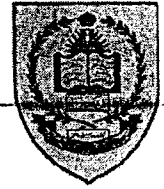




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Item 1	Vacuum Mattress, surgical table	Qty. (1)
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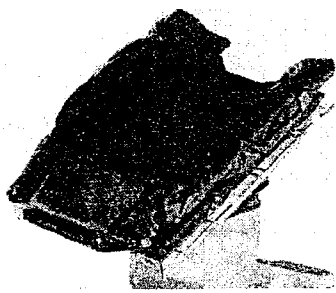
IMPORTANT NOTE:

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	For patient positioning safely in steep supine trendelenburg position, as per figure 		
2	The mattress should be securely fixed on the OR table to prevent the patient from sliding off the table, all fixation accessories should be included		
3	All additional required supports and clamps should be included and priced separately		
4	Size: 100*125 cm or larger		
5	Pump should be included		
6	Reusable and easy to clean and disinfect		



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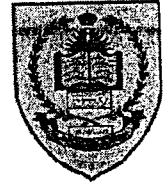
model/catalogue number and country of origin of the offered item(s), full technical description/specifications and any accessories or options included in the offer.

7. 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.
8. 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.
9. 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).
10. DRMS has the right to increase or decrease the awarded quantities by a percentage not exceeding 30% 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.
11. 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 twelvemonths from the date of final acceptance of the equipment by DRMS.



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- iv. *Vendors must specify the origin of all offered items and accessories in the technical offer.*
- v. *Equipment manufactured by reputable companies based in any of the countries mentioned in (3.i) will be taken into consideration regardless of the manufacturing site only:*
- a. *If they are approved for sale in the same country of origin (an original and officially endorsed free-sale certificate from an authorised body must be included in the offer.*

OR

- b. *If they are approved for sale in at least three of the countries mentioned in (3.1) (an original and officially endorsed free-sale certificate from an authorised body in those countries must be included in the offer).*
- vi. *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 shall be rejected.*

3. *Warranty for item 1 & 2:*

- i. *Offers must include a full warranty including spare parts and labour for a period of a minimum of 24 months from the date of installation.*
- ii. *If at any time during the warranty period the item becomes inoperative due to a technical fault the item must then be repaired by the supplier /local agent within a period of fourteen days from written notification, otherwise the supplier must replace the item with a new identical functioning one and will endure a penalty determined by the Royal Medical Services for each day of the downtime of the system. In case the item was replaced by a new one, the warranty period mentioned in (4.i) above will start from the installation and commissioning date of the new item.*

4. *Power requirements (for item 1): where applicable either single phase 220V, 50Hz V. Systems with external transformers are considered conforming only if clearly stated in the technical specifications.*

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

6. *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*



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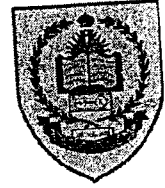


