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**Specification for the purchase of laboratory plastics**

**( LP -2025 )**

1. **Service customer profile.**

ICF "Alliance for Public Health" (hereinafter referred to as the Alliance) is a leading professional organization that, in cooperation with key public organizations, the Ministry of Health and other government bodies, is fighting a number of epidemics, including HIV /AIDS and TB in Ukraine, manages prevention programs and provides high-quality technical support and financial resources to organizations on the ground. The mission of the Alliance is to reduce the spread of infections and mortality and reduce the negative impact of epidemics by supporting public resistance to them in Ukraine, as well as by disseminating effective approaches to prevention and treatment in Eastern Europe and Central Asia.

As an independent legal entity registered in Ukraine since 2003 and after gaining managerial autonomy since January 2009, the Alliance shares the values and remains a member of the global partnership of the Public Health Alliance (an international charitable organization uniting 30 organizations from different countries, with a Secretariat in Hove , United Kingdom of Great Britain and Northern Ireland).

The main programs currently implemented by the Alliance are funded by the Global Fund to Fight AIDS, Tuberculosis and Malaria.

This procurement is being carried out within the framework of the implementation of the program " Sustainable Response to HIV and TB Epidemics in the Context of War and Recovery in Ukraine " for 2024-2026 in accordance with Grant Agreement No. 3644 dated "19" December 2023 (grant name UKR - C - AUA ), between the Global Fund to Fight AIDS, Tuberculosis and Malaria and the ICF "Public Health Alliance". Payment is made without value added tax on the basis of paragraph 26 of subsection 2 of section XX of the Tax Code of Ukraine.

1. **General product description.**
   1. The Alliance reserves the right to increase or decrease the purchase volume by +/- 30%.
   2. Each participant has the right to submit a price offer for all or any individual lot listed below.
   3. A separate winner of the competition will be selected for each Lot.

|  |  |  |
| --- | --- | --- |
| Lot No. | Goods | Quantity, pcs. |
| 1 | Centrifuge tubes , Falcon type , sterile, with conical bottom and screw cap, 50 ml | 122050 |
| 2 | Cryotubes, sterile, with screw cap , with external thread, 2 ml | 254 00 |
| 3 | Universal tip with filter, volume 100 - 1000 μl | 64162 |

\*The participant of the competition has the right to offer an increased quantity of products if this is due to the peculiarities of the manufacturer's packaging.

