

D-Dimer Turbi Latex

REF: 585 001 **50 Test** REF: 585 002 **100 Test** REF: 585 003 **80 Test**
 R1 Diluent 1 X 20 ml R1 Diluent 2 X 20 ml R1 Diluent 2 X 16 ml
 R2 Latex 1 X 5 ml R2 Latex 2 X 5 ml R2 Latex 2 X 4 ml
 Calibrator 1 vial Calibrator 1 vial Calibrator 1 vial

REF: 585 001-1 **50 Test** REF: 585 002-1 **100 Test** REF: 585 004 **500 Test**
 R1 Diluent 1 X 20 ml R1 Diluent 2 X 20 ml R1 Diluent 5 X 40 ml
 R2 Latex 1 X 5 ml R2 Latex 2 X 5 ml R2 Latex 5 X 10 ml
 Without calibrator Without calibrator Without calibrator

Intended Use

In vitro diagnostic reagents for the quantitative determination of D-Dimer in human plasma by means of particle-enhanced turbidimetric immunoassay.

Background

The D-dimer assay is specific for fibrin derivatives. In this assay, the presence of cross linked D-dimer domain is diagnostic for lysis of a fibrin clot, and confirm that thrombin was formed and factor XIII was activated with reactive fibrinolysis. Since fibrinogen derivatives do not contain the cross-linked D-dimer domain, they are not recognized by the D-dimer assay, even when present in high concentration. In other words, Fibrin derivatives in plasma containing D-Dimer (XDP) are specific markers for fibrinolysis, as opposed to fibrinogenolysis. D-dimers are detected by immunoassays using monoclonal antibodies specific for the cross-linked D-dimer domain in fibrinogen.

Test Principle

This D-dimer test is based upon reinforced immunoturbidimetry. Monoclonal anti D-dimer antibodies in the reagent react with the D-dimer antigen in the sample, forming antigen/antibody complexes that increase the working solution turbidity.

Reagents

R1 Reagent1

Tris-HCl 125 mM

R2 Latex reagent

Latex particles coated with mouse anti-human D-Dimer monoclonal antibodies.
Preservatives.

Calibrator

Human serum. D-Dimer concentration is stated on the vial label.

All raw materials of human origin used in the manufacture of this product showed no reactivity when tested for HBsAg, anti-HIV-1/2 and HCV with commercially available test methods. However, this product should be handled as though capable of transmitting infectious diseases

Storage and Stability











Reagents in the original vial are stable to the expiration date stated on the vial label when capped and stored at (2 - 8 °C). Do not freeze reagents. Open vial is stable for 3 months when stored at (2-8 °C).

Precautions and Warnings

For in vitro diagnostic use only. Do not pipette by mouth. Reagents containing sodium azide must be handled with precaution. Sodium azide can form explosive azides with lead and copper plumbing. Since absence of infectious agents cannot be proven, all specimens and reagents obtained from human blood should always be handled with precaution using established good laboratory practices. Disposal of all waste material should be in accordance with local guidelines.

As with other diagnostic tests, results should be interpreted considering all other test results and the clinical situation of the patient

SYMBOLS IN PRODUCT LABELLING

	Authorised Representative		Use by/Expiration Date
	For in-vitro diagnostic use		CAUTION. Consult instructions for use
	Batch Code/Lot number		Manufactured by
	Catalogue Number		(Xi) - Irritant
	Consult instructions for use		Temperature Limitation

Deterioration

The D-Dimer latex reagent should have a white, turbid appearance free of granular particulate. Visible agglutination or precipitation may be a sign of deterioration, and the reagent should be discarded.

The **Reagent1** should be clear and colourless. Any turbidity maybe sign of deterioration and reagent should be discarded.

Specimen Collection and Preservation

Citrated, platelet-poor plasma is used for the d-dimer assay. Citrated, platelet-poor plasma is prepared from venous blood collected into 3.2 % trisodium citrate at a ratio of 9 : 1. The citrate concentration must be adjusted in patients with a HCT >55%. Plasma should be separated as soon as possible after the specimen is obtained. D-dimers are stable for 8 hours in citrated plasma maintained at room temperature, 7days at 2 - 8 °C or 2 months at -20 °C.

