

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

FIRST TRIMESTER DOUBLE MARKER

PATIENT INFORMATION
NAME MRS. KAVITA

PATIENT CODE: 011807200292 DOB: 10/11/88 (DDMMYY)

LMP: 18/04/18 EDD: 26/01/19 PHYSICIAN:

SPECIMEN

SPECIMEN CODE: MOLQ LAB COLLECTION DATE: 21/07/18

RECEIVED: 21/07/18

REFERRING LAB #: MOLQ LAB

REPORTED: 23/07/18

CLINICAL INFORMATION

GESTATIONAL AGE: 13 weeks 0 day by U/S (12 wks 6 days on 20/07/18) MATERNAL AGE AT TERM: 30.2 years

MATERNAL WEIGHT: 51.0 kg MATERNAL RACE: INDIAN

MATERNAL IDDM: Not specified (Non-diabetic assumed)

GESTATION: Twins - Chorionicity unknown SCREENING STATUS: Initial sample

PARA / GRAVIDA: 0 / 1

Assay	Results	MoM	DOWN SYNDROME			TRISOMY 18/13				
PAPP-A	5120 mIU/L	0.91	1	serum screen	age only		Higher	serum screen	background risk	
fßhCG	49.8 ng/ml	1.26	Higher Risk			— 1:10	Risk			— 1:10
Ultrasound CRL	NT MoM	Nasal Bone				1:100	1:100			1:100
Fetus A: 64.9 Fetus B: 69.1	1.0 mm 0.60 0.9 mm 0.52		1:250							
Risk Assessment (at term) Down Syndrome ~1:16800						_ 1:1000				1:100
Age alone Fetus A: Fetus B: Trisomy 18/13		1:943 ~1:33500 ~1:33500 ~1:99000	Lower Risk			- 1:10000	L Lower R Risk			1:1000
Fetus A: Fetus B:		~1:99000 ~1:99000		~1:16800	1:943			~1:99000	1:7285	

Interpretation* (based on partial information supplied)

DOWN SYNDROME Screen Negative

This patient's relative risk for Down syndrome has been determined by dividing the assay results by the average analyte level in twin pregnancies, and then calculating the risk associated with a singleton pregnancy. This relative risk calculation places the patient below the screening cut-off for Down syndrome.

TRISOMY 18/13 Screen Negative