



Date: 23rd July 2024

REQUEST FOR QUOTATION RFQ N° UNFPA/MNG/RFQ/24/008

Dear Sir/Madam,

We hereby solicit your quotation for the supply of below items to UNFPA Mongolia Country office in Ulaanbaatar, Mongolia.

A. List of items:

- Item 1. IUD removal hook – 24 pcs
- Item 2. Medical instruments for vasectomy set – 2 pcs
- Item 3. Vasectomy training model – 2 pcs
- Item 4. Multi-purpose uterus model (IUD insertion, removal) – 6 pcs

Please refer to the detail of the technical requirement from Annex II.

B. Schedule of Requirements:

- UNFPA has the right to place an order for all or some of the items.
- Partial bid is not allowed. The Bidder shall **be** required to quote for all items.
- Thirty (**30**) calendar days upon approval of the order.
- Incoterms 2020: DAP, UN House, Sukhbaatar district, 6th khoroo, United Nations Street, Ulaanbaatar, Mongolia.

I. About UNFPA

UNFPA, the United Nations Population Fund (UNFPA), is an international development agency that works to deliver a world where every pregnancy is wanted, every childbirth is safe and every young person's potential is fulfilled.

UNFPA is the lead UN agency that expands the possibilities for women and young people to lead healthy sexual and reproductive lives. To read more about UNFPA, please go to: [UNFPA about us](#)

II. Questions

Questions or requests for further clarifications should be submitted in writing to the contact person below:

Name of contact person at UNFPA:	<i>Procurement Officer</i>
Tel N°:	976-11-353503, ext 3355
Email address of contact person:	Batsuuri@unfpa.org

The deadline for submission of questions is Monday, 29th July 2024, 6:00pm (GMT +8). Questions will be answered in writing and shared with all parties as soon as possible after this deadline.

III. Eligible Bidders

This Request for Quotation is open to all eligible bidders; to be considered an eligible bidder for this solicitation process you must comply with the following:



- A bidder must be a legally constituted company that can provide the requested products and have legal capacity to enter into a contract with UNFPA to deliver in the country, or through an authorized representative.
- A bidder must not have a conflict of interest regarding the solicitation process or with the TORs / Technical Specifications. Bidders found to have a conflict of interest shall be disqualified.
- At the time of Bid submission, the bidder, including any JV/Consortium members, is not under procurement prohibitions derived from the [Compendium of United Nations Security Council Sanctions Lists](#) and has not been suspended, debarred, sanctioned or otherwise identified as ineligible by any [UN Organization](#) or the [World Bank Group](#).
- Bidders must adhere to the UN Supplier Code of Conduct, which may be found by clicking on [UN Supplier Code of Conduct](#).

IV. Content of quotations

Quotations should be submitted in a single email whenever possible, depending on file size.

Quotations should contain:

- Technical proposal, in response to the requirements outlined in the specifications should comply with:
 - Offered item overview form showing the technical specifications relevant to the UNFPA requirement, brand name, model ID, actual photo of items offered.
 - Bidder’s previous experience relevant to the current bidding. List of contracts etc.
- Signed Declaration Form, to be submitted strictly in accordance with the document.
- Price quotation, to be submitted strictly in accordance with the price quotation form.
 - The quotation shall be valid at least for (90) days after the closing date.

All forms must be parts of the quotation **must be signed by the company’s relevant authority** and submitted in PDF format!

V. Instructions for submission

Proposals should be prepared based on the guidelines set forth in Section IV above, along with a properly filled out and signed price quotation form and are to be sent by email to the address indicated below no later than: **Monday, 5th August 2024, at 5:00 PM Ulaanbaatar Time**¹.

Name of contact person at UNFPA:	Procurement Officer
Official Email address:	procurement@unfpa.org.mn

Please note the following guidelines for electronic submissions to UNFPA Mongolia’s secured email address:

