample unit will be eliminated. Evaluation duration will be determined by the purchasing committee.*

   *<u>Any offered item fail in the evaluation/assessment process will be rejected</u>*

5. *Vendors must specify the origin of the offered items and accessories in the technical offer.*

6. *Bidders must submit their reservations/queries regarding tender specifications and/or special terms within the first half of the tender closing period starting from the tender announcement date. Reservations/queries submitted after the end of this period shall be rejected.*

7. *Goods are to be brand new; manufacturing date of the delivered products should not exceed 18 months from the date of the final award.*

8. *Delivery period should be mentioned clearly; the purchasing committee has the right to reject any offer with delivery time exceeding 4 months.*

9. *Warranty:*

   a- *Offers must include a full warranty for a period of a minimum of 24 months from the date of installation or 30 months from the date of receiving the items at the agreed location mentioned in the final order (whichever comes first).*

   *Warranty must include corrective & preventive maintenance activities as per manufacturer recommendations including:*

   - *Required spare parts (free of charge)*
   - *Labour*
   - *Hardware*
   - *Software*
   - *Rechargeable batteries*

   *At the end of the warranty period, The supplier commits to implement final inspection of the submitted goods and submit the reports signed by the site chief engineer stating that the equipment are working properly as well as all preventive maintenance reports during the warranty period.*

   *Warranty & installation are excluded for items mentioned in Attachment no.1 the purchasing committee has the right to exclude any other item not listed in the attachment and does not required installation and warranty.*

   b- *In the case where a delay in installation has occurred as a result of the supplier's dereliction and has exceeded a period of one month from the date of receiving the items, the approved warranty period will automatically be considered as 24 months from the date of installation.*

   c- *If at any time during the warranty period the item becomes inóperative due to a technical fault the item must then be repaired by the supplier /local agent within a period of 5 working days from written notification, warranty will be extended according to downtime period.*

d- *If the delay exceeds 30 days the supplier must replace the item with a new identical functioning one (within the same delivery period mentioned in the final offer). In case the item was replaced by a new one, the warranty period mentioned in 9.a) above will start from the installation and commissioning date of the new item.*

e- *Local agent/ supplier is committed to transport and install any awarded item during the warranty period on free of charge basis to any location inside the country whenever required by Royal Medical Services, this should not include pre-installation requirements in the new site.*

10. *All offered items (main unit) should be fully designed, manufactured, and labelled by their real original manufacturer in which all related testing, research, development and approvals went through.*

    *Any relabelled products for the main unit (white-label manufacturing, OEM, or repackaging) are rejected.*

11. *One set of operation manual(s) and one set of service manual(s) including schematics and a spare-part list must be delivered with each unit, CD/DVD is acceptable. For large tenders, a certain agreed percentage of manuals per item may be agreed upon.*

12. *Where applicable, pre-installation shall be the sole responsibility of the supplier. Pre-installation shall include removal of old system(s), any civil work, electrical work or site modification(s) necessary to accommodate the new system(s) according to manufacturers' specifications and safety standards in addition to the work required for bringing back the site to the same working conditions as before installing the new system(s).*

13. *Power requirements: where applicable either single phase 220V, 50Hz or 3-phase 380V. Systems with external transformers are considered conforming only if clearly stated in the technical specifications.*

14. *Technical offers must include clear original technical brochures/catalogues for all offered items.*

15. *Offers must include fully detailed technical offers and compliance sheets as a soft copy ( either Microsoft office or Microsoft excel format) in addition to a hard copy, mentioning the exact*

*model/catalogue number and country of origin of the offered item(s), full technical description/specifications and any accessories or options included in the offer.*

*Offer must clearly indicate the origin of the offered item and the country where the manufacturer of the offered items is based.*

16. *Compliance sheets must be as per the tabular format of the technical specifications in the tender documents, listing the required specifications on one column and a Yes or NO response to each point in the adjacent column, with reference to page and line numbers in the relevant technical brochure. Offers not complying with this term shall be rejected.*

