

SAVITHRI R.

Date of birth: 11 July 2001
Examination date: 07 June 2024

Address: Chennai Hospital no.: 2400088144

Referring doctor: Dr Nandhini Kumari Quality Scans
Address: Chennai

Maternal characteristics and history

Ethnic origin: South Asian (Indian, Pakistani, Bangladeshi).
Parity: 0.
Maternal weight: 80.0 kg.
Smoking in this pregnancy: no; Diabetes Mellitus: no.
Conception: spontaneous;
last period: 02 March 2024

EDD by dates: 07 December 2024

First Trimester Ultrasound

Gestational age: 13 weeks + 2 days from CRL

EDD by scan: 11 December 2024

Crown-rump length (CRL) 71.0 mm 
Nuchal translucency (NT) 1.60 mm

Chromosomal markers:

Nasal bone: present.

Maternal Serum Biochemistry

Sample 2400088144, taken on: 07 June 2024, analysed on: 11 June 2024.

Free β -hCG	29.19 IU/l	Roche	equivalent to	1.196 MoM
PAPP-A	4.371 IU/l	Roche	equivalent to	1.114 MoM

FMF Operator: NANDHINIKUMARI BASKAR, FMF Id: 61379

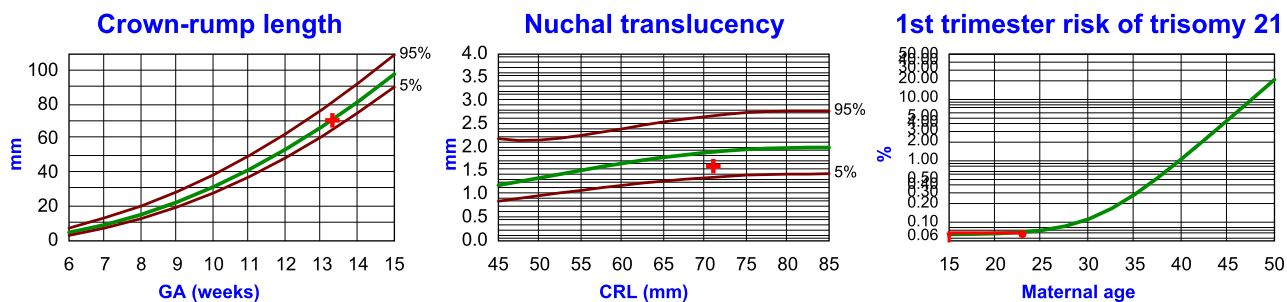
Condition	Background risk	Adjusted risk
Trisomy 21	1 in 1451	<1 in 20000
Trisomy 18	1 in 17123	<1 in 20000
Trisomy 13	<1 in 20000	<1 in 20000

The background risk for aneuploidies is based on maternal age (22 years). The adjusted risk is the risk at term, calculated on the basis of the background risk, ultrasound factors (fetal nuchal translucency thickness, nasal bone) and maternal serum biochemistry (PAPP-A, free beta-hCG).

Biochemical marker medians used to calculate MoMs are corrected as necessary according to several maternal characteristics including racial origin, weight, height, smoking, method of conception and parity.

The estimated risk is calculated by the FMF-2018 software (version 4.6) and is based on findings from extensive research coordinated by the Fetal Medicine Foundation (UK Registered charity 1037116). The risk is only valid if the ultrasound scan was performed by a

sonographer who has been accredited by the Fetal Medicine Foundation and has submitted results for regular audit (see www.fetalmedicine.org).



Comments

INTERPRETATION : The first trimester screening for the given sample is found **SCREEN NEGATIVE** .

Please Note: The above interpretation is based on a cut off of 1:250 for T21 , 1:100 for T13 & T18

Reviewed By

Dr. Suresh Bhanushali MD (Path)
Consultant Pathologist

Notes

1. Quality of the Down's syndrome screening program (Biochemical values, MoMs and Risk assessments) monitored by UKNEQAS on an ongoing basis.
2. This interpretation assumes that patient and specimen details are accurate and correct.
3. Lilac Insights does not bear responsibility for the NT & CRL measurements.
4. This is a risk estimation test and not a diagnostic test. An increased risk result does not mean that the fetus is affected and a low risk result does not mean that the fetus is unaffected.