

**NOTICE ON CHANGE
OF DELIVERY REQUIREMENTS — COMPARATIVE DATA TABLE AND
SCHEDULE OF REQUIREMENTS**

TENDER: RFQ

BENEFICIARY COUNTRY: Serbia

REFERENCE NUMBER: UNOPS-EP-2016-G-001

DESCRIPTION: Purchase of Dental Equipment In Support Of Women Entrepreneurship

UNCCS CODE: n/a

POSTING DATE: 28 January 2016

- 1) The following sentence is added to the **Delivery requirements — Comparative Data Table** in the section **“Delivery place and Incoterm rules”** (pg. 26):

“Offload, assembly and installation of the equipment on the beneficiaries’ premises are included in the price.”

Now therefore, Delivery requirements — Comparative Data Table is changed to read as follows:

Delivery requirements — Comparative Data Table

UNOPS Requirements		Is quotation compliant? Bidder to complete	Details Bidder to complete Insert details
Delivery time	Bidder shall deliver the goods 45 (Forty Five) calendar days after Contract signature.	<input type="checkbox"/> Yes <input type="checkbox"/> No	
Delivery place and Incoterms rules	DAP (Delivered at Place) as per Incoterms 2010. Offload, assembly and installation of the equipment on the beneficiaries’ premises are included in the price. Four beneficiaries in the Municipalities of Aleksinac, Blace, Brus and Raska. Net of any direct taxes, customs duties and indirect taxes and VAT.	<input type="checkbox"/> Yes <input type="checkbox"/> No	
Consignee details	The exact address the goods should be delivered and offloaded to, as well as all other relevant details, will be	<input type="checkbox"/> Yes <input type="checkbox"/> No	

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	available upon contract signature.		
UNOPS Right to vary requirements	At the time the Contract is awarded, UNOPS reserves the right to vary the quantity of the goods and associated services specified above, provided this does not exceed +/- 20%, without any change in the unit prices or other terms and conditions of the RFQ.	<input type="checkbox"/> Yes <input type="checkbox"/> No	

The offered goods and related services (if applicable) are in accordance with the required specifications and requirements specified in Section III: Schedule of Requirements.

Yes No

ANY DEVIATION MUST BE LISTED BELOW:

Name: _____

Title: _____

Date: _____

Signature: _____



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2) The following sentences are added to the **Note to Section III: Schedule of Requirements (pg. 9)**:

“The Contractor will be responsible for providing after sales support through the standard manufacturer’s warranty.

All the products must be in compliance with the Law on Medicinal Products and Medical Devices (published in the “Official Gazette of the Republic of Serbia” No. 30/2010 dated 7 May 2010) .

All the products must be duly listed in the appropriate registers administered by the Medicines and Medical Devices Agency in Serbia (Agencija za lekove i medicinska sredstva).”

Now therefore, Note to Section III: Schedule of Requirements (pg. 9) is changed to read as follows:

NOTE: The offered equipment must comply with requirements laid down in the Technical specifications (Form C: Technical Quotation Form - Technical specifications for goods – Comparative Data Table). The required specifications present minimum requirements. Bidders may offer items with more advanced specs and such offers are accepted as appropriate.

Manufacturer’s technical literature submitted with the offer must comply with the written specifications of the Bidder. In the event that there are differences between the submitted manufacturers’ literature and written Bidder specification reasons for that must be explained in the Bidder specifications. Manufacturer’s technical literature (brochures, booklets, instructions, etc.) should be submitted. The manufacturers’ technical literature should be marked in an appropriate manner (i.e. model number).

Bidder warrants that offered equipment is new and unused. Bidder shall further warrant that none of the supplies have any defect arising from design, materials or workmanship.

The Contractor will be responsible for providing after sales support through the standard manufacturer’s warranty.

Where applicable, items must be in compliance with the Law on health safety of products for general use Republic of Serbia.

All the products must be in compliance with the Law on Medicinal Products and Medical Devices (published in the “Official Gazette of the Republic of Serbia” No. 30/2010 dated 7 May 2010) .

All the products must be duly listed in the appropriate registers administered by the Medicines and Medical Devices Agency in Serbia (Agencija za lekove i medicinska sredstva).