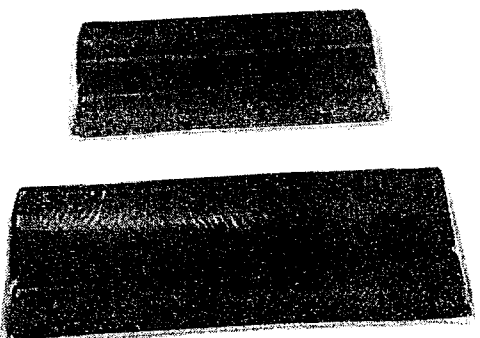
SPECIAL TERMS

- 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.
1. Required certificates:
- 2.1 For equipment of US origin, a copy of a certificate of FDA approval & the relevant 510K clearance for selling to US healthcare facilities for the offered model must be submitted with the technical offer.
 - 2.2 For equipment of other origins, a copy of either a CE certificate with the relevant CE number (MDD)/TÜV/BSI/UL OR a certificate of FDA approval & the relevant 510K clearance for selling to US healthcare facilities for the offered model must be submitted with the technical offer.
 - 2.3 Only for class I medical equipment, submission of a copy of Declaration of Conformity certificate (MDD) for the offered model shall be accepted.
 - 2.4 With each offer, bidders must provide a formally endorsed document issued by the manufacturer stating that the bidder is the sole certified agent / distributor for the offered item.
 - 2.5 In all of the above cases (except 2.4) certificates must be formally endorsed by JFDA.
2. Country of origin:
- i. The country of origin of the main part (s) of the system must be one of the following:
USA, Canada, Japan, UK, Sweden, Finland, Denmark, Switzerland, Belgium, Germany, France, Netherlands, Spain, Norway, Italy, Ireland, Austria, New Zealand, Australia & Czech Republic.
 - ii. Accessories and consumables (as determined by the purchasing committee) may be manufactured in other countries and/or by different manufacturers.
 - iii. All offered items must be approved for sale in the same country of origin.



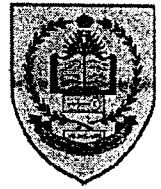
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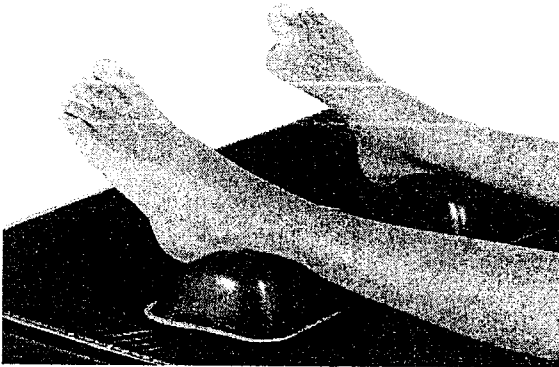
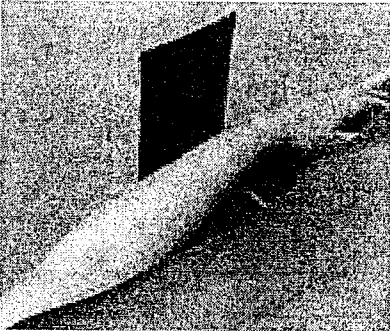
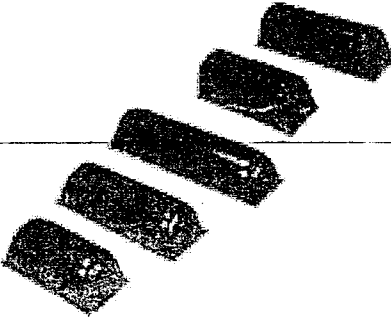
	Minimum Requirements	Qty	Compliance (Y/N), Notes	Brochure Page No.
5	<p>Knee Pad, to protect the knees against pressure damage in the prone position in various procedures Approximate size (510*150*40 mm)</p> 	2		



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	Minimum Requirements	Qty	Compliance (Y/N), Notes	Brochure Page No.
2	<p>Heel Pads, provide support and protection for the heels during long surgeries</p> <p>Approximate size (180*100*70)</p> 	1 pair		
3	<p>Arm Retainer Pad, designed to Perspex/ acrylic arm supports</p> 	2		
4	<p>Flat Bottom Chest Roll, provide support and stability during surgery</p> <p>Approximate size (510*150*150 mm)</p> 	4		



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Item 3	Gel Pad Positioner	Qty. (1 set)
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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	Compliance (Y/N), Notes	Brochure Page No.
1	Prone Head Rest; anatomically designed so that the patient head can comfortably rest face down with respiratory tubes in place Approximate size (280*240*140 mm) 	1		



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	Minimum Requirements	Compliance (Y/N), Notes	Brochure Page No.
3	Allows easy movement of the leg when placing it in the desired position		
4	Lifting capacity \geq 150 kg		
5	Range of movement: - Lithotomy position between +80 to -30 or better - Abduction range between +25 to -8 or better		
6	Handle to Provide easy adjustment of the stirrups		
7	Reusable boot pad		
8	With fixing clamp		



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Item 2	Stirrups, Surgical table, Pair	Qty. (1)
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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	Compatible with the available surgical tables in OR department in King Hussein Hospital (Eschmann model T20 A+ & Trumpf model Mars)		
2	Boot design stirrups, designed to cover and protect the head of the fibula and the peroneal nerve, as per figure 