

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

FIRST TRIMESTER DOUBLE MARKER

PATIENT INFORMATION

NAME MRS PADMINI FETUS B

PATIENT CODE: 10359222

DOB: 08/06/93 (DDMMYY)

LMP: 13/05/18 EDD: 21/02/19 PHYSICIAN:

SPECIMEN

SPECIMEN CODE: MOLQ LAB COLLECTION DATE: 17/08/18

RECEIVED: 17/08/18

REFERRING LAB #: MOLQ LAB

REPORTED: 18/08/18

CLINICAL INFORMATION

GESTATIONAL AGE: 13 weeks 1 day by U/S (12 wks 4 days on 13/08/18) MATERNAL AGE AT TERM: 25.7 years MATERNAL WEIGHT: 76.0 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	6550 mIU/L	1.81		serum screen	age only	4.40	Higher	serum screen	background risk
fßhCG	296.2 ng/ml	7.69	Higher Risk			— 1:10	Risk		1:10
Ultrasound CRL	. NT MoM	Nasal Bone				1:100	1:100		1:100
Fetus A: 60.6 Fetus B: 60.6	1.6 mm 1.02		1:250				1.100		1.100
Risk Assessment (at term) Down Syndrome ~1:994						1:1000	_		1:1000
Age alone	OITIC	1:1340							
Fetus A:		~1:1990	Lower			- 1:10000	L Lower		1:1000
Fetus B:		~1:1990	Risk				R Risk		111000
Trisomy 18/13		~1:99000	L				L		
Fetus A: Fetus B:		~1:99000 ~1:99000		~1:994	1:1340			~1:99000	1:11074

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