- The following reference must be included in the email subject line: **RFQ N^o UNFPA/MNG/RFQ/24/008 – Medical items**. Proposals, including both technical and financial proposals, that do not contain the correct email subject line may be overlooked by the procurement officer and therefore not considered.
- The total email size may not exceed **20 MB (including email body, encoded attachments and headers)**. Where the technical details are in large electronic files, it is recommended that these be sent separately before the deadline.
- Please do **NOT** send the emails containing your offer to any other email address (not even as a carbon copy (CC) or blind carbon copy (BCC)); otherwise UNFPA will not be able to guarantee confidentiality and fair and transparent handling of your bid. UNFPA reserves the right to reject bids sent via the appropriate channel but copied or blind copied to other email addresses.
- When submitting electronic offers, Bidders will receive an auto-reply acknowledging receipt of the **first** email. Should your offer require you to submit more than one email, in the body of this first email, bidders are requested to list the number of messages, which make up their technical offer and the number of messages, which make up

¹ <http://www.timeanddate.com/worldclock/city.html?n=69>



their financial offer. If you do not receive any auto-reply for the first email from UNFPA's email system, please inform Procurement official at: batsuuri@unfpa.org.

- Any quotation submitted will be regarded as an offer by the bidder and does not constitute or imply acceptance of the quotation by UNFPA. UNFPA is under no obligation to award a contract to any bidder as a result of this RFQ.

VI. Overview of Evaluation Process

Quotations will be evaluated based on the compliance with the technical specifications and the total cost of the goods (as per price quote).

The evaluation will be carried out in a two-step process by an ad-hoc evaluation panel. Technical proposals will be evaluated for technical compliance prior to the comparison of price quotes.

VII. Award

In case of a satisfactory result from the evaluation process, UNFPA shall award a Purchase Order to the lowest priced bidder whose bid has been determined to be substantially compliant with the bidding documents.

VIII. Right to Vary Requirements at Time of Award

UNFPA reserves the right at the time of award of Contract to increase or decrease, by up to 20%, the volume of goods specified in this RFQ without any change in unit prices or other terms and conditions.

IX. Payment Terms

UNFPA payment terms are net 30 days upon receipt of shipping documents, invoice and other documentation required by the contract.

X. Fraud and Corruption

UNFPA is committed to preventing, identifying, and addressing all acts of fraud against UNFPA, as well as against third parties involved in UNFPA activities. UNFPA's Policy regarding fraud and corruption is available here: [Fraud Policy](#). Submission of a proposal implies that the Bidder is aware of this policy.

Suppliers, their subsidiaries, agents, intermediaries and principals must cooperate with the UNFPA Office of Audit and Investigations Services as well as with any other oversight entity authorized by the Executive Director and with the UNFPA Ethics Advisor as and when required. Such cooperation shall include, but not be limited to, the following: access to all employees, representatives' agents and assignees of the vendor; as well as production of all documents requested, including financial records. Failure to fully cooperate with investigations will be considered sufficient grounds to allow UNFPA to repudiate and terminate the Agreement, and to debar and remove the supplier from UNFPA's list of registered suppliers.

A confidential Anti-Fraud Hotline is available to any Bidder to report suspicious fraudulent activities at [UNFPA Investigation Hotline](#).

XI. Zero Tolerance

UNFPA has adopted a zero-tolerance policy on gifts and hospitality. Suppliers are therefore requested not to send gifts or offer hospitality to UNFPA personnel. Further details on this policy are available here: [Zero Tolerance Policy](#).

XII. RFQ Protest

Bidder(s) perceiving that they have been unjustly or unfairly treated in connection with a solicitation, evaluation, or award of a contract may submit a complaint to the UNFPA Head of Office, Khalid Sharifi at ksharifi@unfpa.org. Should the supplier be unsatisfied with the reply provided by the UNFPA Head of the Business Unit, the supplier may contact the Chief, Supply Chain Management Unit at supplychain@unfpa.org.

XIII. Disclaimer



United Nations Population Fund
UNFPA Mongolia Country Office
Email: contact@unfpa.org.mn
Website: www.unfpa.org

Should any of the links in this RFQ document be unavailable or inaccessible for any reason, bidders can contact the Procurement Officer in charge of the procurement to request for them to share a PDF version of such document(s).



