

The Supplier's Obligation to Report on the Performance of the Framework Agreement in Joint Procurement Procedures in Healthcare

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American Chamber of Commerce in Croatia Američka gospodarska komora u Hrvatskoj

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Introduction

During the implementation of joint public procurement procedures in healthcare, in mid-2023, suppliers began to notice in the procurement documentation a provision on the supplier's obligation to deliver to the contracting authority a report on the performance of the framework agreement.

At the same time, the reporting obligation is not specified in the PD section that prescribes the conditions that the supplier must fulfill, only in the forms attached to the PD or in the section on the definition of prices.

Example:

The selected bidder shall submit to the Contracting Authority a semi-annual report on the performance of the Framework Agreement, in electronic form, for each individual User.

The report that the selected bidder shall submit must at least contain information about the User, the type and name of the delivered goods, the quantity, the unit and total prices, and must be in the form of a bill of quantities. The reports must be submitted in an electronically editable form (Excel table).

In addition to the delivery of semi-annual reports, the Selected Bidder shall also notify the Contracting Authority in writing when the total contractual obligations of all Users under the Framework Agreement reach 75% of the Framework Agreement value and after they reach the Framework Agreement value.

Given that framework agreements are concluded for a term of 2 years, such a provision creates additional obligations for an economic entity that are in no way included in the bid price.

The economic entity must:

- submit 4 semi-annual reports during the term of the framework agreement;
- control the performance of all individual contracts and when the total value of individual contracts reaches 75% or 100% of the framework agreement value;
 - submit a report on the fulfillment of 75% of the framework agreement value;
 - submit a report on 100% fulfillment of the framework agreement.

Pursuant to Article 313, paragraph 2 of the Public Procurement Act, the **public contracting authority** is obliged to control whether the performance of the public procurement contract is in accordance with the conditions specified in the procurement documentation and the selected bid.

Therefore, the obligation to control the performance of the contract/framework agreement is a legal obligation of the contracting authority, not of the bidder.



Furthermore, the Instructions for the implementation of the joint public procurement procedure issued by the Ministry of Health in May 2022 explicitly state:

"The joint contracting authority/central body is responsible for controlling the performance of the Framework Agreements concluded, the need for possible changes during their term, the receipt and control of performance guarantees, and the timely planning and initiation of a new joint procedure in order to ensure the continuity of contracts without time gaps from the expiration of the valid until the conclusion of new framework agreements for the same procurement categories."

It is unclear why, after a year of joint public procurement, the Ministry gave up its own recommendations to the central authorities and started to impose reporting obligations in joint procurement procedures in which it is the principal body¹. In addition to its own procedures, it also began to demand the same from other central bodies in charge of joint procurement in healthcare, which is contrary to the recommendations of May 2022.

The actions the central bodies have undertaken are also inconsistent in relation to:

- bidder's requests to change procurement documentation in the part that refers to the reporting obligation;
- the Ministry of Health's instructions on the inclusion of the supplier's reporting obligation in the procurement documentation.

A number of bidders warned the contracting authorities that according to the Public Procurement Act, Article 313, it is the contracting authority's obligation to control that the performance of the contract/framework agreement is in accordance with the conditions prescribed in the documentation and, accordingly, requested the deletion of this obligation from the PD; however, the majority of the contracting authorities did not accept the request and explained that both the bidder and the contracting authority are obliged to control the performance of the contract.

Some contracting authorities saw the foundation of such an obligation in Article 11 of the Regulation on internal organization of the State Office for Central Public Procurement, which mentions the scope of work of the Department for the Management of Contracts and Framework Agreements². That article states that the contracting authority is obliged to control the performance of contracts and framework agreements, but since it does not clearly define the mode of control, the contracting authorities and the State Office for Central Public Procurement believe that the

CHIF Procurement of special groups of medications, publication number in EPPC - 2023/S 0F2-0022606



¹Implantation material and consumables for interventional neuroradiology for healthcare institutions in the Republic of Croatia, publication number in EPPC - 2023/S 0F2-0013906;

