Procurement of equipment for General Hospital in Loznica

NO. IOP/61-2021/UHI

Clarification No.1

Issued on 5th of October 2022

Question 1:

Lot 9 – Hospital Furniture

We are a procurement company in Turkey and we are planning to participate your tender.
We have analyzed the technical spesifications document for Lot 9 and we have some questions about the technical specifications document to optimise the calculation of the cost and production.
1.Do you have drawings or sample photograph of the demanded goods and if you have could you please share them with us.
2.What is the diameter and thickness of metal parts and their type (L,square,round etc.)?
We look forward to hear from you."

Answer 1:
1. No, we do not have drawings or sample photograph of the demanded goods.
2. Please, be precise about which item within Lot 9 you ask for the diameter and thickness of metal parts and their type (L, square, round etc.).

Question 2:

Lot 5 – Radiology Imaging

Item No. 5
Under ID 5.21 is requested: Biopsy convex probe.
Does the Purchaser mean by this request that bidder should offer convex probe with a biopsy guide?

Answer: Yes, system should support convex probe with biopsy guide.

Under ID 5.22 is requested: Volume probes (convex array, micro convex array, linear array).
Are all probe types listed in bracket mandatory, or they are only mentioned as examples? Is it acceptable to offer a system which supports volume convex and microconvex probes?

Answer: Yes, it is acceptable.

Item No. 9
Purchaser asked for minimal technical features under ID 9.39-9.41. same features that are already requested under ID 9.36-9.38. As it is obviously that it is technical mistake, Purchaser should change tables with technical specification accordingly with deleting ID 9.39.-9.41."

Answer: Yes, it is a technical mistake, bidder should offer one linear probe whose characteristics are listed under Id line 9.38. Mentioned will be amended through Amendment no. 1.

Question 3:

1. We kindly ask the Purchaser to confirm that Manufacturer's Sales Authorization and Manufacturer's After Sales Authorization as well as the Manufacturer's Technical Statement requested in Tender documents can also be signed by the authorized EU or Regional representative of the Manufacturer.

Answer:

Yes, it would be also acceptable to submit the Manufacturer's Authorization and the Manufacturer's After Sales Authorization signed by the European or Regional representative of the Manufacturers.

2. In the Tender documents the Purchaser requests Manufacturer's Sales Authorization and Manufacturer's After Sales Authorization and that service company shall employ minimum number of qualified persons – certified by the manufacturer of equipment for servicing - 1(one) per item model offered. Please confirm that these documents should be issued only by manufacturers of the main equipment i.e. Items listed in the Price Schedule sheet of Technical Specifications of each LOT and not by the manufacturers of additional equipment listed under items?

Answer:

Manufacturer`s Authorisation for Sales and Aftersales, for Bidder and Service Company and service certificate for item model offered should be submitted only from manufacturer of the main system (equipment).

3. The Purchaser requests delivery period of maximum 120 days from the date of the advance payment. Considering that Covid-19 restrictions are still in force in some countries, worldwide lack of semiconductors and unpredictable situation with gas and energy supply due to crisis in Ukraine, delivery times became longer and longer with many vendors and production facilities. We kindly ask the Purchaser to change the requirement to: Delivery period of maximum 180 days from the date of the advance payment.

Answer:

Purchaser accepts explanation and revised request for a delivery period of maximum of 180 days. Mentioned will be amended through Amendment no. 1.

4. In the General Technical Requirements the Purchaser requested that the equipment offered should be manufactured in compliance with Quality Standard ISO 9001 certification for Manufacturer(s). We kindly ask the Purchaser to allow Quality Standard ISO 9001 or ISO 13485 certification for Manufacturer(s).

Answer:
Yes, it is allowed.

Question 4:

Lot - Hospital Furniture

In the Technical Specifications the Purchaser allowed Manufacturers Statement to be used as a proof only in case where requested parameter is not stated in official manufacturer data sheet. Considering that in LOT9_HospitalFurnirure a lot of items are furniture that will be custom made according to the required technical specification, therefore there are no official manufacturer data sheet/catalogues, could you please confirm that for those items the Manufacturers Statement can be used as proof for all the parameters of the above-mentioned items?

Answer 4:

Yes, we confirm that the manufacturer's Statement can be used as proof for items where the Supplier doesn't have manufacturer data sheets/catalogs, having in mind that in Lot 9 - HospitalFurnirure a lot of items are furniture that will be custom made according to the required technical specification.

Public Procurement Committee