|  |
| --- |
| **Lot #1** - Centrifuge tubes , Falcon type , sterile, with conical bottom and screw cap, 50 ml |
| 1. General description of the product and its technical characteristics   1.1 Product description:  - must be sterile, graduated, polypropylene,  - each falcon in individual packaging,  - centrifugation at not less than 3000 rpm,  - there must be a field for recording,  - have a screw cap,  bottom type : conical,  - without a " skirt ",  - volume: 50 ml.  Medical products are intended for multifunctional laboratory diagnostics In Vitro .    2. Quality requirements2.1 The product must be approved for marketing and/or operation (application) in accordance with the legislation. Medical devices must comply with the requirements of the Technical Regulations.2.2 The product must be approved for marketing and/or (application) in accordance with the legislation. Medical devices must comply with the requirements of the Technical Regulations\*. The supplier must provide a certified copy of the Declaration of Conformity of Medical Devices to the Technical Regulations\* or the Certificate of State Registration, other relevant documents, a copy of the quality certificate. \* - Resolution of the Cabinet of Ministers of Ukraine dated 02.10.2013 No. 754 “On Approval of the Technical Regulations on Medical Devices for Diagnostics in vitro ".  \*\*- other permits, copies of documents of conformity confirming the completion of the conformity assessment procedure and marking with the national mark of conformity in accordance with the requirements of the Technical Regulations.   3. Primary packaging3.1 Packaging of goods must ensure the preservation of the quality and integrity of the goods; comply with established international standards and ensure its preservation during transportation, loading, unloading and storage.3.2 Packaging and labeling must comply with the requirements of current legislation. The packaging of the goods must contain the following information: manufacturer's name, product name, manufacturer's name, expiration dates, batch number, number of tips contained in the rack, storage requirements, registration certificate number of the Ministry of Health.3.3 Availability of instructions for the use of the medical product/technical description of the medical product in Ukrainian.    4. Expected delivery date of goods: by 30.11.202 5   5. Shelf lifeAt the time of delivery of products to the logistics warehouse, the residual shelf life must be no less than 70% of the total. In the event of a product with a shorter remaining shelf life, additional approval must be obtained from the customer.   6. General requirements for products and documentation 6.1 Compliance with the requirements of the manufacturer's ISO 13485 and/or ISO 9001 quality management system, confirmed by providing copies of certificates 6.2 Documents confirming the completion of the conformity assessment procedure in accordance with the requirements of the Technical Regulations 6.3 Compliance with the requirements of European standards (if any) 6.4 Quality certificate 6.5 Other permits for the offered product |
| **Lot #2** - Sterile cryotubes with screw cap , with external thread, 2 ml |
| 1. General product description and technical specifications   1.1 Product description:  Volume : 2 ml  • Sterile  • Free from human DNA , free from DNase human , free from PCR inhibitors and pyrogens , free from RNase human  • Polypropylene​  • Graduated  • Transparent  • Color Lids : Neutral  Cap type : Screw , with external thread  • Availability of a field for recording :  • Flat-bottomed  Application temperature : from at least -70 °C to + 121 °C or more  Packaging type : Packaging  • Quantity units per package : no more than 100  Medical products are designed for multifunctional laboratory diagnostics In Vitro .  2. Quality requirements2.1 The product must be approved for placing on the market and/or operation (application) in accordance with the legislation. Medical devices must comply with the requirements of the Technical Regulations.2.2 The product must be approved for placing on the market and/or (application) in accordance with the legislation. Medical devices must comply with the requirements of the Technical Regulations\*. The supplier must provide a certified copy of the Declaration of Conformity of Medical Devices to the Technical Regulations\* or the Certificate of State Registration, other relevant documents, a copy of the quality certificate. \* - Resolution of the Cabinet of Ministers of Ukraine dated 02.10.2013 No. 754 “On Approval of the Technical Regulations on Medical Devices for Diagnostics in vitro ".  \*\*- other permits, copies of documents of conformity confirming the completion of the conformity assessment procedure and marking with the national mark of conformity in accordance with the requirements of the Technical Regulations.   3. Primary packaging3.1 Packaging of goods must ensure the preservation of the quality and integrity of the goods; comply with established international standards and ensure its preservation during transportation, loading, unloading and storage.3.2 Packaging and labeling must comply with the requirements of current legislation. The packaging of the goods must contain the following information: manufacturer's name, product name, manufacturer's name, expiration dates, batch number, storage requirements, registration certificate number of the Ministry of Health.3.3 Availability of instructions for the use of the medical product/technical description of the medical product in Ukrainian.   4. Expected delivery date of goods: by 30.11.202 5   5. Shelf lifeAt the time of delivery of products to the logistics warehouse, the residual shelf life must be no less than 70% of the total. In the event of a product with a shorter remaining shelf life, additional approval must be obtained from the customer.   6. General requirements for products and documentation 6.1 Compliance with the requirements of the manufacturer's ISO 13485 and/or ISO 9001 quality management system, confirmed by providing copies of certificates 6.2 Documents confirming the completion of the conformity assessment procedure in accordance with the requirements of the Technical Regulations 6.3 Compliance with the requirements of European standards (if any) 6.4 Quality certificate 6.5 Copy of the metrological verification certificate for the equipment (or confirmation that such a certificate will be issued and provided together with the delivery of the product) 6.6 Other permits for the offered product |
| **Lot No. 3** - Universal tip with filter, volume 100 - 1000 μl |
| 1. General description of the product and its technical characteristics   1.1 Product description:• Dosage range 100-1000 μl  • Sterility: Sterile, apyrogenic , free of RNA and DNAase  • Presence of a filter: Yes• Material: polypropylene• Packaging type: Packaging• Number of units: no more than 100 pcs. / pack . in a stand  Medical products are intended for multifunctional laboratory diagnostics In Vitro .  Medical products must be universal - must be suitable for working with pipette dispensers single-channel from different manufacturers   2. Quality requirements2.1 The product must be approved for placing on the market and/or operation (application) in accordance with the legislation. Medical devices must comply with the requirements of the Technical Regulations.2.2 The product must be approved for placing on the market and/or (application) in accordance with the legislation. Medical devices must comply with the requirements of the Technical Regulations\*. The supplier must provide a certified copy of the Declaration of Conformity of Medical Devices to the Technical Regulations\* or the Certificate of State Registration, other relevant documents, a copy of the quality certificate. \* - Resolution of the Cabinet of Ministers of Ukraine dated 02.10.2013 No. 754 “On Approval of the Technical Regulations on Medical Devices for Diagnostics in vitro ".  \*\*- other permits, copies of documents of conformity confirming the completion of the conformity assessment procedure and marking with the national mark of conformity in accordance with the requirements of the Technical Regulations.   3. Primary packaging3.1 Packaging of goods must ensure the preservation of the quality and integrity of the goods; comply with established international standards and ensure its preservation during transportation, loading, unloading and storage.3.2 Packaging and labeling must comply with the requirements of current legislation. The packaging of the goods must contain the following information: manufacturer's name, product name, manufacturer's name, expiration dates, batch number, number of tips contained in the rack, storage requirements, registration certificate number of the Ministry of Health.3.3 Availability of instructions for the use of the medical product/technical description of the medical product in Ukrainian.3.4 Packaging of the Goods: 1 rack no more than 100 tips.   4. Expected delivery date of goods: by 30.11.202 5   5. Shelf lifeAt the time of delivery of products to the logistics warehouse, the residual shelf life must be no less than 70% of the total. In the event of the presence of goods with a shorter residual shelf life, there must be additional approval with the customer.   6. General requirements for products and documentation6.1 Compliance with the requirements of the manufacturer's ISO 13485 and/or ISO 9001 quality management system, confirmed by providing copies of certificates 6.2 Documents confirming the completion of the conformity assessment procedure in accordance with the requirements of the Technical Regulations6.3 Compliance with the requirements of European standards (if any)6.4 Quality certificate6.5 Other permits for the offered goods |
|  |