PRICE QUOTATION FORM

Name of Bidder:	
Date of the quotation:	Click here to enter a date.
Request for quotation N°:	UNFPA/MNG/RFQ/24/008
Currency of quotation:	
Validity of quotation:	<i>(The quotation shall be valid for a period of at least 3 months after the submission deadline.)</i>

Price Quotation Form

No	Product Name & Description	Unit Price (DAP)	Unit of measure	Quantity	Total price (DAP)	Delivery schedule
Item 1.	IUD removal hook		Each	24		
Item 2.	Medical instruments for vasectomy set		Each	2		
Item 3.	Vasectomy training model		Each	2		
Item 4.	Multi-purpose uterus model (IUD insertion, removal)		Each	6		
Grand total						

- DAP point: United Nations Population Fund (UNFPA) Mongolia country office, UN House, UN Street-14, Ulaanbaatar 14201, Mongolia
- For local suppliers, please indicate in the Vendor's comment section below if the unit price includes the Value Added Tax (VAT). Otherwise, no indication will be understood as the unit price are exclusive of VAT.

<i>Vendor's Comments:</i>

I hereby certify that the company mentioned above, which I am duly authorized to sign for, has reviewed RFQ UNFPA/MNG/RFQ/24/008 including all annexes, amendments to the RFQ document (if applicable) and the responses provided by UNFPA on clarification questions from the prospective service providers. Further, the company accepts the General Conditions of Contract for UNFPA, and we will abide by this quotation until it expires.

	Click here to enter a date.
Name and title	Date and place



DECLARATION FORM

UNFPA.MNG.RFQ.24.008

The undersigned, being a duly authorized representative of the Company represents and declares that:

		YES	NO
1.	The Company and its Management ² have not been found guilty pursuant to a final judgement or a final administrative decision of any of the following:		
	a. Fraud;	<input type="checkbox"/>	<input type="checkbox"/>
	b. Corruption;	<input type="checkbox"/>	<input type="checkbox"/>
	c. conduct related to a criminal organization;	<input type="checkbox"/>	<input type="checkbox"/>
	d. money laundering or terrorist financing;	<input type="checkbox"/>	<input type="checkbox"/>
	e. terrorist offences or offences linked to terrorist activities;	<input type="checkbox"/>	<input type="checkbox"/>
	f. sexual exploitation and abuse;	<input type="checkbox"/>	<input type="checkbox"/>
	g. child labour, forced labour, human trafficking; or	<input type="checkbox"/>	<input type="checkbox"/>
	h. irregularity (non-compliance with any legal or regulatory requirement applicable to the Organization or its Management).	<input type="checkbox"/>	<input type="checkbox"/>
2.	The Company and its Management have not been found guilty pursuant to a final judgment or a final administrative decision of grave professional misconduct.	<input type="checkbox"/>	<input type="checkbox"/>
3.	The Company and its Management are not: bankrupt, subject to insolvency or winding-up procedures, subject to the administration of assets by a liquidator or a court, in an arrangement with creditors, subject to a legal suspension of business activities, or in any analogous situation arising from a similar procedure provided for under applicable national law.	<input type="checkbox"/>	<input type="checkbox"/>
4.	The Company and its Management have not been the subject of a final judgment or a final administrative decision finding them in breach of their obligations relating to the payment of taxes or social security contributions.	<input type="checkbox"/>	<input type="checkbox"/>

² "Management" means any person having powers of representation, decision-making or control over the Organization. This may include, for example, executive management and all other persons holding downstream managerial authority, anyone on the board of directors, and controlling shareholders.



5.	The Company and its Management have not been the subject of a final judgment or a final administrative decision which found they created an entity in a different jurisdiction with the intent to circumvent fiscal, social or any other legal obligations in the jurisdiction of its registered office, central administration, or principal place of business (<i>creating a shell company</i>).	<input type="checkbox"/>	<input type="checkbox"/>
6.	The Company and its Management have not been the subject of a final judgment or a final administrative decision which found the Company was created with the intent referred to in point (5) (<i>being a shell company</i>).	<input type="checkbox"/>	<input type="checkbox"/>

The UNFPA reserves the right to disqualify the Company, suspend or terminate any contract or other arrangement between the UNFPA and the Company, with immediate effect and without liability, in the event of any misrepresentation made by the Company in this Declaration.

It is the responsibility of the Company to immediately inform the UNFPA of any changes in the situations declared above.

This Declaration is in addition to, and does not replace or cancel, or operate as a waiver of, any terms of contractual arrangements between the UNFPA and the Company.

Signature:

Date:

Name and Title:

Name of the Company:

UNGM N°:

Postal Address:

Email:



**ANNEX I:
General Conditions of Contracts:
De Minimis Contracts**

This Request for Quotation is subject to UNFPA's General Conditions of Contract: De Minimis Contracts, which are available in: [English](#), [Spanish](#) and [French](#)

Please note that a PDF version of the applicable General Conditions of Contracts must be provided.