17. *Qualifications and after sales service: The technical bid should contain all the necessary documents (training certificates) to prove the capability of the bidders for supplying and installing a trouble free equipment meeting the quality standards and technical specifications of the manufacturer and the ability of the bidders for providing efficient after sales service conducted by authorized certified biomedical service engineers with minimum 2 years experience in the same field.*

18. *Accessories and consumables:*

    18.1 *Any accessories and consumable items necessary to operate the offered system must be clearly identified and priced separately.*

    *All offered accessories and consumables must be approved by the manufacturer.*

    18.2 *Technical offers must include a priced list for all accessories and consumables related to the required equipment as a hard copy in addition to a soft copy (either Microsoft office or Microsoft excel format) with prices fixed for a period of five years from the date of installation and commissioning with a maximum annual increase of 2%, any essential item not listed will be considered free of charge.*

    18.3 *Accessories and consumables must be priced according to their delivery destination either to Queen Alia International Airport or to RMS Main Medical Stores.*

    18.4 *Where applicable, a start-up kit of accessories and consumable items must be provided with each system on a free-of-charge basis.*

19. *Spare Parts:*

    19.1 *Technical offers must include a comprehensive and priced list of all spare parts related to the awarded equipment (including rechargeable batteries) as a hard copy in addition to a soft copy (either Microsoft office or Microsoft excel format) valid for a minimum period of five years with a maximum annual increase of 2%, commencing at the end*

*date of the warranty period, any essential item not listed will be considered free of charge.*

19.2 *Spare parts must be priced according to their delivery destination either to Queen Alia International Airport or to RMS Main Medical Stores.*

19.3 *Prices of spare parts should be reasonable and will be taken into consideration during the purchasing process; the purchasing committee has the right to eliminate any offered item with unreasonable high prices of spare parts.*

19.4 *Delivery period of required spare parts should not exceed 2 months from the date of the final order.*

20. *Spare parts, consumables and accessories availability must be guaranteed for a minimum period of ten years starting from the date of installation and commissioning.*

21. *Local agent is committed to sign a service contract with Royal Medical Services to perform all relevant maintenance for the awarded equipment (whenever required by DRMS within a period that does not exceed one year from the end date of the warranty period and if mentioned clearly in the specifications) without exceeding the original mentioned cost and as per mentioned terms within minimum required period.*

22. *Tender Awards:*

22.1 *For the final list of offers having a chance of winning the award, the awarding process shall be based on the accumulative value of both the offered item and its' running cost (Total Cost of Ownership) over a period of seven years from the date of installation and commissioning. Only offers with the lowest total cost of ownership over a period of seven years from the date of installation and commissioning shall qualify for the award.*

22.2 *Running cost includes the value of consumables, accessories needed to operate the system over the same period as well as the cost of any service contract (where applicable).*

*23. For PC/Laptop based systems:*

*23.1 Complete restoration medium (CD/DVD/etc.) of the operating system and the application software must be supplied.*

*23.2 Purchased licenses, software keys & dongles must be provided at any time during and after the warranty period on free of charge basis whenever required by DRMS for a minimum period of ten years starting from the date of installation and commissioning.*

*23.3 Where locally supplied computers, laptops & printers are offered, the offered model should be from well known manufacturer.*

*24. Pricing must include services of sale, shipment, transportation, delivery from port to site or to Main Medical Stores, installation, pre-installation (if needed), training, commissioning, warranty and bringing the equipment into service.*

*25. Custom clearance of goods shall be the responsibility of the Jordanian Armed Forces (JAF), however, suppliers shall bear all costs incurred by handling charges and any demurrage charges or extra expenses incurred by the port's corporation (including expenses caused by delay in presenting the necessary shipment documents for either clearing or transporting the goods to the required location mentioned in the final order, delivery note issuing charges, unloading*

*charges, local shipping charges etc.). The supplier is also responsible for providing of all relevant shipping documents, together with the delivery order(s).*

26. *a. DRMS has the right to increase the awarded quantities by a percentage not exceeding 35% after the final order notification with the same prices, terms and conditions of the original contract upon DRMS request and approval of the awarded party.*

*b. DRMS has the right to decrease the awarded quantities by a percentage not exceeding 50% after the final order notification with the same prices, terms and conditions of the original contract upon DRMS request and approval of the awarded party.*

