



Abbott

Abbott GmbH
Max-Planck-Ring 2
65205 Wiesbaden
Postfach 1303, 65011 Wiesbaden

Tel. +49 61 22 580
Fax. +49 61 22 58 1244

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To: Centrul Pentru Achizitii Publice Centralizate In Sanatate

Tender no. ocds-b3wdp1-MD-1695305626407

“Achiziția Testelor pentru tehnologia de examinare de laborator a donatorilor de sânge/componente sanguine și sângelui/componentelor sanguine donat la prima etapă de triere și la a doua etapă de triere, întru realizarea Programului Național „Securitatea transfuzională și autoasigurarea țării cu produse sanguine” conform necesităților pentru anul 2024”.

LETTER OF CONFIRMATION

Herewith, we ABBOTT GMBH having its principal of business at Transfusion Medicine in regard to the Tender “Achiziția Testelor pentru tehnologia de examinare de laborator a donatorilor de sânge/componente sanguine și sângelui/componentelor sanguine donat la prima etapă de triere și la a doua etapă de triere, întru realizarea Programului Național „Securitatea transfuzională și autoasigurarea țării cu produse sanguine” conform necesităților pentru anul 2024”, confirm the following in regard to the Lot 1: “Teste pentru tehnologia de examinare de laborator a donatorilor de sânge/componente sanguine și sângelui/componentelor sanguine donat la prima etapă de triere”:

I. GENERAL REQUIREMENTS; number of tests, accessories / consumables / reagents / calibrators / solutions and needed consumables related to the activity of the laboratory medical device (by technology applied to existing device /by technology applied to another device than the existing one at the customer site).

DELIVERY TRanches FOR 2024 YEAR:

- I. tranche: February.
- II. tranche: April.
- III. tranche: July.
- IV. tranche: October.

1.1 REQUIREMENTS FOR THE REAGENTS:

- a) Stability when placed in a medical device: stability when placing in the medical device at least for 28 days.
- b) Provided by the manufacturer with protection elements against evaporation and leakage.

1.2 REQUIREMENTS FOR THE CONSUMABLES:

- a) Unusable.
- b) Ensures a high degree of accuracy and excludes contamination of samples.
- c) Delivered in secured package, marked and labelled by the manufacturer with the mention of identity data (name, batch / serial number, terms of validity, storage conditions). The identity data displayed on the box must coincide with those on the labels of each component of the set (as appropriate).

1.3 TEST FOR THE DETERMINATION OF HBsAg – 82100 TESTS

Destination: for the examination of donor blood in markers hemotransmissible infections - viral hepatitis B.
Method of application in the test reaction:

- a) Closed type technology;
- b) automated;

Sitz der Gesellschaft: Wiesbaden
Amtsgericht Wiesbaden HRB 31478

Geschäftsführer:
Christian Grapow
Robert Funck, Konstantinos Varlas



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c) chemiluminescent, including with various antigen identification options

Duration of the testing process, including the incubation period - till 30 minutes.

Diagnostic product:

1) Test for the screening of blood donated and intended for transfusion or as a raw material for the production of the human plasma preparation, of a generation that will ensure the detection of:

- a) In human serum / plasma,
- b) qualitative HBsAg,
- c) common characteristics for the genetic variations of HBV virus worldwide known, including for the Eastern European region.

2) Architect HBsAg test has 100% sensitivity on samples of blood donors;

3) Architect HBsAg test specificity is 99,91% inclusive, on samples a blood donor (is more than tender spec minimum requirement 99.88%);

4) Architect HBsAg test with analytical sensitivity (detection limit) is between 0.017 and 0.022 IU/ml. (It's much more sensitive than tender spec. 0.13 IU / ml.).

Components accompanying the diagnostic product: mandatory present all the necessary components, in sufficient quantities for the application of the test in the test reaction, according to the instructions for use of the product.

Form of packing: set delivered in secure packaging, marked and labeled by the manufacturer indicating identification data (name, batch number / series, expiry terms, storage conditions). The identity data displayed on the box must coincide with those on the labels of each component of the set.

1.4 TEST FOR THE DETERMINATION OF ANTI HCV – 82275 TESTS:

Destination: for the examination of donor blood in markers hemotransmissible infections - viral hepatitis C.

Method of application in the test reaction:

- a) In closed type technology;
- b) automated;
- c) chemiluminescent test, including with various antibody identification options.

Duration of the testing process, including the incubation period - till 30 minutes.