ANNEX II: Technical specification

Item No	Item name	Requirement	Quantity
Item 1	IUD removal hook	<p>Product Description The Universal IUD Hook is a medical instrument designed for intrauterine device (IUD) insertion and removal procedures. The instrument is atraumatic, ensuring minimal tissue damage and patient discomfort during procedures. Its curved design allows for optimal access and maneuverability within the uterine cervix. Classified under EU MDR 2017/745 as Class I device</p> <p>Technical specifications:</p> <ul style="list-style-type: none"> ● Material: High-grade stainless steel (AISI 304) ● Total Length: 10 inches (25 cm) ● Hook Diameter: 4 mm ● Design: Curved for enhanced access and maneuverability, smooth, polished surface to prevent tissue trauma. ● Ergonomics: Handle designed for a comfortable grip and precise control ● Sterilization: Autoclavable, compatible with standard sterilization procedures. ● Packaging & Labelling: <ul style="list-style-type: none"> ◦ Single unit packaging or kit packaging with Symbols used according to ISO 15223. ◦ Labelling as per regulations applied to the device. Shall include name of the device, name and address of the manufacturer, product code or manufacturers reference number. <p>Documentation requirement:</p> <ul style="list-style-type: none"> ● Instructions for Use (IFU) should be provided with comprehensive instructions on the use, cleaning, disinfection and sterilization methods and types for the product. ● Warranty Certificate: At least one-year warranty covering manufacturing defects. ● Manufacturing license <p>Regulatory approvals required:</p> <ul style="list-style-type: none"> ● CE mark Comply to EU Council Directive 93/42/EEC (MDD) or EU Regulations 2017/745 (MDR) with valid EC Certificate and Declaration of Conformity for the CE mark is applicable. ● ISO 13485 Medical Devices - Quality Management Systems ● ISO 14001 Environmental Management ● ISO 50001 Energy Management Systems 	24

		<p>Safety & product Standards Must comply with the following standards (as per latest revisions of the standards):</p> <ul style="list-style-type: none"> ● ISO 7153-1:1991 Surgical Instruments – Metallic Materials ● ISO 17664:2004 Sterilization of Medical Devices - Information to be Provided by the Manufacturer for the Processing of Resterilizable Medical Devices ● ISO 13402:1995 Surgical and Dental Hand Instruments - Determination of Resistance Against Autoclaving, Corrosion, and Thermal Exposure 	
<p>Item 2</p>	<p>Medical instruments for vasectomy set</p>	<p>Product Description A vasectomy set typically includes a range of specialized instruments designed to facilitate the vasectomy procedure, which is a form of male sterilization.</p> <p>Technical specifications:</p> <p>A. Scalpel Handle and Blades</p> <ul style="list-style-type: none"> ○ Scalpel Handle: Size 3, made of stainless steel AISI 304. ○ Scalpel Blades: Size 10 or 15, sterile and disposable. Stainless steel AISI 420. <p>B. Straight Hemostatic Forceps</p> <ul style="list-style-type: none"> ○ Type: Mosquito forceps ○ Size: 5 inches (12.5 cm), straight ○ Material: Stainless steel AISI 410, atraumatic, serrated tips <p>C. Curved Hemostatic Forceps</p> <ul style="list-style-type: none"> ○ Type: Kelly or Crile forceps ○ Size: 5.5 inches (14 cm), curved ○ Material: Stainless steel AISI 410, atraumatic, serrated tips <p>D. Ring-Tip Forceps</p> <ul style="list-style-type: none"> ○ Type: Ring-tipped (also known as Allis forceps) ○ Size: 5.5 inches (14 cm) ○ Material: Stainless steel AISI 410, atraumatic, with fine tips for grasping the vas deferens <p>E. Needle Holder</p> <ul style="list-style-type: none"> ○ Type: Mayo-Hegar or similar ○ Size: 5 inches (12.5 cm) ○ Material: Stainless steel AISI 410, with a locking mechanism <p>F. Suture Scissors</p>	<p>2</p>