27. *a. Bidder is not allowed to submit more than one offer for the same item whether that's solely or in coalition or partnership with other bidders.*

*b. Bidder is allowed to include within their offer optional alternatives for the same offered item from the same manufacturer.*

28. *The supplier must furnish DRMS with a guarantee stamped and legalized by the Notary Public equals to (115%) of the total value of the awarded equipment valid for twelve months from the date of final acceptance of the equipment by DRMS.*

29. *Training:*

*29.1 For items where abroad service training courses are required in technical specifications, offers must include a certified service training program for at least 3 working days at a reputable center abroad recognized by the manufacturer for at least one biomedical engineer or biomedical technician; all costs inclusive, air tickets, , boarding, commuting, accommodation (minimum 3 star hotel on full board basis) and any extra costs.*

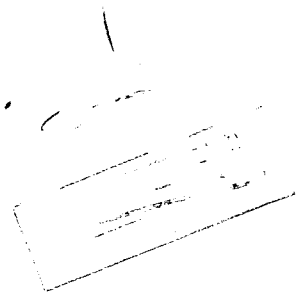
*29.2 For items where abroad <u>user</u> training courses are required in technical specifications, offers must include a certified operator training program at a reputable center abroad recognized by the manufacturer for at least one operator; all costs inclusive, air tickets, boarding, commuting, accommodation (minimum 3 star hotel on full board basis) and any extra costs.*

*29.3 The period of the training courses must be according to the manufacturer's program excluding traveling days and must be stated clearly in the technical offer.*

*29.4 Training Programs must conform to the following standards:*

- *User training must comprise understanding and use of operation manual(s), correct and safe operation of the equipment, as well as user preventive maintenance and calibration.*
- *Service training must comprise: theory, understanding and use of service manual(s), calibration, preventive maintenance procedure, and practical troubleshooting and repair exercises, and must be conducted by professional instructors employed or authorized by the system manufacturer.*
- *Service training must be conducted on a system of identical make, model, and configuration to that purchased by DRMS, and designated by the manufacturer or the local agent for training purposes.*
- *Certificates must be endorsed and officially sealed by the system manufacturer, legally empowering trainees to engage in user and service activities according to operation and service manual(s).*
- *Where applicable, offers must include an on-site user and service training.*

30. For offers submitted in Jordanian dinars, payment will be either by wire transfer or by cheque after final acceptance of good, any other way of payment will be rejected.



| # | Equipment |
|---|---|
| 1 | Adjustable Auto Pipettes |
| 2 | Resuscitation Bag |
| 3 | Laryngoscope Set |
| 4 | Oxygen Flow meter wall type/ single |
| 5 | Regulator Suction with canister, wall vacuum outlet |
| 6 | Oxygen Regulator for Oxygen Cylinders |
| 7 | Pulse Oximeter, Finger type |
| 8 | Oxygen Cylinder |
| 9 | Doppler, portable |
| 10 | Diagnostic set, Portable |
| 11 | Direct Ophthalmoscope, Portable |
| 12 | Otoscope, Portable |
| 13 | Air Mattress System, homecare |
| 14 | Stethoscope |
| 15 | Aneroid Sphygmomanometer |
| 16 | Video Assisted Laryngoscope, portable |
| 17 | Scale, Manual |
| 18 | Wood's Light |
| 19 | Cough Pressure, Normal Saline |
| 20 | Rehabilitation Walking Parallel Bars, non-powered |
| 21 | Therapy Mat |
| 22 | Medical Ball All Size |
| 23 | Dumbbells Rack with complete set of dumbbells |
| 24 | Crutches |
| 25 | Shoulder wheel |
| 26 | Mobile Mirror |
| 27 | Cuff Weights |
| 28 | Walker, different sizes |
| 29 | Patient Elbow Stick |
| 30 | Rehabilitation Training Ladder |
| 31 | Rehabilitation Suspension Frame |
| 32 | Exercise Band All Size (Theraband) |
| 33 | Lens trial set |
| 34 | Wheel Chair |