Reagent Preparation

Spectrum D-dimer reagents (**R1 & R2**) are supplied ready-to-use and stable up to the expiry date labeled on the bottles when properly stored refrigerated at 2 – 8 °C.

D-dimer Calibrator: Reconstitute with distilled water as mentioned on vial label. Mix gently and incubate at room temperature for 10 minutes before use.

Stability: 1 month at -20 °C.

Calibration curve

Generate a calibration curve by serial dilutions of the supplied calibrator to cover the range between 0.2 µl/ml to the upper limit of the calibrator.

Cal. 6: Calibrator
 Cal. 5: 200 µl Cal. 6 + 200 µl saline
 Cal. 4: 200 µl Cal. 5 + 200 µl saline
 Cal. 3: 200 µl Cal. 4 + 200 µl saline
 Cal. 2: 200 µl Cal. 3 + 200 µl saline
 Cal. 1: 200 µl saline

Concentration	Cal.1	Cal.2	Cal.3	Cal.4	Cal.5	Cal.6
(for example: the undiluted C = 5.9 µg /ml)	0.18	0.37	0.74	1.48	2.95	5.9

Quality Control

Control sera are recommended to monitor the performance of manual and automated assay procedures. Each laboratory should establish its own Quality Control Scheme and corrective actions if controls do not meet the acceptable tolerances.

Materials required but not provided

D-Dimer control (Ref : 405 001)

Procedure

1. Bring the reagents and the photometer to 37°C
2. Assay conditions:

Wavelength 578 nm
Temperature 37°C
Cuvette 1cm light path
zero adjustment distilled water

3. Pipette into a cuvette :

Reagent1 (R1)	400 µl
Latex (R2)	100 µl
Calibrator or Sample	40 µl

4. Mix and read absorbance immediately (A1). After 3 minutes of the sample addition, read (A2).

Calculation

Calculate the absorbance difference (A2-A1) of each point of the calibration curve and plot the values obtained against the D-dimer concentration of calibrator dilution. D-dimer concentration in the sample is calculated by interpolation of its (A2-A1) in the calibration curve.

Detection limit

85-115%

Precision

	CV (%)	
	Intra-Run	Inter-Run
Low	1.13	2.31
Medium	1.65	0.98
High	1.17	1.01

Sensitivity

Up to 0.08 µg /ml .

Linearity

Up to 7.5 µg /ml .

specimens showing higher concentration should be diluted 1+2 using physiological saline and the assay is repeated. Results are multiplied by 3.

Interfering Substances

No interference for :

Heparin 50 µg/mL)
Na-citrate (1000 µg/mL)
Heparin (50 µg/mL)
Triglyceride (2500 µg/mL)
EDTA (5 µg/mL)
Haemoglobin (1000 µg/mL)

Bilirubin (20 µg/mL) and has interferences for: Turbidity (> 1.25%).

Expected Values

The determination of reference ranges for D-dimer concentrations of clinically healthy individuals is very difficult.

Suggested value in plasma with this method < 0.5 µg/ml.

These data are to be interpreted as a guide. Each laboratory should establish its own reference intervals.

Waste disposal

Disposal of all waste material should be in accordance with local guidelines.

References

- 1-Bick R.L. et al. thromb res 1992;65:785-90
- 2-Gaffney P.J. Fibrinolysis supplement 2. 1993;7:2-8
- 3- Bover, P. et al. int J Epidemiol 1994;23:2027
- 4- Janssen M.G. et al. Thromb Haemost 1997;77:262-

ORDERING INFORMATION

CATALOG NO.	QUANTITY
585 001	50 Test
585 002	100 Test
585 003	80 Test
585 001-1	50 Test
585 002-1	100 Test
585 004	500 Test



Egyptian Co for Biotechnology - Spectrum Diagnostics (S.A.E)

Obour city industrial area. block 20008 piece 19 A. Cairo. Egypt.

Tel: +202 4489 2248 - Fax: +202 4489 2247

www.spectrum-diagnostics.com

E-mail: info@spectrum-diagnostics.com



MDSS GmbH
Schiffgraben 41
30175 Hannover, Germany



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