

General Information

Intended use:

FOB Turbilatex is a latex turbidimetric assay for the quantitative detection of human haemoglobin in human stool samples.

This assay is simple and widely applicable. Test results aid in a presumptive diagnosis of faecal occult blood (gastrointestinal bleeding).

For professional *in vitro* diagnostic use only.

FOB Turbilatex can be performed on every open chemistry analyser. Please follow the subsequent instructions in order to assure performance characteristics as describes in the instructions for use. This instruction has been validated by CerTest BIOTEC S.L Laboratories.

Additionally, please read the "Instructions for use" for instructions on operating and programming user defined test.

Reagents:

Materials provided by CerTest BIOTEC S.L.:

| Reagents | Quantity | Code |
|---|---|---|
| Turbidimetric reagents (R1 & R2) 200 Det/kit | R1: 2 vials, 2x22 mL R2: 1 vial, 1x13 mL | TL-022FB01 TL-022FB02 |
| Auxiliary Reagents | | |
| Calibration kit | Calibrator: 6 vials, 6x1 mL. | TL-022FB70, TL-022FB71 TL-022FB72 TL-022FB73 TL-022FB74 TL-022FB75 |
| Controls kit | Control C1, 2 vials, 2x1 mL/vial. Control C2, 2 vials, 2x 1 mL/vial. | TL-022FB08 TL-022FB09 |
| Sample diluent kit | 4 vials, 4x125 mL/vial | TL-022UN03E |
| Sample dilutions vials | 1x2 mL/vial 1x2,4 mL/vial | MST-0018MU MST-0019U |

Preparation of reagents:

R1 and R2 are ready to use.

Calibrators are ready to use.

Controls are ready to use

Storage and stability

Kit components must be stored at temperature indicated on the label. Do not freeze.

Reagents are stable up to the expiration date printed on the label, always considering that reagent containers must be properly closed to avoid any contamination, must be kept away from the sunlight and conserved at temperature indicated on the label of each reagent.

Specimen:

Collect enough quantity of human stool samples. These samples should be collected in clean and dry containers (no preservatives or transport media). The samples can be stored in the refrigerator (2-8°C) for 3 days prior to testing. Homogenise stool samples as thoroughly as possible prior to preparation.

The sample dilution vial with diluted sample can be stored for 7 days in the refrigerator (2-8°C) prior to testing.

Use FOB Turbilatex stool collection tubes for sample collections described the instructions for use.

Assay procedure

Application parameter set up:

Specific analyzers settings for FOB Turbilatex must be programmed onto the analyzer, see below. For instructions, consult the Cobas c301 (Roche) analyzer manual and instructions for use provided with the kit.

Loading of reagents:

Load reagents according to the Cobas c301 (Roche) analyzer manual.

Calibration curve establishment:

A 6-points calibration curve can be established in Cobas c301 (Roche) analyzer. For instructions consult analyzer manual.

Calibration stability:

Calibrate the system at least once a month is extremely recommended. Recalibrate the system when reagent lot is change or when the controls are out of the assigned range given in the control label and CoA.

QC controls:

FOB Turbilatex controls C1 and C2 must be assayed each day before running patient fecal sample extract to validate the calibration curve. The controls have assigned value ranges indicated on the label and certificate of analysis supplied. The control measurements must be within the indicated value range to obtain valid results for patient fecal extract. If the control values are out of range, follow next procedures: 1) Repeat QC control measurement, 2) Repeat calibration measurement.

Results:

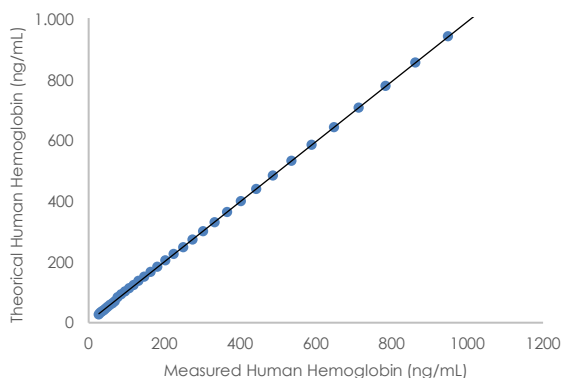
The results are evaluated automatically by the analyzer and presented in ng/mL.