		<ul style="list-style-type: none"> ○ Type: Straight or curved surgical scissors ○ Size: 4.5 inches (11.5 cm) ○ Material: Stainless steel AISI 420 or 440, fine tips <p>G. Iris Scissors</p> <ul style="list-style-type: none"> ○ Type: Curved or straight ○ Size: 4.5 inches (11.5 cm) ○ Material: Stainless steel AISI 420 or 440, fine tips <p>H. Vasectomy Clamp</p> <ul style="list-style-type: none"> ○ Type: Ringed clamp or No-Scalpel Vasectomy (NSV) clamp ○ Size: Varies, typically 5-6 inches (12.5-15 cm) ○ Material: Stainless steel AISI 410, designed to securely hold the vas deferens during the procedure <p>I. Dissecting Forceps</p> <ul style="list-style-type: none"> ○ Type: Adson forceps with teeth or without teeth ○ Size: 4.75 inches (12 cm) ○ Material: Stainless steel AISI 410, fine tips for precise tissue handling <p>J. Vasectomy Hook</p> <ul style="list-style-type: none"> ○ Type: Curved hook ○ Size: 5 inches (12.5 cm) ○ Material: Stainless steel, atraumatic <p>Documentation requirement:</p> <ul style="list-style-type: none"> ● Instructions for Use (IFU) should be provided with comprehensive instructions on the use, cleaning, disinfection and sterilization methods and types for the product. ● Warranty Certificate: At least one-year warranty covering manufacturing defects. ● Manufacturing license <p>Regulatory approvals required:</p> <ul style="list-style-type: none"> ● CE mark Comply to EU Council Directive 93/42/EEC (MDD) or EU Regulations 2017/745 (MDR) with valid EC Certificate and Declaration of Conformity for the CE mark is applicable. ● ISO 13485 Medical Devices - Quality Management Systems ● ISO 14001 Environmental Management ● ISO 50001 Energy Management Systems <p>Safety & product Standards:</p>	
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		<p>Must comply with the following standards (as per latest revisions of the standards):</p> <ul style="list-style-type: none"> ● ISO 7153-1:1991 Surgical Instruments – Metallic Materials ● ISO 17664:2004 Sterilization of Medical Devices - Information to be Provided by the Manufacturer for the Processing of Resterilizable Medical Devices ● ISO 13402:1995 Surgical and Dental Hand Instruments - Determination of Resistance Against Autoclaving, Corrosion, and Thermal Exposure 	
<p>Item 3</p>	<p>Vasectomy training model</p>	<p>Product Description A vasectomy training model, that simulates an adult male genital structure, with a realistic shape and texture, designed for no-scalpel vasectomy operation training.</p> <p>The vasectomy model grants the development of competence-based training sessions, allowing practice and learning skills to develop no-scalpel vasectomy.</p> <p>Technical specifications:</p> <ul style="list-style-type: none"> ● Anatomical model composed by: <ul style="list-style-type: none"> ○ <i>Male torso section</i>, modeling lower abdomen to above-knee stump. ○ <i>Removable scrotal skins</i>. ○ <i>Testicle assembly</i> with tubing attached, simulating the vas deferens. ● External skin tone: light to medium skin tone. ● Male torso section, with at least four (4) anti-slip feet/pads. ● Removable scrotal skin, flexible and penetrable to be pierced and to access the vas deferens simulation tubing. ● Male torso section, Removable scrotal skin and Testicle assembly must be built of materials resistant to the application of disinfectants commonly used in healthcare premises, i.e.: 70% isopropyl alcohol. <p>Accessories:</p> <ul style="list-style-type: none"> ● Two (2) reusable removable scrotal skins <ul style="list-style-type: none"> ○ One (1) included in the standard configuration. ○ The rest to be used as spare parts. ● Two (2) reusables set of two testicle assemblies <ul style="list-style-type: none"> ○ One (1) set included in the standard configuration. ○ The rest to be used as spare parts. ● Two (2) sets of vas deferens simulation tubing with securing/fastening mechanism. <ul style="list-style-type: none"> ○ One (1) included in the standard configuration. ○ The rest to be used as spare parts. <p>Documentation requirement:</p>	<p>2</p>