*3. Delivery terms.*

***3.1. Delivery on DAP terms to the address of the Alliance warehouse, Kyiv/Kyiv region.***

3 .2. It is desirable that the delivery was carried out in one batch no later than November 30, 2025. In case of delivery in several batches, you must indicate your proposed delivery schedule in Appendix No. 3 to the specification, column "Delivery time, batches".

3.3. Participants are invited to provide their own forecasts regarding the delivery times of partial and full orders, as well as deliveries in multiple batches (Appendix No. 3)

*4. Payment terms*

4.1. Payment terms: 50% - advance payment within 10 (ten) business days from the date of signing the Agreement, 50% - balance payment within 10 (ten) business days from the date of completion of acceptance of the Goods.

4.2. The supply contract will be concluded and payments will be made in:

* in Ukrainian hryvnias for residents of Ukraine, which will be the equivalent in dollars according to the official exchange rate of the National Bank of Ukraine on the day of invoicing for the supply of each individual batch of Goods.

**The supply of this product is exempt from VAT!**

**Attention potential resident suppliers!** In accordance with the provisions of Clause 26, Subsection 2, Section XX of the Tax Code of Ukraine: transactions for the supply of goods (except for excisable goods) and the provision of services to the customs territory of Ukraine are exempt from value added tax, if such goods/services are paid for at the expense of grants ( sub-grants ) provided in accordance with the programs of the Global Fund to Fight AIDS, Tuberculosis and Malaria in Ukraine, which are implemented in accordance with the Law of Ukraine "On the Implementation of the Programs of the Global Fund to Fight AIDS, Tuberculosis and Malaria in Ukraine" (No. 4999-17 dated 11.08.2013). The procedure for carrying out such operations is determined by the Resolution of the Cabinet of Ministers of Ukraine “Some issues of importing goods into the customs territory of Ukraine and supplying goods and services in the customs territory of Ukraine paid for by grants ( sub-grants ) of the Global Fund to Fight AIDS, Tuberculosis and Malaria in Ukraine” dated April 17, 2013 No. 284.

