# **MOLQ LABORATORY (A UNIT OF MOLECULAR QUEST HEALTHCARE** PVT. LTD.)

# FIRST TRIMESTER DOUBLE MARKER

PATIENT INFORMATION NAME MRS BABITA W/O NARENDER PATIENT CODE: 01180912386

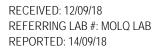
DOB: 20/04/94 (DDMMYY) LMP: 13/06/18 EDD: 25/03/19 PHYSICIAN:

#### **SPECIMEN**

SPECIMEN CODE: MOLQ LAB COLLECTION DATE: 12/09/18

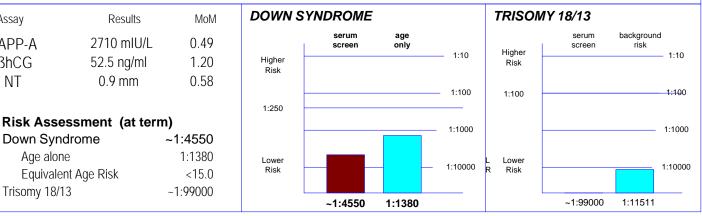
## **CLINICAL RESULTS**

Assay Results PAPP-A 2710 mIU/L 0.49 fßhCG 52.5 ng/ml 1.20 0.9 mm NT 0.58**Risk Assessment (at term)** Down Syndrome ~1:4550 Age alone 1:1380



#### **CLINICAL INFORMATION**

GESTATIONAL AGE: 12 weeks 2 days by U/S (12 wks 3 days on 13/09/18) MATERNAL AGE AT TERM: 24.9 years MATERNAL WEIGHT: 42.0 kg MATERNAL RACE: INDIAN MATERNAL IDDM: Not specified (Non-diabetic assumed) **GESTATION:** Singleton SCREENING STATUS: Initial sample PARA / GRAVIDA: 0 / 1



#### (based on partial information supplied) Interpretation\*

DOWN SYNDROME

# **Screen Negative**

The risk of Down syndrome is LESS than the screening cut-off.

**TRISOMY 18/13** 

### **Screen Negative**

The risk of trisomy 18 is less than the screening cut-off.