HSCC I LTD

Dated: 24.01.2023

Call for Objection Notice

Reference : Annexure 6 Proprietary Article Certificate

Subject: "Call for Objection" against Procurement of **Vacuusport for Upper limb** for Expansion of Sports Injury Center New Delhi, where OEM is M/s. Weyergans High Care AG, Address –An Gut Boisdorf 8, 52355 Dueren, Germany under proprietary article certificate.

Vacuusport for Upper limb is intended to be procured on proprietary article basis as per provision of GFR 2017, directly from M/s. Trusted Resources Pvt. Ltd. Kailsah House 4B/27 Tilak Nagar New Delhi, an authorized dealer/distributor/agent of M/s. Weyergans High Care AG, against their proposal submitted and by accepting that /s. Weyergans High Care AG, Address –An Gut Boisdorf 8, 52355 Dueren, Germany are the sole manufacturer/supplier of item which satisfies all requirements of being a proprietary item.

Relevant documents are published on Institute's website & CPP Portal for inviting objection(s)/claim(s)/comment(s), if any, from eligible manufacturer/supplier, before accepting the claim of earlier said manufacturer and accordingly making procurement under proprietary article.

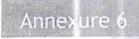
Objections(s)/claim(s)/comment(s) should be sent to the office of GM Procurement HSCC (I) LTD in a sealed envelope with above mentioned subject & reference number, or by e-mail at proc@hsccltd.co.in and hsccsic9@gmail.com in within 15 days from the date of publication of this notice i.e on before 06/02/2023 up to 05:00PM.

After due date, it will be assumed that no manufacturer/supplier has any objection/claim against above said equipment article & same will be consider as proprietary article.

Pawan Kumar Bhatia GM, (Projects) HSCC(I)Ltd.

Encl: Related Documents:

- 1. PAC Certificate by Manufacturer
- 2. PAC Certificate by Department
- 3. Technical Specification



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Annexure 6: Proprietary Article Certificate (Refer Para 4.6.1)

Valid for the Current Financial Year

File N	Number and Date I				a	
1	Description of article			Vacusport for Upper Extremity		
2	Forecast of quantity/annual requirement			01		
3	Approximate estimated value for above quantity			40,00,000/-		
4	Maker's name and address			M/s Weyergans High Care AG Address- An Gut Boisdorf 8, 52355 Duer Germany		
5	Name(s) of authorised dealers/stockists			M/s Trusted Resources Pvt. Ltd New Delhi		
6	I approve the above purchase on PAC basis and certify that: Note- Tick to retain only one out of (b), (c-1) or (c2) whichever is applicable and out others. Please do confirm (a) by ticking it – without which PAC certificate w invalid.					
6(a)	This is the only firm who is manufacturing/stocking this item. AND					ι
6(b)	the second data and the se					
-	No other make/brand will be suitable for following tangible reasons (like					
6 (c-1)		pares):	0	·K		
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Weyergans High Care AG - An Gut Boisdorf 8 - 52355 Düren The Director Sports Injury Center, Safdarjung Hospital Ansari Nagar East, Near to AIIMS Metro Station New Delhi 110029 India

Weyergans High Care AG An Gut Boisdorf 8 - 52355 Düren Germany Tel. +49-(0)2421-9678-0 Fax +49-(0)2421-9678-20 info@wcyergans.de www.weyergans.com

Proprietary Certificate

Dear Sirs.

We herewith certify that the "VACUMED A.N.S Intermittent Vacuum Therapy" device is solely developed and manufactured by Weyergans High Care AG and is a proprietary article.

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PUBL

DL: 1/11/2022

RAJ KUMAR Area: DEL MI Regn No. 6536

EXP.

Weyergans High Care AG is the only manufacturer and patent holder for the VACUMED A.N.S. device. The patent for this Intermittent Vacuum Therapy device is hold at the Germany Patent Office in Munich under the reference numbers 102013111208.0 and 19912611.9 -45 and is valid world-wide with the following Applications & Technical Specifications.

Intermittent Negative Pressure Therapy for Arm, Neck & Shoulder

The intended use of the equipment is to treat

- Repetitive Strain Injury Syndrome Mouse Arm
- Complex Regional Pain Syndrome
- Carpal Tunnel Syndrome
- Post T aumatic/ Post Operative Swelling
- Wound Healing Acceleration
- Odema Therapy



27th June 2022



Spankasse Aachen - IBAN: DE59 3905 0000 0047 2845 00 - BIC: AAC 5D E 33 Commerzhank AG Köln- IBAN: DE47 3708 0040 0101 7780 00 - BIC: DRES DE FF 370 VAT No. DE 121678028 - HRB-Düren 2678 - DIN EN ISO 9001 - DIN EN ISO 13485



The Interm ttent Vacuum Therapy Equipment for upper body must have:

- Negative Pressure adjustable from -20 to 100 mbar
- Intervals of negative pressure and atmospheric pressure adjustable from 1 to 20 see

 The height and angle of the negative pressure tube can be individually adjustable for optimal ergonomics and patient comfort.

- Dimensions: 1050 X 780 X 1030-1280 mm (W, D, H) and approx weight 80 Kg.
- Should be medical product class IIa.
- Should have pre programmed protocols for different medical conditions
- Should have a control panel to select program
- Input Voltage: 230 V AC/ 800 Watt.

For any further queries, please do not hesitate to contact us.