Implantation material and consumables for invasive and interventional cardiology for healthcare institutions in the Republic of Croatia, publication number in EPPC - 2023/S 0F2-0013913;

Pacemakers with associated electrodes and additional accessories for pacemaker implantation and testing, publication number in EPPC - 2023/S 0F2-0013936

²Ministry of Health Implantation material and consumables for interventional neuroradiology for healthcare institutions in the Republic of Croatia, publication number in EPPC – 2023/S 0F2-0013906; UHC Zagreb Medications in the CHIF lists with generic parallels for healthcare institutions in the Republic of

Croatia I-XII, publication number in EPPC - 2023/S 0F2-0032745;

mentioned provision of the Act does not exclude the possibility of imposing additional control obligations upon the bidder.

Some contracting authorities nevertheless accepted the economic entity's request³ and removed the disputed provision from the joint procurement documentation, while some did not even include it in their documentation⁴.

Although it is logical to standardize the basic provisions of procurement documentation (which are not specific to the subject of procurement) in all joint public procurement procedures, this is not the case even when the same contracting authority carries out two or more joint procurement procedures. That is, the same central authority is further inconsistent in its actions as it imposes reporting obligations in some procedures but not in others⁵. Control over the performance of the contract/framework agreement is, in fact, the responsibility of the contracting authority. It is also a fact that most contracting authorities' IT systems do not allow for the control of contract performance, so they transfer their own legal obligation to economic entities.

The fulfillment of such a reporting obligation will impose additional adaptive technical and organizational requirements on economic entities associated with significant additional unplanned operating costs.

Although the specific report template (number of rows, number of columns) is not defined, it can be assumed that drafting such a report is a very complex, time-consuming, and technically demanding job that requires data control and processing in each contract, each item of the bill of quantities, and each item of which there are dozens/hundreds. The fulfillment of this provision requires a complex IT infrastructure and significant human resources.

Sestre milosrdnice UHC – Implantation material and consumables for endoscopy and endoscopic devices for healthcare institutions in the Republic of Croatia, publication number in EPPC – 2023/S 0F2-0027112 – reporting is required



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³ CHC Rijeka – Reagents, tests and other consumables for laboratory diagnostics for healthcare institutions in the Republic of Croatia, publication number in EPPC – 2023/S 0F2-0029799

⁴ E.g., Dubrava UH – Implantation material and consumables for neurosurgery for healthcare institutions in the Republic of Croatia, publication number in EPPC – 2023/S 0F2-0020639

 $^{^5}$ E.g., Sestre milosrdnice UHC – Ear implants (including artificial cochlea) for healthcare institutions in the Republic of Croatia 2023/S 0F2-0010636 – no reporting required

Recommendations

AmCham believes that the provisions obliging suppliers to submit reports are not in accordance with the provisions of the Public Procurement Act for the following reasons.

First of all, such provisions are contrary to the provisions of Article 313, paragraph 2 of the Act, which clearly prescribes control of the performance of contracts awarded in public procurement procedures as an obligation of public contracting authorities. Contracting authorities are not authorized to transfer such an obligation to bidders, especially not in the way that has been seen in recent public procurement practice. AmCham believes that the aforementioned clear provision of the Public Procurement Act cannot be derogated by the provision of the Regulation on the internal organization of the State Office for Central Public Procurement, a subordinate legal act.

Additionally, and as already stated, the bidders observed that reporting obligations are often not clearly stated in the PD as one of the conditions of participation but, for example, as a form attached to the PD. Such actions of the contracting authority could be considered contrary to the rules under Article 200 of the Public Procurement Act on the way of structuring procurement documentation, and at the same time, they ignore the fact that the inclusion of the reporting obligation in a particular procedure is de facto a condition for participation. That is, complying with the requirements of the reporting obligation in practice represents a significant financial, organizational, and time burden for the bidders, which they cannot adequately reflect in the pricing segment of their bids due to the aforementioned practice of PD structuring.

The inclusion of such obligations is particularly difficult for small and medium-sized enterprises. Due to the impossibility of complying with the reporting obligation and the financial and organizational burden it implies, such businesses could effectively be prevented from participating in (joint) public procurement procedures. This would violate the fundamental principles of public procurement prescribed in Article 4, paragraph 1 of the Public Procurement Act, especially the principle of equal treatment and the principle of prohibition of discrimination.