Performance characteristics

The following results have been obtained during the validation of FOB Turbilatex on the Cobas c301 (Roche) analyzer.

Linearity:

FOB Turbilatex on Cobas c301 (Roche) instrument using calibrator kit is linear in the calibration range of 0-1000 ng/mL.



Measuring range:

FOB Turbilatex assay measuring range is 10-1000 ng/mL on the Cobas c301 analyser. Samples higher concentrated than 1000 ng/mL must be diluted for proper quantification by the user, using additional sample buffer.

Prozone effect

Using the reported parameters, no hook effect was observed up to 10 ng/mL. Samples with Haemoglobin concentration of 10 ng/mL give a typical positive result >1000 ng/mL.

Detection limit

Limit of detection (LOD): 8 ng/mL (*). The lower limit of detection of FOB Turbilatex was determined on 20 samples and 2 sample replicates as the mean value + 2·SD.

Limit of quantification (LOQ): 10 ng/mL (*). The lower limit of quantification is defined as the lowest actual amount of analysis that can be reliably detected; imprecision is < 20% as CV% on the Cobas c301 (Roche) instrument.

(* Data obtained by the analyser Biolis 24i (Tokyo Boeki)

Precision

FOB Turbilatex was tested with three different controls levels.

| | Low (50 ng/mL) | Medium (100 ng/mL) | High (500 ng/mL) |
|---------------------|-------------------|-----------------------|---------------------|
| N | 20 | 20 | 20 |
| Mean (ng/mL) | 47.8 | 104.0 | 494.1 |
| SD (ng/mL) | 3.0 | 6.0 | 12.2 |
| CV (%) | 6.2 | 5.7 | 2.4 |

Method comparison

Results obtained with FOB Turbilatex on the Cobas c301 (Roche) instrument were compared with those obtained with EKEN FOB Latex.

| | Sensitivity | Specificity |
|-------------------------------------|-------------|----------------|
| FOB Turbilatex vs FOB Latex® | 96% | >99% |

Shipping damage

Please notify your distributor, if this product was received damaged.

Symbols key

| | | | |
|------------|----------------------------------|------------|------------------------|
| IVD | For in vitro diagnostic use only | | Keep dry |
| | Consult instructions for use | | Temperature limitation |
| REF | Catalogue number | LOT | Lot number |
| | Use by | | Manufacturer |
| | Contains sufficient for <n> test | DIL | Sample diluent |
| | Keep out of the sunlight | | |

Manufacturer

CERTEST BIOTEC

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San Mateo de Gállego, Zaragoza (SPAIN)
www.certest.es

NOTES

Please refer to the instructions for use for the detailed information about the test on the following:

Synthesis; Principle; Precautions; Reagents; Specimen collection; Interpretation of results.

Cobas c301, Roche / Application parameters

| ASSAY PARAMETERS | |
|-------------------------|---------------------------|
| Std. No | 6 |
| R1 | 100 µL |
| Sample | 10 µL |
| R2 | 22 µL |
| Others | NA |
| Reaction mode | End point |
| Primary wavelength | 520 nm |
| Secondary wavelength | None |
| Direction | Increase |
| Reagent blank lecture | 12 cycle |
| Final Lecture | 40 cycle |
| Reaction time | 10 min |
| Linear range | 0-1000 ng/ml |
| CALIBRATION | |
| Calibration Method | Linear |
| Calibration set | 5 calibrators + Blank |
| Blank | Calibrator 1 (0 ng/ml) |
| Calibrator 1 | Calibrator 2 (50 ng/ml) |
| Calibrator 2 | Calibrator 3 (100 ng/ml) |
| Calibrator 3 | Calibrator 4 (250 ng/ml) |
| Calibrator 4 | Calibrator 5 (750 ng/ml) |
| Calibrator 5 | Calibrator 6 (1000 ng/ml) |
| STEPS | |
| Addition R1 | |
| Addition Sample | |
| Incubation | |
| Addition R2 | |
| Blank Lecture | Cycle 12 |
| Incubation | |
| Final lecture | Cycle 40 |