| | |
|---|---|
| 35 | *Commode Chair* |
| 36 | *Bassinet (Baby Cot)* |
| 37 | *Resuscitation Cart (Crash Cart)* |
| 38 | *Medication Cart* |
| 39 | *Cart, Drawers* |
| 40 | *Examination Couch, Manual* |
| 41 | *Gynaecology Examination Table, Manual* |
| 42 | *Examination Table, Neonates, Manual* |
| 43 | *Intravenous Pole, Mobile Stand* |
| 44 | *DDA Cabinet* |
| 45 | *Stainless steel Multipurpose Trolley* |
| 46 | *Cabinet, Instrument, Operation Theatres* |
| 47 | *Dressing Cart* |
| 48 | *Stainless steel wire shelving unit* |
| 49 | *Paper Trolley* |
| 50 | *Stainless Steel Sink (Clean up counter)* |
| 51 | *Scopes Cabinet* |
| 52 | *Mayo Table* |
| 53 | *Table, Instrument* |
| 54 | *Stool, Adjustable, Doctor* |
| 55 | *Stool, Adjustable, Operation Theatres* |
| 56 | *Carts, linen/laundry, soiled, Double* |
| 57 | *Closed distribution trolley* |
| 58 | *Step Ladder, Conductive, Double* |
| 59 | *Step, Surgeon, Single* |
| 60 | *Kick Bucket* |
| 61 | *Mobile Stand for Oxygen Cylinder* |
| 62 | *Stainless Steel Wire Basket, 1 STU* |
| 63 | *Cart, Plaster* |

# عدد جراحية لقسم التجميل في مركز التاهيل الملكي

| | | |
|---|---|---|
| **Item 3** | Arnaud craniofacial 10distractor 1.5,30 mm with driver | **Qty. (4)** |

**IMPORTANT NOTE:**

*Vendors are required to fill out the product details, compliance information, and brochure reference page number in the columns below*

**TECHNICAL SPECIFICATIONS:**

The unit must meet or exceed the requirements listed in the table below.

| **Product Details** | |
|---|---|
| Name of Manufacturer | |
| Model/ catalogue number | |
| Country of Origin for the offered model | |
| Country where the manufacturer is based | |
| Delivery time | |
| Full warranty period | |
| FDA clearance OR CE Mark | |

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| | | |
|---|---|---|
| **Item 4** | Kaye scissors | **Qty. (2)** |

**IMPORTANT NOTE:**

*Vendors are required to fill out the product details, compliance information, and brochure reference page number in the columns below*

**TECHNICAL SPECIFICATIONS:**

The unit must meet or exceed the requirements listed in the table below.

| **Product Details** | |
|---|---|
| Name of Manufacturer | |
| Model/ catalogue number | |
| Country of Origin for the offered model | |
| Country where the manufacturer is based | |
| Delivery time | |
| Full warranty period | |
| FDA clearance OR CE Mark | |

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| | | |
|---|---|---|
| **Item 5** | Trepstat facelift dissection ,saptulated tip | **Qty. (2)** |

**IMPORTANT NOTE:**

*Vendors are required to fill out the product details, compliance information, and brochure reference page number in the columns below*

**TECHNICAL SPECIFICATIONS:**

The unit must meet or exceed the requirements listed in the table below.

| **Product Details** | |
|---|---|
| Name of Manufacturer | |
| Model/ catalogue number | |
| Country of Origin for the offered model | |
| Country where the manufacturer is based | |
| Delivery time | |
| Full warranty period | |
| FDA clearance OR CE Mark | |

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| | | |
|---|---|---|
| **Item 6** | Osteotome straight 2mm | **Qty. (4)** |

**IMPORTANT NOTE:**

*Vendors are required to fill out the product details, compliance information, and brochure reference page number in the columns below*

**TECHNICAL SPECIFICATIONS:**

The unit must meet or exceed the requirements listed in the table below.

| **Product Details** | |
|---|---|
| Name of Manufacturer | |
| Model/ catalogue number | |
| Country of Origin for the offered model | |
| Country where the manufacturer is based | |
| Delivery time | |
| Full warranty period | |
| FDA clearance OR CE Mark | |

| | | |
|---|---|---|
| **Item 7** | Mitchells trimmer | **Qty. (4)** |

**IMPORTANT NOTE:**

*Vendors are required to fill out the product details, compliance information, and brochure reference page number in the columns below*

