T385 COLUMN

CLARIFICATIONS No.8

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ICB No. IOP/3-2015/NCE/1

Project: Public Sector Research and Development

Subproject: New Capital Equipment

Procurement: Procurement of Capital Equipment

| | Questions dated: 08/09/2015 | | |
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| If in situation when foreign company is supplier, customs clearance procedure and payments (including custom duties and other import taxes) are under Purchaser responsibility. Therefore, Purchaser prepares all documentation. It is recommended that non-resident supplier does not send the goods to Serbia until they obtain the consent from Purchaser which says that Purchaser prepared the documentation so that the goods would not be detained at customs. After customs clearance, the supplier proceeds the transport to the delivery place. | On the page 32 of the tender dossier there is a condition that "The Purchaser shall not pay any amount of Custom Duties and other Import Taxes". But if non-resident company supplies equipment on the DAP terms, all customs clearance procedure and payments (including custom duties and other import taxes) are under Purchaser responsibility. Please clarify. | Page 32 ITB 14.8 (b) (i) and (c) (v) | · N |
| See Corrigendum no. 5 | On the page 44 of the tender dossier there is requirement to provide for Bidder Certificates such as ISO 9001:2008. As we are official representative of the Manufacturer for the sale and technical support, can we provide only Manufacturer's ISO 9001:2008 | Tender dossier page 44 | Н |
| | Questions dated: 04/09/2015 | | |
| Answers | Questions | Reference in PN/ Tender dossier | Qtns. No. |

| in your bid. | ISO 13485:2003 specifies requirements for a quality management system where an organization needs to demonstrate its ability to provide medical | |
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| It is not necessary to submit copy of manufacturer's ISO 13485 and ISO 14001 | Considering that: | |
| 9001:2000 certification for Manufacturer(s), ISO 13485 and ISO 14001 in order to achieve the international accreditation according to EN ISO/IEC 17025:2005, EN 45003. | 2.2. The equipment offered should be manufactured in compliance with Quality Standard ISO 9001:2000 certification for Manufacturer(s), ISO 13485 and ISO 14001 in order to achieve the international accreditation according to EN ISO/IEC 17025:2005, EN 45003. | Tender dossier, General Technical Requirements, 2. Equivalency of Standards and Codes |
| The equipment should be manufactured in compliance with Quality Standard ISO | Reviewing the Bidding Document, we have found that under General Technical Requirements, 2. Equivalency of Standards and Codes you have requested that: | |
| | Can we use our Bank variant? | |
| | This guarantee will expire: (a) if the Applicant is the successful bidder, upon our receipt of copies of the contract agreement signed by the Applicant and the performance security issued to the Beneficiary in relation to such contract agreement; or (b) if the Applicant is not the successful bidder, upon the earlier of (i) our receipt of a copy of the Beneficiary's notification to the Applicant of the results of the bidding process; or (ii) twenty-eight days after the end of the Bid Validity Period; but in any cases the guarantee will expire not later than 01.03.2016. | |
| The answer is yes. You can use your Bank variant. | But our Bank (taking into account 05.10.15 as a new deadline for bid submission and 120 + 28 days) suggests modified variant: | 3 "Form of Bid Security/Bank Guarantee" |
| | "This guarantee will expire: (a) if the Applicant is the successful bidder, upon our receipt of copies of the contract agreement signed by the Applicant and the performance security issued to the Beneficiary in relation to such contract agreement; or (b) if the Applicant is not the successful bidder, upon the earlier of (i) our receipt of a copy of the Beneficiary's notification to the Applicant of the results of the bidding process; or (ii) twenty-eight days after the end of the Bid Validity Period." | Tender dossier page 57-58 |
| | On the page 58 of the tender dossier "Form of Bid Security/Bank Guarantee" there are options for guarantee explorations: | |

| See Corrigendum no. 1 | On the page 44, article 2.3.1, the ordering party stipulated that the way of asserting professional capacity should be by delivery of certificates such as ISO 9001:2008 and 27001:2005 (or 27001:2013). - Would the ordering party find it acceptable to change this part of the competition documentation, and request only the delivery of the certificate ISO 9001:2008 since the majority of potential bidders do not have ISO 27001:2005 certificate? In this way, the ordering party would secure more potential bidders, competitiveness, as well as the possibility to receive the best offer. | Tender dossier page 44., article 2.3.1 | 7 |
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| See Corrigendum no. 1 and no. 5 Bidders should submit ISO 9001:2008. | On the page 44, article 2.3.1, the ordering party stipulated that the way of asserting professional capacity should be by delivery of certificates such as ISO 9001:2008 and 27001:2005 (or 27001:2013). - Does the ordering party, under "such as", refer to the possibility of delivery of either one or the other certificate? In case that it is possible to deliver similar certificates, we want to ask the ordering party to specify which certificates, apart from the listed ones, would be accepted as "such as". | Tender dossier page 44., article 2.3.1 | 6 |
| See Corrigendum no. 5 | On the page 44, article 2.3.1, the ordering party stipulated that the way of asserting professional capacity should be by delivery of certificates such as ISO 9001:2008 and 27001:2005 (or 27001:2013). - By this, does the ordering party refer to ISO certificates issued on the name of the potential bidder or on the name of the manufacturer whose equipment the potential bidder is offering? | Tender dossier page 44., article 2.3.1 | 5 |
| | devices and related services that consistently meet customer requirements and regulatory requirements applicable to medical devices and related services, and The ISO 14000 family of standards provides practical tools for companies and organizations of all kinds looking to manage their environmental responsibilities. Is it necessary to submit manufacturer's ISO 13485 and ISO 14001 in our bid? If it is, could you please clarify if that requirement applies to all lots? Or ISO 13485 is required only if the offered equipment are medical devices? Could you also please clarify if it is acceptable that bidder's ISO 14001 is submitted, instead of manufacturer's ISO 14001 certificates? | | |

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