When forming your price offer, please pay attention to the content of clause 26, subsection 2, section XX "Transitional provisions" of the Tax Code of Ukraine.

Thus, in accordance with the norms of the Tax Code of Ukraine for transactions of supply of goods / provision of services carried out to implement the programs of the Global Fund to Fight AIDS, Tuberculosis and Malaria in Ukraine (and which are exempt from VAT), the supplier is granted a tax credit:

"In the event of transactions exempted under this paragraph, the provisions of paragraph 198.5 of Article 198 of this Code and the provisions of Article 199 of this Code shall not apply."

**5. Composition of the participant's competitive proposal.**

Each participant in the competitive process must provide the following documentation and materials:

a) a copy of documents confirming the state registration of the competition participant.

b) copies of valid documents allowing the use of this product on the territory of Ukraine (items 2 and 6 in the product description)

c) technical description/instructions/technical data sheet for the proposed products (clause 3.3 in the product description).

d) completed and signed annexes to the specification:

- completed competition participant form (see Appendix No. 1) ;

- technical specification (see Appendix No. 2 – additionally in Word format );

- table of the applicant's price offer (see Appendix No. 3 - additionally in Word format );

- Composition of final beneficiary owners tenderer ( see Appendix No. 4).

e ) any other documents that, in your opinion, may be useful in making a decision.

**6. Criteria for evaluating price offers:**

a) compliance of the participant's application with the technical specification;

b) compliance of the participant's application with organizational requirements;

c) proper product quality, documented;

d) reasonable price;

e) delivery times

**Appendix No. 1**

**Specification for the purchase of laboratory plastics ( LP - 2025)**

**General information**

Please fill in the table below.

|  |  |  |
| --- | --- | --- |
| 1. | Full company name |  |
| 2. | Company legal address |  |
| 3. | Physical address of the company |  |
| 4. | Head of the company: position, name |  |
| 5. | Contact phone number of the company head |  |
| 6. | Contact person for this tender proposal |  |
| 7. | Contact person phone number |  |
| 8. | Contact person's fax number |  |
| 9. | Contact person email address |  |
| 10. | Company page on the Internet |  |
| 11. | Bank details for concluding a supply contract |  |

Signed by me, \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_,

holding the position\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_(head of the enterprise)

on behalf of the company \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

\_\_\_\_\_\_\_ (day) \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_ (month) 20\_\_\_\_\_\_\_\_ (year).

\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_ (signature)

**Appendix No. 2**

**Specification for the purchase of laboratory plastics ( LP - 2025)**

**Technical specification**

|  |  |
| --- | --- |
| **Centrifuge tubes , Falcon type , sterile, with conical bottom and screw cap, 50 ml** | **Specify the name and TM of the product:** |
| .1 Product Description:  - must be sterile, graduated, polypropylene,  - each falcon in individual packaging,  - centrifugation at not less than 3000 rpm,  - there must be a field for recording,  - have a screw cap,  bottom type : conical,  - without a " skirt ",  - volume: 50 ml.  Medical products are designed for multifunctional laboratory diagnostics In Vitro . | **Write down the proposed characteristics:** |
| **Documents provided** | |
| The supplier must provide a certified copy of the Declaration of Conformity of Medical Devices with the Technical Regulations\* or the Certificate of State Registration, other relevant documents, a copy of the quality certificate.  \* - Resolution of the Cabinet of Ministers of Ukraine dated 02.10.2013 No. 754 “On Approval of the Technical Regulations on Medical Devices for Diagnostics in vitro ".  \*\*- other permits, copies of conformity documents confirming the completion of the conformity assessment procedure and marking with the national conformity mark in accordance with the requirements of the Technical Regulations. | **Please write down the documents provided:** |
| Availability of instructions for use of the medical device/technical description of the medical device in Ukrainian. | **yes / no:** |
| Shelf life  At the time of delivery of products to the logistics warehouse, the remaining shelf life must be no less than 70% of the total. In the event of the presence of goods with a shorter remaining shelf life, there must be additional agreement with the customer. | **indicate the total and remaining shelf life:** |
| General requirements for products and documentation  6.1 Compliance with the requirements of the manufacturer's ISO 13485 and/or ISO 9001 quality management system, confirmed by providing copies of certificates 6.2 Documents confirming completion of the conformity assessment procedure in accordance with the requirements of the Technical Regulations 6.3 Compliance with the requirements of European standards (if any) 6.4 Quality certificate 6.5 Other permits for the offered product | **Please write down the documents provided:** |

