



ACCENT-300 D-DIMER

DIAGNOSTIC KIT FOR DETERMINATION OF D-DIMER CONCENTRATION

INTRODUCTION

FDPs (Fibrin and Fibrinogen Degradation Products) are generic name for several degradation products that are formed as the result of plasmin mediated, proteolytic degradation of fibrin and fibrinogen. FDP assay become important test to diagnose or monitor fibrinolytic disorder, especially disseminated intravascular coagulation (DIC). But FDP assay also detects the degradation products of fibrinogenolysis. D-dimer assay is more specific for fibrinolysis because it can only be produced as the result of plasmin mediated, proteolytic degradation of fibrin.

METHOD PRINCIPLE

This D-dimer assay is a turbidimetric assay that utilizes antibody coated latex particles, in the presence of D-dimer, the particles aggregate to form larger aggregates and light scattering increases. The increase in scattered light is proportional to the amount of D-dimer in sample.

REAGENTS

Package	
1-Reagent	1 x 48 ml
2-Reagent	1 x 18 ml
D-Dimer Diluent	2 x 40 ml

Buffer (1-Reagent), latex (2-Reagent) and D-Dimer Diluent stored at 2-10°C are stable until expiry date printed on the package. The reagents are stable for 4 weeks on board the analyser at 2-10°C. Do not freeze the reagents. Protect from light and contamination!

Concentrations in the test

Tris(hydroxymethylo)aminomethane	0.38 mol/l
suspension of latex particles sensitized with	0.2 w/v%
anti-D-Dimer antibodies (mouse)	0.2 ₩/ ٧/0

Warnings and notes

- Products for in vitro diagnostic use only.
- The reagents must be used only for the purpose intended by suitably qualified laboratory personnel, under appropriate laboratory conditions.
- Products from human source have been tested for the HbsAg and antibodies to HCV and HIV and found to be non-reactive. However this material should be handled as thought capable of transmitting infectious disease.
- Products contain sodium azide (< 0.1%) as a preservative. Avoid contact with skin and mucous membranes.
- Allow the reagents to equilibrate at the room temperature before use.
- Swirl the latex reagent (2-Reagent) well before use.
- Do not mix different lots of reagent.
- Do not add new reagent to the remaining reagent.
- Pay attention not to contaminate cuvettes with dust or detergents.
- Sometimes turbidity appears in buffer reagent (1-Reagent) but it has no influence on assay results.
- Immunoassay cannot deny non-specific reaction and rarely occurs prozone effect when assay samples containing unusually high D-dimer level.

SPECIMEN

Plasma.

Nine volumes of fresh blood are collected in one volume of 0.11M trisodium citrate, followed by centrifugation at 3000 x g for 10-30 minutes. Use supernatant as plasma sample.

Samples containing more than 20 μ g/ml D-dimer should be reassayed, using a 1:10 sample dilution with D-Dimer Diluent.

If the measurement is done after 8 hours since separation of plasma from blood, the samples should be kept in refrigerator for up to 4 days. After 4 days the samples should be kept in freezer for up to 2 months.

Nevertheless it is recommended to perform the assay with freshly collected samples!

PROCEDURE

These reagents may be used in automatic analyser ACCENT-300. 1-Reagent and 2-Reagent are ready to use. For reagent blank 0.9% NaCl is recommended.

APPLICATION

Parameters			
No.	55	Prim.Wave.	578
Test	D-DIM	Sec.Wave.	
Method	Kinetic	Sample Vol.	6
Direction	Ascend	R1 Vol.	180
Unit	µg/ml	R2 Vol.	60
Decimals	2	Line. Limit	20
Incubation	5	Antigen Check	ν
Reaction	3 18	Substrat	0
R1 Blank		Mix. R Blank	
Lower	0	Lower	0
Upper	0	Upper	0
Response		Linearity	
Lower	-2.5	Lower	
Upper	2.5	Upper	
S 1- 37 - 1	20	E-11 Marra	
Sample Vol.	30	Full Name	D-DIM
Dilution	10	Print No.	55
Calibration			

Calibration	
Rule	Spline
Replicates	1
Interval	0
Sensitivity	0
Correlation	0
Difference	2.5
Blank Response	0 2.5
Coefficient Difference	0
Non-linear SD	0

REFERENCE VALUES¹

 plasma
 < 0.5 μg/ml</th>

 It is recommended for each laboratory to establish its own reference ranges for local population.

QUALITY CONTROL

For internal quality control it is recommended to use the CORMAY D-DIMER CONTROLS (Cat. No 4-459) with each batch of samples. For the calibration of automatic analysers systems the CORMAY D-DIMER CALIBRATOR (Cat. No 4-259) is recommended. The calibration curve should be prepared every 4 weeks, with change of reagent lot number or as required e.g. quality control findings outside the specified range.

PERFORMANCE CHARACTERISTICS

These metrological characteristics have been obtained using the automatic analysers BS-400 and TBA80FR. Results may vary if a different instrument is used.

- Sensitivity: 0.3 µg/ml.
- Linearity: up to 20 µg/ml.

For higher concentrations dilute the sample with D-Dimer Diluent in the ratio of 1 to 10 and repeat the assay. Multiply the result by 11.

Specificity / Interferences

Haemoglobin up to 4.6 g/dl, conjugated bilirubin up to 19.6 mg/dl, free bilirubin up to 18.4 mg/dl, RF up to 500 IU/ml do not interfere with the test.

Precision

Repeatability (run to run)	Mean	SD	CV
n = 10	[µg/ml]	[µg/ml]	[%]
level 1	2.50	0.05	1.97
level 2	9.20	0.54	5.91

Reproducibility (day to day)	Mean	SD	CV
n = 80	[µg/ml]	[µg/ml]	[%]
level 1	2.60	0.11	4.38
level 2	8.99	0.48	5.36

Method comparison

A comparison between CORMAY kit (y) and another commercially available kit based on latex turbidimetric assay (x) using 54 samples gave following results:

 $y = 0.991 x + 0.028 \mu g/ml;$ R = 0.9989

(R – correlation coefficient)

WASTE MANAGEMENT

Please refer to local legal requirements.

LITERATURE

1. Alan H. B. Wu, Tietz Clinical Guide to Laboratory Tests, W.B. Saunders Company, 4th edition, 332 (2006).

Date of issue: 05. 2012.

MANUFACTURER

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