**TECHNICAL SPECIFICATIONS:**

The unit must meet or exceed the requirements listed in the table below.

| **Product Details** | |
|---|---|
| Name of Manufacturer | |
| Model/ catalogue number | |
| Country of Origin for the offered model | |
| Country where the manufacturer is based | |
| Delivery time | |
| Full warranty period | |
| FDA clearance OR CE Mark | |

JORDAN ARMED FORCES

The DIRCETROATE OF ROYAL MEDICAL SERVICES

THE INSTITUE OF BIOMEDICAL TECHNOLOGY

| | | |
|---|---|---|
| Item 3 | Dandy artery forceps lateral curved | Qty. (20) |

**IMPORTANT NOTE:**

*Vendors are required to fill out the product details, compliance information, and brochure reference page number in the columns below*

**TECHNICAL SPECIFICATIONS:**

The unit must meet or exceed the requirements listed in the table below.

| Product Details | |
|---|---|
| Name of Manufacturer | |
| Model/ catalogue number | |
| Country of Origin for the offered model | |
| Country where the manufacturer is based | |
| Delivery time | |
| Full warranty period | |
| FDA clearance OR CE Mark | |

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| | | |
|---|---|---|
| **Item ٩** | Malleable brain re-tractor set (all size ) | **Qty. (2 set )** |

**IMPORTANT NOTE:**

*Vendors are required to fill out the product details, compliance information, and brochure reference page number in the columns below*

**TECHNICAL SPECIFICATIONS:**

The unit must meet or exceed the requirements listed in the table below.

| **Product Details** | |
|---|---|
| Name of Manufacturer | |
| Model/ catalogue number | |
| Country of Origin for the offered model | |
| Country where the manufacturer is based | |
| Delivery time | |
| Full warranty period | |
| FDA clearance OR CE Mark | |

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| | | |
|---|---|---|
| **Item 10** | Tie wire gauge 28,316 annealed stainless steel | **Qty. (30 roll /10m )** |

**IMPORTANT NOTE:**

*Vendors are required to fill out the product details, compliance information, and brochure reference page number in the columns below*

**TECHNICAL SPECIFICATIONS:**

The unit must meet or exceed the requirements listed in the table below.

| **Product Details** | |
|---|---|
| Name of Manufacturer | |
| Model/ catalogue number | |
| Country of Origin for the offered model | |
| Country where the manufacturer is based | |
| Delivery time | |
| Full warranty period | |
| FDA clearance OR CE Mark | |

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| | | |
|---|---|---|
| **Item 11** | Rein clips and applicators | |

**IMPORTANT NOTE:**

*Vendors are required to fill out the product details, compliance information, and brochure reference page number in the columns below*

**TECHNICAL SPECIFICATIONS:**

The unit must meet or exceed the requirements listed in the table below.

| **Product Details** | |
|---|---|
| Name of Manufacturer | |
| Model/ catalogue number | |
| Country of Origin for the offered model | |
| Country where the manufacturer is based | |
| Delivery time | |
| Full warranty period | |
| FDA clearance OR CE Mark | |

| | **Minimum Requirements** | **Qty.** | **Y/N) , Notes (Compliance** | ***Brochure Page No.*** |
|---|---|---|---|---|
| .1 | Rein clips | 40 | | |
| .2 | Rein applicator | 4 | | |

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Item 12 **Microsurgery set** **Qty. (2) Set**

**IMPORTANT NOTE:**

*Vendors are required to fill out the product details, compliance information, and brochure reference page number in the columns below*

***All tools should be laser printed with the manufacturer name and (made in "country of origin").***

**TECHNICAL SPECIFICATIONS:**

The unit must meet or exceed the requirements listed in the table below.