|  |  |
| --- | --- |
| **Lot #2 - Sterile cryotubes with screw cap , with external thread, 2 ml** | **Specify the name and TM of the product:** |
| .1 Product Description:  Volume: 2 ml  • Sterile  • Free from human DNA, free from human DNases , free from PCR inhibitors and pyrogens, free from human RNases  • Polypropylene​  • Graduated  • Transparent  • Color Lids : Neutral  Cap type : Screw , with external thread  • Availability of a field for recording :  • Flat-bottomed  Application temperature : from at least -70 °C to + 121 °C or more  Packaging type : Packaging  • Quantity units per package : no more than 100  Medical products are designed for multifunctional laboratory diagnostics In Vitro . | **Write down the proposed characteristics:** |
| **Documents provided** | |
| The supplier must provide a certified copy of the Declaration of Conformity of Medical Devices with the Technical Regulations\* or the Certificate of State Registration, other relevant documents, a copy of the quality certificate.  \* - Resolution of the Cabinet of Ministers of Ukraine dated 02.10.2013 No. 754 “On Approval of the Technical Regulations on Medical Devices for Diagnostics in vitro ".  \*\*- other permits, copies of conformity documents confirming the completion of the conformity assessment procedure and marking with the national conformity mark in accordance with the requirements of the Technical Regulations. | **Please write down the documents provided:** |
| Availability of instructions for use of the medical device/technical description of the medical device in Ukrainian. | **yes / no:** |
| At the time of delivery of products to the logistics warehouse, the remaining shelf life must be no less than 70% of the total. In the event of the presence of goods with a shorter remaining shelf life, there must be additional agreement with the customer. | **indicate the total and remaining shelf life:** |
| General requirements for products and documentation  6.1 Compliance with the requirements of the manufacturer's ISO 13485 and/or ISO 9001 quality management system, confirmed by providing copies of certificates 6.2 Documents confirming completion of the conformity assessment procedure in accordance with the requirements of the Technical Regulations 6.3 Compliance with the requirements of European standards (if any) 6.4 Quality certificate 6.5 Copy of the metrological verification certificate for the equipment (or confirmation that such a certificate will be issued and provided together with the delivery of the goods) 6.6 Other permits for the offered goods | **Please write down the documents provided:** |

|  |  |
| --- | --- |
| **Lot #3 - Universal tip with filter, volume 100 - 1000 μl** | **Specify the name and TM of the product:** |
| Product Description:  • Dosing range 100-1000 μl  • Sterility: Sterile, pyrogen-free , free of RNA and DNAase  • Filter available: Yes • Material: polypropylene • Packaging type: Packaging • Number of units: no more than 100 pcs. / pack . in a stand  Medical products are intended for multifunctional laboratory diagnostics In Vitro .  Medical products must be universal - must be suitable for working with pipette dispensers single-channel from different manufacturers | **Write down the proposed characteristics:** |
| **Documents provided** |  |
| The supplier must provide a certified copy of the Declaration of Conformity of Medical Devices with the Technical Regulations\* or the Certificate of State Registration, other relevant documents, a copy of the quality certificate.  \* - Resolution of the Cabinet of Ministers of Ukraine dated 02.10.2013 No. 754 “On Approval of the Technical Regulations on Medical Devices for Diagnostics in vitro ".  \*\*- other permits, copies of conformity documents confirming the completion of the conformity assessment procedure and marking with the national conformity mark in accordance with the requirements of the Technical Regulations. | **Please write down the documents provided:** |
| Availability of instructions for use of the medical device/technical description of the medical device in Ukrainian. | **yes / no:** |
| Shelf life  At the time of delivery of products to the logistics warehouse, the remaining shelf life must be no less than 70% of the total. In the event of the presence of goods with a shorter remaining shelf life, there must be additional agreement with the customer. | **indicate the total and remaining shelf life:** |
| General requirements for products and documentation  6.1 Compliance with the requirements of the manufacturer's ISO 13485 and/or ISO 9001 quality management system, confirmed by providing copies of certificates 6.2 Documents confirming completion of the conformity assessment procedure in accordance with the requirements of the Technical Regulations 6.3 Compliance with the requirements of European standards (if any) 6.4 Quality certificate 6.5 Other permits for the offered product | **Please write down the documents provided:** |

