

KOS Diagnostic Lab

(A Unit of KOS Healthcare)



Dr. Vinay Chopra MD (Pathology & Microbiology) Chairman & Consultant Pathologist

Dr. Yugam Chopra MD (Pathology) CEO & Consultant Pathologist

ng/mL

0.00 - 500.00

NAME : Mr. SUBHASH GUPTA

AGE/ GENDER : 74 YRS/MALE **PATIENT ID** : 1569749

COLLECTED BY REG. NO./LAB NO. :012408030051

REFERRED BY **REGISTRATION DATE** : 03/Aug/2024 04:32 PM BARCODE NO. :01514377 **COLLECTION DATE** : 03/Aug/2024 04:39PM CLIENT CODE. : KOS DIAGNOSTIC LAB REPORTING DATE : 03/Aug/2024 06:16PM

2550H

CLIENT ADDRESS : 6349/1, NICHOLSON ROAD, AMBALA CANTT

Test Name Value Unit **Biological Reference interval**

HAEMATOLOGY

D-DIMER (QUANTITATIVE)

D - DIMER (QUANTITATIVE)

by EFIA (FLUORESCENT ENZYME IMMUNOASSAY)

During coagulation sequence of reactions occuring in the body in response to variety of external and/or internal stimuli. The enzymaticcascade reaction reaction terminates in the conversion of fibrinogen to fibrin by enzyme thrombin. The fibrin gel is then converted to a stable fibrin clot. The fibrin network is dissolved by enzyme plasmin to generate cross-linked FIBRIN DEGRADATON PRODUCTS. D-DIMER is the smallest plasmin resistant molecular unit present within FDP.

INCREASED D-DIMER IS SEEN IN FOLLOWING CONDITIONS:

1. Door Voin Thrombosis (DVT)

1.Deep Vein Thrombosis (DVT)

2. Venous Thromboembolism

3. Recent Surgery

4.Trauma

5.Infection

6.Liver disease 7.Pregnancy

8. Eclampsia

9.Heart Disease

10.Some cancers

11.Elderly

NOTE:

1. A normal or low D-dimer helps to rule out clotting as cause of symtoms.

2. D- DIMER is approximately 6 hours in circulation of individuals with normal renal functions. Patients with stabilized clots and not going active fibrin deposition and plasmin activation may not give detectable D-Dimer elevation, anti-coagulant therapy.

3. In Pulmonary Embolism (PE), the larger the clot size, higher the expected level of of circulating D-Dimer. Conversely, theamount of D – DIMER release from very small clots may be diluted by circulation and may not give detectable increase.

4. Fibrionolysis is a highly regulated process and in dynamic delicate balance. In case of hereditary, acquired deficiency and dysfunction of fibrinogen, the rate of fibrinolysis will be altered thereby not giving detectable D-Dimer level.

5. False positive may be seen with high levels of rheumatoid factor, bilirubin, lipemic sera and haemolysed blood

RECHECKED.

*** End Of Report ***



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