		<ul style="list-style-type: none"> ● Bidders shall furnish the following documents as a part of the Bid, in English language: <ul style="list-style-type: none"> ○ Manufacturer brochure or data sheet including at least all technical specifications required. ○ Instructions for use including procedures for cleaning, disinfection and storage conditions. ○ List of all common spare parts and accessories with part numbers. ○ Manufacturer authorization letter. ○ Manufacturer letter of commitment including: <ul style="list-style-type: none"> ■ Installation and commissioning on site would be performed by the manufacturer or the representative in the country. ■ At least one (1) year of full onsite warranty. ■ At least five (5) years of spare parts availability. ■ Training on use, cleaning and disinfecting for the healthcare staff, in Mongolian language. ● Bidders awarded, during the commissioning shall deliver the following documents, in English, Mongolian and/or another requested language: <ul style="list-style-type: none"> ○ Instructions for use including procedures for cleaning, disinfection and storage conditions. ○ List of all common spare parts and accessories with part numbers. ○ Warranty certificate. ○ Manufacturer letter with at least five (5) years of spare parts availability. <p>Regulatory approvals required:</p> <ul style="list-style-type: none"> ● Bidders shall furnish the following documents as a part of the Bid: <ul style="list-style-type: none"> ○ Valid manufacturing license, issued by the corresponding authority in the manufacturing country. <p>Safety & product Standards:</p> <ul style="list-style-type: none"> ● Valid ISO 9001, Quality management systems certificate, issued by Conformity assessment bodies, notified bodies or by bodies accredited by regulatory authorities. 	
Item 4	Multi-purpose uterus model (IUD insertion, removal)	<p>Product Description</p> <p>An uterus training model, that simulates an adult female genital structure, with a functional shape and texture, is designed for multi-purpose training.</p> <p>The uterus training model grants the development of competence-based training sessions, allowing practice and learning skills to develop skills like vaginal examination, pap smear sample collection, IUD insertion, removal of retained products of conception by using manual vacuum aspiration (MVA).</p>	6

		<p>Technical specifications:</p> <ul style="list-style-type: none"> ● Model representing female abdominal section. ● Design allows configuring the model in two modes: <ul style="list-style-type: none"> ○ Simulation, to be fastened around the body of a person. ○ Training, to be used as a standalone educational material. ● Inside the model, the uterus is adjustable allowing it to be configured in a retroverted and anteverted angles. ● At least two sized uterus: <ul style="list-style-type: none"> ○ Small, representing an early post-abortion uterus. ○ Large, representing a large uterus like a multiparous and/or a post abortion uterus. ● Must be built of easily cleanable materials. <p>Accessories:</p> <ul style="list-style-type: none"> ● One (1) carrying case, from the same manufacturer or expressly recommended by the manufacturer of the multi-purpose uterus model. ● One (1) strap to fasten the model around the body. ● One (1) system or device to keep the multi-purpose uterus model in position during training sessions. <p>Documentation requirement:</p> <ul style="list-style-type: none"> ● Bidders shall furnish the following documents as a part of the Bid, in English language: <ul style="list-style-type: none"> ○ Manufacturer brochure or data sheet including at least all technical specifications required. ○ Instructions for use including procedures for cleaning, disinfection and storage conditions. ○ List of all common spare parts and accessories with part numbers. ○ Manufacturer authorization letter. ○ Manufacturer letter of commitment including: <ul style="list-style-type: none"> ■ Commissioning on site would be performed by the manufacturer or the representative in the country. ■ At least one (1) year of full onsite warranty. ■ At least five (5) years of spare parts availability. ■ Training on use, cleaning and disinfecting for the healthcare staff, in Mongolian language. ● Bidders awarded, during the commissioning shall deliver the following documents, in English, Mongolian and/or another requested language: <ul style="list-style-type: none"> ○ Instructions for use including procedures for cleaning, disinfection and storage conditions. ○ List of all common spare parts and accessories with part numbers. ○ Warranty certificate. 	
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		<ul style="list-style-type: none">○ Manufacturer letter with at least five (5) years of spare parts availability. <p>Regulatory approvals required:</p> <ul style="list-style-type: none">● Bidders shall furnish the following documents as a part of the Bid:<ul style="list-style-type: none">○ Valid manufacturing license, issued by the corresponding authority in the manufacturing country. <p>Safety & product Standards:</p> <ul style="list-style-type: none">● Valid ISO 9001, Quality management systems certificate, issued by Conformity assessment bodies, Notified bodies or by bodies accredited by regulatory authorities.	
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