In view of all the above, AmCham proposes that provisions obliging suppliers to submit reports not be included in the procurement documentation in the future.

Annex 1:

| Contracting authority | Subject of procurement | EPPC publication number | Deadline for delivery | Clause on delivery of reports | Request to delete clause accepted |
|-----------------------|---|-------------------------------|--------------------------|---|--|
| Ministry of Health | Implantation material and consumables for interventional neuroradiology | 2023/S 0F2- 0013906 | 5/10/2023 | Yes | No |
| Ministry of Health | Implantation material and consumables for invasive and interventional cardiology | 2023/S 0F2- 0013913 | 5/10/2023 | Yes | / |
| Ministry of Health | Pacemakers with associated electrodes and additional accessories for pacemaker implantation and testing | 2023/S 0F2- 0013936 | 5/10/2023 | Yes | |
| Ministry of Health | Consumables for medically assisted fertilization, gynecology and human reproduction | 2023/S 0F2- 0035583 | 10/9/2023 | No | |
| Ministry of Health | Consumables for nuclear medicine | 2022/S 0F2- 0023686 | 11/22/2022 | No | |
| Ministry of Health | Laundry procurement, rental, washing and ironing | Not published | | | |
| UHC ZAGREB | Medications in the CHIF lists that have generic parallels | 2023/S 0F2- 0032745 | 9/18/2023- 9/25/2023 | Yes | No |
| UHC ZAGREB | Other medications that are not included in the categories under numbers 1 and 17. | 2023/S 0F2- 0031090 | 9/6/2023- 9/20/2023 | Yes | No |
| UHC ZAGREB | Implantation material and consumables for cardiac surgery and extracorporeal blood flow | 2023/S 0F2- 0013804 | 5/12/2023 | No | |
| UHC ZAGREB | Implantation material for vascular surgery | 2023/S 0F2- 0013630 | 5/10/2023 | No | |
| UHC ZAGREB | Implantation material and consumables for electrophysiology | 2023/S 0F2- 0013309 | 5/15/2023 | No | |
| UHC ZAGREB | Implantation material and consumables for electrophysiology – repeated | 2023/S 0F2- 0040893 | 11/6/2023 | No | |

| UHC ZAGREB | Ophthalmic implantation material and consumables | 2023/S 0F2- 0012569 | 5/3/2023 | No | |
|------------------------------|---|------------------------|-----------|-----|---|
| UHC ZAGREB | Ophthalmic implantation material and consumables – repeated | 2023/S 0F2- 0040887 | 11/6/2023 | No | |
| UHC ZAGREB | Implantation material and consumables for interventional radiology | 2023/S 0F2- 0013590 | 5/16/2023 | No | |
| UHC ZAGREB | Implantation material and consumables for interventional radiology – repeated procedure part I | 2023/S 0F2- 0040472 | 11/2/2023 | No | |
| UHC ZAGREB | Implantation material and consumables for interventional radiology – repeated procedure part II | 2023/S 0F2- 0040887 | 11/7/2023 | No | |
| UHC ZAGREB | Medical/infectious waste disposal services | Not published | | | |
| Sestre milosrdnice UHC | Implantation material and consumables for endoscopy and endoscopic devices | 2023/S 0F2- 0027112 | 8/8/2023 | Yes | / |
| Sestre milosrdnice UHC | Ear implants (including artificial cochleae) | 2023/S 0F2- 0010636 | 4/24/2023 | No | / |
| Sestre milosrdnice UHC | Gastroenterological implantation material and consumables | 2023/S 0F2- 0027404 | 8/17/2023 | Yes | / |
| Sestre milosrdnice UHC | Consumables for oncology and radiotherapy for healthcare institutions in the Republic of Croatia | 2023/S 0F2- 0024282 | 7/18/2023 | Yes | / |
| Sestre milosrdnice UHC | Implantation material for traumatology (fracture devices, screws, plates for fixation, systems for installing plates and screws and others) | 2023/S 0F2- 0026087 | 7/31/2023 | Yes | / |
| UHC Osijek | Catheters, drains, probes and cannulas | 2023/S 0F2- 0023003 | 7/6/2023 | Yes | / |
| UHC Osijek | Consumables for ENT | 2023/S 0F2- 0018054 | 6/5/2023 | Yes | / |
| UHC Osijek | Surgical suture material | 2023/S 0F2- 0018982 | 6/7/2023 | Yes | / |
| UH Split | Laparoscopic instruments and consumables for electrosurgery | 2023/S 0F2- 0012650 | 5/8/2023 | Yes | / |



