

Survey for Coagulation (COA12, COA4, COA2)

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Instructions for Use

Notes:

The usual precautions in the laboratory for potentially hazardous samples apply for these samples. The blood donations used for production were found to be non-reactive for HBsAg, anti-HIV 1/2 and anti-HCV.

Various tasks of the proficiency testing scheme may occasionally be given over to qualified subcontractors. However, ESfEQA is responsible to the participants for the subcontractor's work.

Results from sample analysis may only be disclosed to colleagues from other laboratories after the testing period has been concluded.

By registering for and participating in this EQA participants agree to accept the general terms and conditions of ESfEQA GmbH. These can be retrieved online at www.esfeqa.eu.

1. Intended Use

The samples are intended for use as quantitative control material for External Quality Assessment (EQA) in medical laboratories for the following analytes:

Antithrombin III (as AT III activity and concentration), aPTT, Fibrinogen, D-Dimer, Protein C, Protein S, Prothrombin Time (as Quick and INR values as well as in seconds), Thrombin Time

2. Product Description

The samples are lyophilized material prepared from human blood plasma.

3. Storage and Stability

The samples should be stored upright at 2-8 °C. They are stable at least until the deadline for data submission as indicated below.

After reconstitution, samples are stable in tightly recapped vials for 24 hours at 2-8 °C or 15-25 °C.

4. Sample Preparation and Analysis

Remove the screw cap from the vial and pipette exactly 1 ml of distilled or deionized water (15-25 °C) to the lyophilisate. Keep the control sample for about 15 minutes at rest in a light-protected place, gently agitate the vial occasionally until the lyophilisate is completely dissolved.

The samples should be treated the same way as patient samples and analyzed in accordance with the instructions of the instrument and reagent manufacturer.

D-Dimer: We request all participants to report their D-Dimer results as D-Dimer FEU (Fibrinogen equivalent units) and ask laboratories which obtain their test results in D-Dimer units (DDU) to convert their DDU result into an FEU value by using the reagent specific conversion factor that is expected to be provided in the instruction for use of the reagent kit (the conversion factor is approximately 2).

5. Dates and Submission of Test Results

Testing Periods: Please refer to the sample labels. Results can be submitted anytime within the testing period indicated on the sample labels.

For the evaluation of survey results, please indicate the reagent and method used - indication of the instrument is optional - and submit your results electronically to ESfEQA at <https://teqa-labv2.esfeqa.eu>.

Contact your local distributor of ESfEQA programs or ESfEQA directly if you need assistance for the registration, configuration and result submission in TEQA.

Quantitative results are generally reported with a value and a unit. The participant determines the number of digits for reporting. Results specified as e.g. '< below measuring range' or '< 0.02' are not valid. If the analyzer system displays such results, they shall be interpreted as following: For results within the measuring range and below the Limit of Quantification (LoQ) the obtained value should be reported. For results below the Limit of Detection (LoD), this limit should be reported. For samples that have analyte concentrations above the test range, the sample can be diluted (if recommended for particular applications) or the upper test range limit can be reported as result. Click or tap here to enter text.

6. Deadline for Data Submission

The deadline for submitting results is indicated on the sample labels (time zone GMT +1).

7. Reports and Certificates

The data will be evaluated by ESfEQA. The individual laboratory reports and certificates can be retrieved online at <https://teqa-labv2.esfeqa.eu>.