| **Product Details** | |
|---|---|
| Name of Manufacturer | |
| Model/ catalogue number | |
| Country of Origin for the offered model | |
| Country where the manufacturer is based | |
| Delivery time | |
| Full warranty period | |
| FDA clearance OR CE Mark | |

| | **Minimum Requirements** | **Qty.** | **Y/N) , (Compliance Notes** | ***Brochure Page No.*** |
|---|---|---|---|---|
| **1** | Jeweler's forceps | 2 | | |
| **2** | Vessel dilator | 1 | | |
| **3** | Castrofiejo curved dissection scissors | 1 | | |
| **4** | Catrofiejo straight cutting scissors | 1 | | |
| **5** | Castrofiejo needle holder | 1 | | |
| **6** | Curved jeweler's forceps | 1 | | |
| **7** | Micro-artery clamps | 2 | | |

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| | | |
|---|---|---|
| **Item 13** | Lone –star hook retractors | **Qty. (8)** |

**IMPORTANT NOTE:**

*Vendors are required to fill out the product details, compliance information, and brochure reference page number in the columns below*

**TECHNICAL SPECIFICATIONS:**

The unit must meet or exceed the requirements listed in the table below.

| **Product Details** | |
|---|---|
| Name of Manufacturer | |
| Model/ catalogue number | |
| Country of Origin for the offered model | |
| Country where the manufacturer is based | |
| Delivery time | |
| Full warranty period | |
| FDA clearance OR CE Mark | |

*Jordan Armed Forces – Arab Army*

*Directorate of pharmacy and medical supply*

*Special terms for medical instruments*

( For Item 3 – 13)

- Offers not complying with any of the special terms or the technical specifications shall be considered non-conforming with tender requirements.

- Any vendor providing FORGED documents shall be disqualified from the current tender and banned from participating in any future RMS tenders

- Offers submitted by vendors with previously complaints or unsolved issues will be excluded from the current tender

1. The vendor is responsible to ensure through official documents that classified medical instrument are manufactured in conformity with applicable quality system standards (ISO) and must provide the latest copy of the certificates of the following standards formally endorsed by the chamber of commerce of the manufacturing origin with the technical offer:

A- ISO 13485:2016

Medical devices — Quality management systems — Requirements for regulatory purposes

ISO 13485:2016 specifies requirements for a quality management system where an organization needs to demonstrate its ability to provide medical devices and related services that consistently meet customer and applicable regulatory requirements.

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B- ISO 9001:2015

Quality management systems — Requirements

specifies requirements for a quality management system aims to enhance customer satisfaction through the effective application of the system, including processes for improvement of the system and the assurance of conformity to customer and applicable statutory and regulatory requirements.

C- ISO 7153-1:2016

Surgical instruments — Materials — Part 1: Metals
specifies metals commonly used to manufacture various types of standard surgical instruments, including but not limited to those used in general surgery, orthopaedics and dentistry.

D- ISO 7151:1988

Surgical instruments — Non-cutting, articulated instruments — General requirements and test methods
Specification of basic requirements for as well physical characteristics as workmanship, of the steel grades used and heat treatment of component parts, excluding rivets, screws and parts manufactured of material grade M.

E- ISO 7740:1985

Instruments for surgery — Scalpels with detachable blades — Fitting dimensions
Lays down the dimensions of two sizes of fitting features for detacheable scalpel blades and the handles with which they are used. It secures a good fitting and interchangeability of detachable blades for scalpels manufactured in different countries or by different manufacturers

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F- ISO 7741:1986

Instruments for surgery — Scissors and shears — General requirements and test methods

This standard deals with materials, heat treatment and hardness of component parts, corrosion resistance, workmanship and cutting ability of scissors and shears used in the surgery and defines the test methods.

G- ISO 13402:1995

Surgical and dental hand instruments — Determination of resistance against autoclaving, corrosion and thermal exposure

Describes test methods to determine the resistance of stainless steel surgical and dental hand instruments against autoclaving, corrosion and thermal exposure.

2. Offered items should be from reputable well known manufacturers and excellent experience in the field previously evaluated and accepted in Main Hospitals at Royal Medical Service . Bidder must submit samples with the technical offer from each brand that was submitted in the offer to be assessed and /or evaluated by RMS

   A. Evaluation duration and quantities will be determined by the purchasing committee.
   B. Offers which do not include such samples will be considered non-conforming.
   C. Offers which fail the evaluation/assessment process will be excluded from the tender.
   D. Samples will be returned to the Bidder at the end of the evaluation/assessment process except for samples from Bidders winning the final award where they will be the property of RMS and will be kept as a reference for the receiving committee.