Diagnostic product:

1) Test for screening blood donated and intended for transfusion or as a raw material for the production of preparations from human plasma generation that will ensure the detection of:

- a) in human serum / plasma,
- b) quality of antibodies to viral hepatitis C virus,

2) Architect Anti HCV test has 100% sensitivity on samples of blood donors;

3) Architect Anti HCV test specificity is 99,93% inclusive, on samples a blood donor (is more than tender spec minimum requirement 99.84%).



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4) Assay will detect the combination of circulating antibodies to Core viral antigens Core, NS3, NS4 proteins, including at least 3 subtypes characteristic of the European region.

Components delivered together with the diagnostic product: mandatory present all the necessary components, in sufficient quantities for the application of the test in the test reaction, according to the instructions for use of the product.

Form of packing: set delivered in secure packaging, marked and labeled by the manufacturer indicating identification data (name, batch number / series, expiry terms, storage conditions). The identity data displayed on the box must coincide with those on the labels of each component of the set.

1.5 TEST FOR ANTI-TREPONEMA PALLIDUM ANTIBODIES – 82275 TESTS

Destination: for the examination of donor blood in markers hemotransmissible infections - anti-Treponema Pallidum antibodies.

Method of application in the test reaction:

- a) Closed type technology;
- b) automated;
- c) chemiluminescent test, including with various antibody identification options.

Duration of the testing process, including the incubation period - till 30 minutes.

Diagnostic product:

- 1) Test is designed for the qualitative detection of antibodies to Treponema Pallidum in human plasma, used in the screening of donated blood and intended for transfusion or raw material for the production of human plasma preparations;
- 2) Architect Syphilis TP test has 100% sensitivity on samples of blood donors;
- 3) Architect Syphilis TP test with specificity 99,94% inclusive, on samples of blood donors (is more than tender spec minimum requirement 99.88%).

Components delivered together with the diagnostic product: mandatory present all the necessary components, in sufficient quantities for the application of the test in the test reaction, according to the instructions for use of the product.

Form of packing: set delivered in secure packaging, marked and labeled by the manufacturer indicating identification data (name, batch number / series, expiry terms, storage conditions). The identity data displayed on the box must coincide with those on the labels of each component of the set.

1.6 TEST FOR DETERMINATION OF HIV AB/AG COMBO – 81575 TESTS

Destination: for the examination of donor blood in markers of hemotransmissible infections - Ag-HIV-1 P24 antigen and anti-HIV1 / HIV2 antibodies.

Method of application in the test reaction:

- a) Closed type technology;
- b) automated;
- c) chemiluminescent test, including with various identification options of antibody.



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Diagnostic product:

1) Test for screening blood donated and intended for transfusion or as a raw material for the production of human plasma preparations, of a generation that will ensure the detection of:

- a) in human serum/plasma,
- b) simultaneous qualitative of HIV-1 p24 Ag and anti-HIV1 / HIV2 antibodies,
- 2) Architect HIV Ag/Ab Combo test has 100% sensitivity on samples of blood donors;
- 3) Architect HIV Ag/Ab Combo test with specificity 99.89% inclusive, on samples of blood donors (is more than tender spec minimum requirement 99.88%).
- 4) Architect HIV Ag/Ab Combo test 0,87 IU/ml analytical sensitivity test (detection limit) inclusive, for P24 antigen. (It's much more sensitive than tender spec. 2 IU/ml).

Components delivered together with the diagnostic product: mandatory present all the necessary components, in sufficient quantities for the application of the test in the test reaction, according to the instructions for use of the product.

Form of packing: set delivered in secure packaging, marked and labeled by the manufacturer indicating identification data (name, batch number / series, expiry terms, storage conditions). The identity data displayed on the box must coincide with those on the labels of each component of the set.

II. THE OFFER INCLUDE THE FOLLOWING TECHNOLOGIES:

2.1 The technology applied to other device than the one existing in the institution, with all components: the laboratory medical device, number of tests, accessories / consumables / reagents / calibrators / solutions and other mandatory items, related to the activity of the medical laboratory device, required in the laboratory examination process of the donated blood, corresponding to:

- 1) HBsAg – 82100 tests;
- 2) Anti HCV antibodies – 82275 tests;
- 3) Antibodies against Treponema Pallidum –82275 tests;
- 4) Ag-HIV-1 P24 and anti-HIV1 / HIV2 antibodies – 81575 tests.

The offer includes devices ARCHITECT i2000SR, Abbott (2 units), with the year of production 2021 year (SN: ISR64824; ISR 64825) - with all components (number of tests, accessories / consumables / reagents / calibrators / solutions and other mandatory products, including device activity) required in the process laboratory examination of donated blood.

