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ORGAN

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Name : **SUFIA BIBI**
Refr.By : **MO. S.D.HOSPITAL**

ID No. : **BRB-27**

Sex / Age : Female / 54 Year
Received On : 02/08/2018
Reported On : 02/08/2018
Lab No. : 1364 **2018/08/BRB/2**

DEPARTMENT OF SEROLOGY

INVESTIGATION	RESULT	UNIT	REFERENCE VALUE
HIV(I &II) Ab & P24 ANTIGEN (Method : Chemiluminescent Microparticle Immunoassay (CMIA))	: 0.11	S/CO	< 1.00 NON REACTIVE > 1.00 REACTIVE
Hepatitis B Surface Antigen (HBsAg) (Method : Chemiluminescent Microparticle Immunoassay (CMIA))	: 0.29	S/CO	< 1.00 NON REACTIVE > 1.00 REACTIVE
ANTI HCV (Method : Chemiluminescent Microparticle Immunoassay (CMIA))	: 0.09	S/CO	< 1.00 NON REACTIVE > 1.00 REACTIVE

* End of Report *

NOTE : THIS IS SCREENING TEST. REACTIVE RESULT SHOULD BE CONFIRMED BY WESTERN BLOT OR PCR METHOD(FOR HIV TEST), RT PCR METHOD(FOR HBSAG), AND A DEFINITIVE DIAGNOSIS IS BASED ON CLINICAL HISTORY.

***THIS IS A SCREENING TEST AND THERE MAY BE FALSE POSITIVE AND FALSE NEGATIVE RESULT DUE TO VERIOUS REASONS.

INTERPRETATION FOR HCV :

- Anti HCV Antibodies appear in serum 2-6 months after infection. However, the window period may exceed to 1 year.
- * A reactive test for Anti-HCV Ab implies infection with HCV but not infectivity or immunity.
- * Anti HCV may be negative in those cases of HCV infection where disease is acute, self-limiting and transient.
- * A non-reactive result does not exclude the possibility of exposure to or infection with HCV.
- * False reactive results may be due to non-specific binding to membrane or in patients with Autoimmune liver disease.
- * All reactive samples are to be confirmed by supplemental assays and a definitive diagnosis is based on clinical history.

Immuno Assay Instryments : Abbott ARCHITECT (Highly Innovative Technology), Made in Germany.

Reagent-Kit : Abbott 2nd Generation Kit(HBSAG TEST), Made in Germany.

Reagent-Kit : Abbott 3rd Generation Kit(HCV TEST), Made in Germany.

Reagent-Kit : Abbott HIV 4th Generation Antigen / Antibody Combo Kit, Made in Germany. US FDA and European FDA Approved.

OSS

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