*Special terms for medical instruments*

E. The purchasing committee has the right to require a sample for any instrument previously installed in RMS Hospitals.

3. A. Bidder is not allowed to submit more than one offer for the same item whether that's solely or in coalition or partnership with other bidders.

   B. Bidder is allowed to include within their offer optional alternatives for the same offered item from the same manufacturer.

4. Required certificates:
   A. For instruments of USA origin, a copy of a certificate of FDA approval must be submitted with the technical offer.
   B. For instruments of other origins, a copy of either a CE certificate (MDD)/TUV OR a certificate of FDA approval must be submitted with the technical offer.
   C. With each offer, bidders must provide a formally endorsed document issued by the manufacturer stating that the bidder is the authorized agent/ distributor for the offered item.
   D. In all of the above cases (except C) certificates must be formally endorsed by JFDA .
   E. Any vendor not submitting all required certificates will be eliminated.

5. All Offered items should be supplied with free sale certificate said confirmation should be stating clearly that the item is freely sold in the country of origin .
6. All items should be engraved or etched with manufacturing origin ,company logo and code number .
7. Each instrument set will be awarded as a listed set which including its specific sterilization baskets and trays .
8. Bidders must submit their reservations/queries regarding tender specifications and/or special terms within the first third of the tender closing period starting from the tender announcement

date. Reservations/queries submitted after the end of this period will be rejected

9. The supplier must furnish DRMS with a guarantee letter stamped and legalized by the Notary Public equals to (115%) of the total value of the awarded equipment valid for (24) months from the date of final acceptance of the instruments by DRMS which shall cover rusting, manufacturing defects, wear and tear and corrosion.
10. Offers must include clear original catalogues for all offered items. Images and part numbers of offered items must be provided, highlighted and outlined clearly.
11. Offers must include fully detailed information as a soft copy (either Microsoft office or Microsoft excel format) in addition to a hard copy , mentioning the exact model/catalogue number and country of origin of the offered item(s), full description/specifications. and any accessories or options included in the offer.
12. Official website of the manufacturer must be provided. The website must clearly demonstrate the history profile and manufacturing features of the company. In case of inconsistency between the information on the website with the provided information of the local agent regarding any offered product, the purchsing committee has the right to eliminate that product from the awarding process.
13. Any complementary parts , accessories and consumable items necessary for the proper operation of surgical Instruments must be included in the offer , priced separately and to be be approved by the manufacturer.
14. A. complementary parts availability must be guaranteed and can be purchased without the need replace the entire system of surgical instruments

    B. Bidder will bear any responsibility resulting from the awarding of insufficient items or sub-items.
15. Pricing must include services of sale, shipment, transportation, delivery from port to site or to Main Medical Stores .
16. A. custom clearance of goods shall be the responsibility of the Jordanian Armed Forces (JAF), however, suppliers shall bear all

costs incurred by handling charges and any demurrage charges or extra expenses incurred by the port's corporation (including expenses caused by delay in presenting the necessary shipment documents for either clearing or transporting the goods to the required location mentioned in the final order, delivery note issuing charges, unloading charges, local shipping charges etc.).

B. The supplier is also responsible for providing of all relevant shipping documents , together with the delivery order(s).

17. Delivery period should be mentioned clearly; the purchasing committee has the right to reject any offer with delivery time exceeding 4 months.
18. Goods are to be brand new; manufacturing date of the delivered products should not exceed 18 months from the date of the final award.
19. A. DRMS has the right to increase the awarded quantities by a percentage not exceeding 35% after the final order notification with the same prices, terms and conditions of the original contract upon DRMS request and approval of the awarded party.

    B. DRMS has the right to decrease the awarded quantities by a percentage not exceeding 50% after the final order notification with the same prices, terms and conditions of the original contract upon DRMS request and approval of the awarded party.

20. For offers submitted in Jordanian dinars, payment will be either by wire transfer or by cheque after final acceptance of good, any other way of payment will be rejected.