1. Medical device for examination of donor blood in markers of hemotransmissible infections:

- a) Closed type technology analyzer;
- b) automated technology analyzer;
- c) analyzer with chemiluminescent marker identification technique, including various antibody identification options;
- d) analyzer with the technique of concomitant identification of HBsAg markers, Anti-HCV antibodies, Anti-Treponema Pallidum antibodies, HIV Ag / Ab and optionally of other markers, such as: Anti HbC, HTLV, Cov-2IgG.

2. Accessories / consumables / reagents / calibrators / solutions and other mandatory products required in the testing process - provided for the total number of tests required: 328225 tests.

3. The capacity to load at the ARCHITECT i2000SR the samples for testing:

- a) 135 samples;
- b) the possibility of continuous loading starting from 1 (one) sample;



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4. Test processing speed – 200 tests per hour.

5. ARCHITECT i2000SR system functionality:

- a) automatic pipetting for reagents / calibrators / solutions applied in the offered technology, as well as samples, including sustained software management (permanent inventory of reagents / calibrators / solutions);
- b) automatic loading / removal for racks or other similar components obligatorily required in the testing process;
- c) automatic dilutions, including pre- and post-dilutions;
- d) automatic waste disposal, including sustained software management (permanent inventory of waste (residues));
- e) permanent monitoring of the status of the samples in the work list;
- f) built-in quality control (QC) system that will ensure the analysis of the quality of each sample, reagents / calibrators / solutions applied in the offered technology, qualification of the testing stages, with auditory and visual notification of errors during operation;
- g) provided with the specially designed module for emergency tests;
- h) provided with database management system;
- i) provided with built-in laser barcode readers;
- j) provided with mobile barcode laser readers;
- k) interconnected with the laboratory data management program in SIA Blood Service;

6. Requirements for the software database management system and its functionality:

- a) Available software applications with the latest generation accessories, including the Microsoft Windows Operating System;
- b) Receiving / monitoring / printing in electronic form or manually entering the results of the analyzes, the results of the quality control, the messages on the devices and events for the maintenance of the devices;
- c) On-line storage of analysis results, quality control results, device messages and device maintenance events, their record and reporting;
- d) User and certificate management;
- e) Submission of the results of the analysis / quality control in the computer system, including the notification of the errors appeared during the operation;
- f) Configuration of the operation of the connected devices.
- g) Viewing real-time statistics (production statistics) / statistics generated by the database management system.
- h) Accessories related to the device but mandatory for its operating conditions, if necessary:
 - 1) RS-232C port;
 - 2) USB 2.0 port;
 - 3) USB 3.0 port;
 - 4) network port (RJ-45).
- i) UPS with the capacity to provide the power supply for at least 60 minutes;
- j) water preparation system, including tank with minimum capacity for 300 samples in testing;

Documents/certificates of RGT submitted with the offer:

- 1. EC Certificate / Declaration of Conformity Certificate for the reagents, consumables, solutions.
- 2. Manufacturer Letter confirming each technical requirement of the eligibility criteria for the offered product, issued by the original manufacturer of the product, signed and stamped.
- 3. Presence of the instruction to use of the product from the manufacturer, in Romanian language, signed by the economic operator.



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4. Confirmation to present the quality certificate for each lot, in each tranche delivery.

5. Confirmation that upon delivery the shelf life of the test will be not less than 80% from its total shelf life.

Declaration of Conformity certificate for offered devices is included.

-Confirmation letter of each technical requirement from the eligibility criteria for the 2 (two) devices offered for performing laboratory examinations of donated blood.

- We confirm to provide with 2 (two) identical devices as model and performance (SN: ISR64824; ISR 64825).

- We confirm to present during the activity of the devices, of the copies of the manual service, electronic diagram, technical passports, including description of technical parts and components a device.

- Training of the personnel involved in the operation of the free medical device, that will be provided by our local distributor, within 10 days of the device being put into operation.

- Confirmation to deliver free of charge consumables, calibrators, working solutions, controls.

- Confirmation to provide free of charge assurance of the functionality, maintenance, repair of medical devices 2 (two) units, for the entire period of the number of tests contracted.

- Confirmation to support bidder to provide free of charge interconnection with the laboratory data management program in SIA Blood service during 10 days from the moment of commissioning and written notification of the beneficiary.



Kind regards,
Murat MERDAL
Division Vice President
Europe, Middle East & Africa
Abbott Transfusion Medicine

 **Abbott**
Abbott GmbH
Max-Planck-Ring 2
65205 Wiesbaden-Delkenheim