D-Dimer Assay



C€ REF C179D R1 1x30ml+R2 1x10ml Turbidimetry

Configuration

The D-Dimer Assay reagent is provided in packaging configuration:

Store 2 - 8°C

INTENDED USE

In vitro test for the quantitative determination of D-Dimer in human plasma on automated clinical chemistry analyzers for Clinical laboratories.

PRINCIPLE OF THE METHOD

Increase of D-Dimer in blood testifies the blood clot is formed and fibrinlytic activity has functioned. It is known that high value of D-Dimer is indicated in diseases such as malignant tumor, vascular disease.

Principle The D-dimer contained in the sample reacts with the latex sensitized with anti-human D-dimer monoclonal antibody (mouse) and forms aggregates, which are determined optically for calculation of D-dimer concentration.

REAGENTS

Diluent (R1)	Tris buffer 100 mM, NaN3 0.1%, pH 8.3 Surfactant 0.95 g/L.
Antibody (R2)	Suspension of anti-human D-Dimer mouse monoclonal antibody coated, pH 7.5, Latex particles (0.2%).PBS Buffer 0.1 mmol/l

CALIBRATION

The assay is calibrated to the Reference Material CRM 470/RPPHS (Institute of Reference of Materials and Measurements, IRMM). It must be used the CALIBRATOR (Cat. No. 179CAL) to calibrate the reagent. The reagent (both mono reagent and bireagent) should be recalibrated every month, when the controls are out of specifications, and when changing the reagent lot or the instrument settings

QUALITY CONTROL

It is recommended to use D-Dimer Quality Control (Cat. No. 179CTL) or BioRad / Siemens level 1 and level 2 for daily quality control.

Two levels of controls should be assayed at least once a day.

Values obtained should fall within a specified range

Each laboratory should establish its own internal Quality Control.

PREPARATION

Reagents: Ready to use

It is recommended to use the D-Dimer Calibrators (Cat. No. 179CAL) for calibration. Dissolve the lyophilized D-Dimer calibrator with 1ml distilled water, balance 10 minutes in room temperature. then dilute to 6 levels calibrators as follows.

Calibrator dilution	1	2	3	4	5	6
Calibrator (µL)		10	25	50	75	100
NaCl 9 g/L (µL)	100	90	75	50	25	-
Factor	0	0.1	0.25	0.5	0.75	1.0

Calibration frequency

Recalibration is recommended:

- · As a blank calibration after 24 hours
- As a blank calibration after reagent bottle change
 As a two point calibration every 30 days if the reagent always on-board
 As a two point calibration after reagent lot change
- As a two point calibration if required following quality control

Procedures

Calibration verification: Not necessary

STORAGE AND STABILITY

All the components of the kit are stable until the expiration date on the label when stored tightly closed at 2-8°C and contaminations are prevented during their use. Do not use reagents over the

Reagent deterioration: The presence of particles and turbidity.

Do not freeze; frozen Antibody or Diluent could change the functionality of the test

The final concentration of the components is below the limits imposed by Regulation (EC) No. 1272/2008 - CLP (and subsequent amendments) and Directive 88/379/CEE and subsequent amendments to the classification-packaging and labeling of dangerous substances

SAMPLES

In order to obtain a plasma sample for measurement, mix 1 part of citric acid (0.11mol/L) with 9 parts of venous whole blood to avoid air bubbles. Centrifuge immediately, not less than 3000 rpm (1500*g), centrifuge for 10 minutes, and collect the upper plasma.

It can be stored for 4 hours (fresh) under +15-25°C; it can be stored for 1 month under -18°C.

PROCEDURE

Reagent preparation:

R1: Ready for use: R2: Ready for use Reaction parameters + Reaction curve

Basic parameters of automatic biochemical instruments

Two-point endpoint method or fixed-time method :

	Blank	Sample/calibrator
	DIATIK	Sample/Cambrator
Sample/calibrator	-	5 μL
Dist. Water	5 µL	-
Reagent 1	300 µL	300 μL
Mix, incubate for 3 - 5 min., then add	:	
Reagent 2	100 μL	100 μL
Mix read absorbance (A1) within 20	sec incubate for 5 min	then read absorbance (A2) and

Gesan has instruction sheets for several automatic analyzers. Instructions for many of them are available on request.

CALCULATIONS

The analyzer automatically calculate the analyte concentration of each sample.

REFERENCE VALUES⁴

Samples with a D-Dimer concentration ≤ 0.50 µg/ml are considered normal. Use of an 8 µg/ml cut-off value is recommended for the VTE exclusion

The reference range should be determined by each hospital to confirm with the characteristics of the region being tested.

PERFORMANCE CHARACTERISTICS

0.2 μg/mL~20.0 μg/mL

Determine the samples with higher concentrations via the rerun function. In the event of a rerun the upper limit of the assay ranges is increased to approximately 20 μg/mL. These values are dependent on the lot specific values of the calibrators in use.

EP5	CV (%)				
	1.13 µg/mL	3.22 μg/mL	7.28 μg/mL		
Total	8.2%	5.2%	3.5%		
Within Run	1.7%	1.5%	1%		
Between Run	2.2%	1.9%	2.4%		
Between Day	7.7%	4.6%	2.3%		

6. Accuracy: Results obtained using this reagent (y) were compared to those obtained using an immunoturbidimetric method from Bayer. 20 samples ranging from 0.2 to 20 μ g/mL of D-Dimer were assayed. The correlation coefficient (r) was 0.97 and the regression equation v = 1.16x - 12.2

The results of the performance characteristics depend on the used analyzer.

INTERFERENCES

- 1. Lipemia (Intralipid): no interference up to 250 mg/dl of intralipid.
- Haemoglobin: no interference up to 500 mg/ dl.
- 3. Bilirubin: no interference up to 40mg/ dl.

1. Clinical diagnosis should not be made on findings of a single test result, but should integrate both clinical and laboratory data.

BIBLIOGRAPHY

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