

Survey for HCV Molecular Testing (HCVm)

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Instructions for Use

Notes:

The samples should be considered as potentially infectious. The usual precautions in the laboratory for potential hazardous samples apply for these samples. The samples should be handled by trained personnel only.



WARNING: This product contains human source material and should be considered as potentially infectious.

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 qualitative and quantitative control material for External Quality Assessment (EQA) in medical laboratories for the following parameters:

HCV RNA qualitative
HCV RNA quantitative

2. Product Description

The lyophilized samples are prepared from human material with additives for formulation and stabilization. The samples are based on human serum. Positive samples contain inactivated whole virus.

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. Once the lyophilized survey samples are reconstituted (with sterile, nuclease-free ddH₂O), the analytical components will be stable for 8 hours when stored tightly capped at 2-8°C.

4. Sample Preparation and Analysis

In order to ensure that all lyophilized sample material is at the bottom of the tube, briefly spin the tube with the lyophilized control (e.g., 30 seconds at 14.000 rpm) before opening.

Reconstitute each lyophilized survey sample by pipetting the volume of Molecular/PCR grade ddH₂O (sterile, nuclease-free) indicated on the sample labels. Allow the control to reconstitute for 10-15 minutes, gently swirl and invert the closed tubes to ensure homogeneity.

After reconstitution of the samples, they should be handled/treated like patient samples (serum samples), and nucleic acid extraction and testing should be conducted in accordance with the instructions of the instrument and reagent manufacturers.

The **unit for reporting** quantitative HCV RNA results in TEQA is '**log₁₀ IU/mL**'.

It is recommended that participants using devices or methods, which provide quantitative values, **also submit values for the corresponding qualitative parameter.** This will enable the participants to compare their results with all laboratories.

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.

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>.