Signed by me, \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_, holding the position of\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_(head of the enterprise)

on behalf of the company \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

\_\_\_\_\_\_\_ (date) \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_ (month) 20\_\_\_\_\_\_\_\_ (year). \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_ (signature)

**Appendix No. 3**

**Specification for the purchase of laboratory plastics ( LP - 2024)**

**Price offer**

Each participant is invited to form their price offers in the form of the table below.

When filling out the table, please note the following:

1. The price for the products is provided on the terms of delivery in accordance with the requirements of clause 4.1 of the specification.
2. The price of the Goods must include the cost of the product itself, packaging/container, labeling and delivery.
3. Price is provided:

* in US dollars;
* taking into account all applicable taxes, fees and costs in accordance with the specified delivery conditions in accordance with the legislation of Ukraine;
* **excluding VAT** .

In accordance with the requirements of the Resolution of the Cabinet of Ministers of Ukraine dated April 17, 2013 No. 284 "Some issues of importation of goods into the customs territory of Ukraine and supply of goods and services in the customs territory of Ukraine paid for by grants ( sub-grants ) of the Global Fund to Fight AIDS, Tuberculosis and Malaria in Ukraine".

1. Payments will be made in Ukrainian hryvnias according to the official exchange rate of the National Bank of Ukraine on the date of invoice.

|  |  |  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- | --- | --- |
| No. | Goods | Product brand, manufacturer, model | Country of origin | Number of units ( pcs .) | Price per 1 unit, USD excluding VAT | Total, US dollars excluding VAT | Delivery time, batches |
| 1 |  |  |  |  |  |  |  |
| 2 |  |  |  |  |  |  |  |
| 3 |  |  |  |  |  |  |  |

Signed by me, \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_,

holding the position\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_(head of the enterprise)

on behalf of the company \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

\_\_\_\_\_\_\_ (day) \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_ (month) 20\_\_\_\_\_\_\_\_ (year).

\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_ (signature)

**Appendix No. 4**

**Specification for the purchase of laboratory plastics ( LP - 2024)**

Based on Article 650-1 of the Civil Code of Ukraine, by signing this form, we confirm the following assurances that are relevant for concluding an agreement with the ICF "Alliance of Public Health":

1. The Company (the legal entity submitting this and other documents for participation in the tender, specified below), any of its directors (members of the board of directors), participants of the Company, its ultimate beneficial owner/owners, its officers or employees of the Company, or any agent, affiliate or other person acting on behalf of the Company, is not currently subject to any US sanctions administered by the Office of Foreign Assets Control of the US Treasury or the US Department of State, the United Nations Security Council, the European Union, Her Majesty's Treasury of the United Kingdom or other authorized sanctioning authority.
2. The Company, and/or a participant of the Company, and/or the ultimate beneficial owner of the Company are not included in the sanctions list of the National Security and Defense Council of Ukraine (in accordance with Article 5 of the Law of Ukraine “On Sanctions”).
3. No personal special economic and other restrictive measures (sanctions) have been applied to the Company's goods, services and/or works in accordance with Article 5 of the Law of Ukraine "On Sanctions".

# Composition of the ultimate beneficial owners of the tenderer

|  |  |  |  |  |
| --- | --- | --- | --- | --- |
| **Name of organization/Full name of individual** | **Registration code / passport data** | **Registration address** | **Citizenship** | **Is the organization/person listed on the sanctions lists of the US, the European Union, or Ukraine?** |
|  |  |  |  |  |
|  |  |  |  |  |
|  |  |  |  |  |

# *[signature] [position]*

Authorized to sign the commercial offer for and on behalf of:

*[ name companies ]*