| | Material for existing ECC | 2022/5 052 | 10/11/2022 | | |
|---|---|---------------------------|------------|--|----------|
| UH Split | Material for existing ECG monitors and defibrillators and other accessories for monitoring cardiac function | 2023/S 0F2- 0036786 | 10/11/2023 | Yes | |
| UH Split | Material for closing and suturing wounds (splitters, cutters – staplers) | 2023/S 0F2- 0012506 | 5/3/2023 | Yes | / |
| CHC Rijeka | Reagents, tests and other consumables for laboratory diagnostics | 2023/S 0F2- 0029799 | 9/15/2023 | Yes - in the consulta tion phase | Accepted |
| CHC Rijeka | Devices and instruments for infusion, biopsy, puncture and administration of cytostatics | 2023/S 0F2- 0013162 | 5/22/2023 | Yes | / |
| CHC Rijeka | Tests and consumables for pathology and cytology | 2023/S 0F2- 0042782 | 10/16/2023 | No | |
| CHC Rijeka | Surgical instruments | 2023/S 0F2- 0041906 | 11/6/2023 | No | |
| Dubrava UH | Implantation material for plastic surgery | 2023/S 0F2- 0018256 | 6/7/2023 | No | |
| Dubrava UH | Neurosurgical implantation material and consumables | 2023/S 0F2- 0020639 | 6/26/2023 | No | |
| Dubrava UH | Consumables, devices and instruments for anesthesiology and intensive care medicine | 2023/S 0F2- 0033380 | 9/22/2023 | No | |
| Dubrava UH | Consumables for dentistry | Not published | | | |
| Dubrava UH | Consumables for neurology | 2023/S 0F2- 0039611 | 10/27/2023 | No | |
| Merkur UH | Consumables for hemodialysis | 2023/S 0F2- 0040163 | 11/3/2023 | Yes | |
| Merkur UH | Medical consumables – plaster | 2023/S 0F2- 0015071 | 5/17/2023 | Yes | |
| Merkur UH | Medical gases | 2023/S 0F2- 0040226 | 10/27/2023 | No | |
| Fran Mihaljević UHID | Material for hygienic needs and care (including antiseptics and disinfectants) | 2023/S 0F2- 0027325 | 9/29/2023 | No | |
| Clinic for Orthopedic Surgery Lovran | Procurement and delivery of medications for a one-year period | 2023/S 0F2- 0020387 | 9/14/2023 | No | |
| Children's Hospital | PRODUCTS FOR PARENTERAL NUTRITION | 2023/S 0F2- 0035903 | 9/14/2023 | No | |



| Croatian Institute for Transfusion Medicine | Consumables for transfusiology | 2023/S 0F2- 0021366 | 5/24/2023 | No | |
|--|---|------------------------|-----------|-----|----|
| Croatian Institute of Public Health | Reagents, tests and other consumables for microbiology for healthcare institutions in the Republic of Croatia | 2023/S 0F2- 0034252 | 9/26/2023 | No | |
| Croatian Institute of Public Health | Cleaning and maintenance equipment and agents | 2023/S 0F2- 0023829 | 7/17/2023 | No | |
| Croatian Institute of Public Health | Rapid antigen tests | 2023/S 0F2- 0023028 | 7/4/2023 | No | |
| Croatian Health Insurance Fund | Procurement of special groups of medications | 2023/S 0F2- 0022606 | 7/5/2023 | Yes | No |

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