AMENDMENT No. 4 TO PROCUREMENT DOCUMENTS

Issued on 04th of November 2022

PROCUREMENT OF EQUIPMENT FOR GENERAL HOSPITAL IN LOZNICA

NO. IOP/61-2021/UHI

In accordance with the Clause 8. Part 1. Bidding Procedures, Section I. Instructions to Bidders, Contents of Bidding Documents, Amendment of Bidding Documents of the Procurement Documents, Public Investment Management Office, No. 11 Nemanjina street, Republic of Serbia, as the Purchaser, hereby notifies all persons concerned for **Procurement of equipment for General Hospital in Loznica, no. IOP/61-2021/UHI**, that there has been an amendment made in the Procurement Documents.

The following provisions of Tender Documents are hereby replaced as follows:

1. Tender Documents are changed in Technical Specification, Lot 10 –Sterilization, Item 20. Steam sterilizer (autoclave):

- **ID 20.15 which read as follows:**

"Outer dimensions: 470 x 400x 630 cm ± 5%"

It is replaced and reads as follows:

"Outer dimensions: 470 x 400x 630 mm ± 5%"

2. Tender Documents are changed in Technical Specification, Lot 3 – Anesthesia and ICU, Item 6 – Intensive care monitoring,

- ID 6.9. which read as follows:

"Thermal recorder that can be connected to any of the monitors"

It is replaced and reads as follows:

"One thermal recorder in total that can be connected to any of the monitors"

3. Tender Documents are changed in Technical Specification, Lot 7 – Mobile Diagnostic equipment, Item 9 – Channel ECG:

- ID 9.16. which reads as follows:

"It must be constructed with the EU safety standards (class CF according to the guidance IEC 60601-2-25:2011) and have a CE mark qualification"

It is replaced and reads as follows:

"It must be constructed with the EU safety standards (class CF according to the guidance IEC 60601-2-25)"

4. Tender Documents are changed in Technical Specification, Lot 7 – Mobile Diagnostic equipment, Item 9 – Channel ECG:

- ID 9.17. which reads as follows:

"It must be constructed with the EU safety standards (class CF according to the guidance IEC 60601-2-25:2011) and have a CE mark qualification.":

ID 9.17. - It must be constructed with the EU safety standards (class CF according to the guidance IEC 60601-2-25:2011) and have a CE mark qualification – **will be deleted.**

5. Tender Documents are changed in Technical Specification, Lot 7 – Mobile Diagnostic equipment, Item 10 – 12 Channel ECG Analyses programm with measurement, interpretation and rolling stand:

ID 10.1. which reads as follows:

"Appropriate for hospital use to newborns, infants and kids. It must provide the "CE" mark and comply with the international safety standards, in particular by the EN 60601-1-25:2015 for ECG."

It is replaced and reads as follows:

"Appropriate for hospital use to newborns, infants and kids. It must comply with the international safety standards, in particular by the EN 60601-2-25: for ECG"

Public Procurement Committee