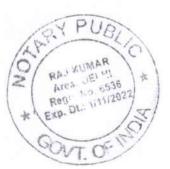
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We are looking forward to hearing from you soon.

Yours sincerely,

WEYERGANS HIGA CARE AG

ppa. Dorothee Sieger (authorized signatory)





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VACCUSPORT FOR UPPER LIMB

DPR Item No.: 60 Cost per equipment: Rs 40,00,000/-No. Of units: 1 Total cost: Rs 40,00,000/- (As in DPR)

It is an effective way to decongest post-traumatic or post-operative swelling, therefore accelerating regeneration and rehabilitation in patients and athletes.

TECHNICAL SPECIFICATION

- 1. Negative pressure adjustable from 20->100 mbar(-) or -20 to> -100 mbar.
- 2. Intervals of negative pressure and atmospheric pressure should be adjustable from 1-20 sec each.
- 3. The height and angle of negative pressure tube should be individually adjustable for optimal ergonomics and patient comfort.
- Dimensions 105 x 78 x 103-128 cm (W x D x H) and approx weight of 80kg.
- 5. Nine fully automatic indication-specific treatment programmes should be available.
- 6. The duration of a treatment should be between 25-40 mins, depending on the indication.
- Input voltage: 220-240V AC/800W (Comply with Indian electricity safety certification)
- 8. Should have CE/ US FDA/ Appropriate Indian Medical Safety & Quality Standard certification.

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- 1. GENERAL SPECIFICATIONS (terms and conditions) FOR physiotherapy and sports medicine, C-ARM , BATTERY DRILL, ARTHROSCOPES, MEDICAL EQUIPMENTS AND AUTOCLAVES (as per minutes 01/10/2019,15/10/2019,22/10/2019,31/10/2019 &07/11/2019 ,04/08/2020 over and above general specification by tendering authority)-TO BE FOLLOWED IN ALL MEDICAL EQUIPMENT ITEMS ON SIC EXPANSION.
- 1) System should be ECE / US FDA/BIS approved.
- 2) Inadvertent use of only US FDA in all specifications drafted by medical equipment committee to be omitted or not taken and should be read as and /or others as above in point 1.
- 3) Medical Grade Stainless steel / titanium or other certified material (satisfying above certification) must be used for all instruments and machinery in mandate of medical equipment committee for SIC expansion.
- 4) AERB approved.(For C-Arm)
- 5) 5 year comprehensive onsite warranty of entire system (Spare and labour) including X-ray tube (For C-Arm) and all accessories and civil, electrical and air conditioning works followed by 5 year CMC.
- 6) Company should confirm the availability of spare parts for 10 years from the date of supply of the equipment.
- Company should have 24 x 7 call support facility.
- List of spare parts with cost must be provided.
- 9) Should conform to latest IEC standard for requirement of safety for electromagnetic compatibility.
- 10) Physical Demonstration is essential.
- 11) Warranty 2 years in Arthroscopes subject to physical damage and wear &tear, and 1 year for Hand held Instruments of arthroscopy subject to physical damage and wear &tear ;
- 12) Warranty 5 years + CMC 5 years in all other items of this committee.
- 13) L1 will include CMC cost also;
- 14) Preventive maintenance every 3 months with log book entry;
- 15) The quality of supplied of instrument / equipment should be strictly same as given for physical demonstration / inspection, failing that the company / dealer will be black listed for 5 years for participating in tender process of hospital.
- 16) Turnkey/installation wherever required to be done by company and included in cost.
- 17) Freeze rate of consumables for 10 years
- 18) Technical specifications to be kept on site for 21 days public domain for comments of stakeholders.
- 19) Prebid meeting to be held of all stakeholders and modifications if needed may be considered.
- 20) On site training for equipment usage to be done wherever needed.
- 21) Patient and user Safety and compensatory clause for all medical machinery of this committee.
 - This should be besides the safety checks and mechanism already essential in every medical machinery.
- 22) Service centre authorized by manufacturer should be located in Delhi NCR and should provide service and repair as soon as possible within 24hrs to ensure patient safety.

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- 23) All electrical equipment should conform to latest electromagnetic safety standards for medical equipment.
- 24) The equipment with all its components will have warranty for period of 5 years from the date of handling over the fully functional unit and all the accessories supplied to the institution. Every thing irrespective of nature that is supplied by the vendor will be under warranty. The equipment which has multiple components should be quoted as a whole and by a single vendor.
- 25) During warranty period the desired uptime of 95% of 365 days (24hrs basis) will have to be ensured with maximum of 5 working days of downtime at a stretch. In case the downtime exceeds the 5%limit in a year or more than 5days at a stretch (whichever is applicable) extension of warranty period by double the excess down time period will be carried out.
- 26) CMC (Comprehensive Maintenance Contract) The post warranty (after 5 years) CMC should be comprehensive for all its components (everything irrespective of nature which is supplied by the vendor under guarantee) inclusive of X-Ray Tubes. (For C-Arm) with 95% uptime and extension of CMC period by double the downtime in excess of 5%.
- 27) Participating bidders should submit a good performance certificate from institutions and a certificate that it is submitting the medical item to the user at the lowest cost with a supply order to that effect preferably of a government institution.
- 28) All procedures laid down by CVC to be followed in all items which were proprietary in nature.

Member Secretary (0)

Chairman