

Double Marker
Test

SNEHA SHOKER (147599)

Date of birth: 19 September 1991
 Examination date: 19 September 2025

Address: Ranchi Hospital no.: 2500212850

Referring doctor: Dr Snigdha Chatterjee
 Address: Ranchi

Santevita Hospital

Maternal characteristics and history

Ethnic origin: South Asian (Indian, Pakistani, Bangladeshi).
 Parity: 0.

Maternal weight: 62.7 kg.

Smoking in this pregnancy: no; Diabetes Mellitus: no.

Conception: spontaneous;

last period: 25 June 2025

EDD by dates: 01 April 2026

First Trimester Ultrasound

Gestational age: 12 weeks + 6 days from CRL

EDD by scan: 28 March 2026

Fetal heart activity	visualised	
FHR	160 bpm	—○—
Crown-rump length (CRL)	66.0 mm	—○—
Nuchal translucency (NT)	1.60 mm	

Chromosomal markers:

Nasal bone: present.

Maternal Serum Biochemistry

Sample 2500212850, taken on: 19 September 2025, analysed on: 22 September 2025.

Free β-hCG	20.79 IU/l	Roche	equivalent to 0.634 MoM
PAPP-A	5.196 IU/l	Roche	equivalent to 1.173 MoM

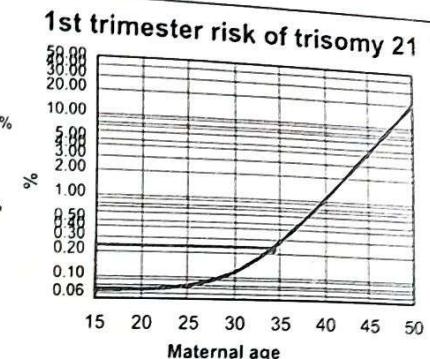
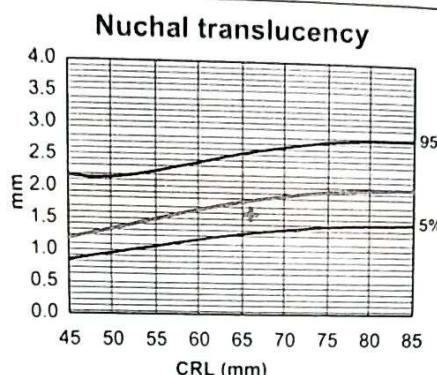
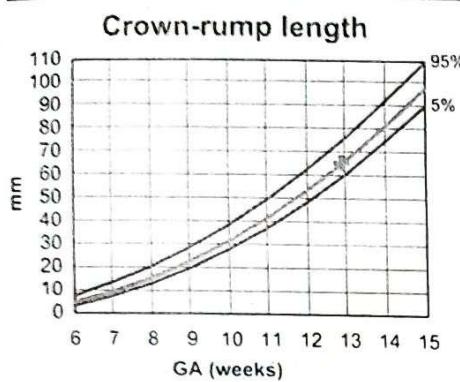
FMF Operator: SHIKHA ANAND, FMF Id: 281678

Condition	Background risk	Adjusted risk
Trisomy 21	1 in 473	1 in 9451
Trisomy 18	1 in 5577	<1 in 20000
Trisomy 13	1 in 13130	<1 in 20000

The background risk for aneuploidies is based on maternal age (34 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, fetal heart rate) 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.

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DEPARTMENT OF IMAGING - ULTRA SONOGRAPHY

Patient name	Mrs. SNEHA SHOKER	Age/Sex	33 Years / Female
Patient Id	2025-1135 / 147599	Visit no	1
Referred by	Dr. Snigdha Chatterjee	Visit date	17/11/2025
LMP date	25/06/2025, LMP EDD: 01/04/2026[20W 5D]		

USG Anomaly Scan Report

Indication(s)

Primigravida

Level I scan- Normal

Double marker done- Low risk T21 1:9451

Real time B-mode ultrasonography of gravid uterus done.

Route: Transabdominal

Single intrauterine gestation

Maternal

Cervix measured 3.4 cm in length.

Internal os closed

Maternal Doppler:

The right uterine artery shows a normal flow pattern.

The left uterine artery shows a normal flow pattern.

The mean pulsatility index of uterine arteries is within normal range.

Right Uterine	1.23	—●— (65%)
Left Uterine	1.32	—●— (73%)
Mean PI	1.275	—●— (69%)

Fetus

Survey

Presentation - Variable

Placenta - Posterior

Lower limit of placenta 2.8cm away from os

Liquor - Adequate

Umbilical cord - Two arteries and one vein

Fetal activity present

Cardiac activity present

Fetal heart rate - 142 bpm

Biometry (mm)

BPD	50.9, 21W 2D	—●— (68%)
HC	190.4, 21W 2D	—●— (63%)
AC	170.1, 22W	—●— (76%)
FL-Rt	36.1, 21W 3D	—●— (59%)

Long bones	Right (mm)	
Tibia	31.4, 21W 5D	—●— (74%)
Fibula	29.6, 20W 4D	—●— (47%)
Humerus	34.4, 21W 5D	—●— (75%)
Radius	29.5, 21W 3D	—●— (57%)
Ulna	32.8, 21W 6D	—●— (78%)



DEPARTMENT OF IMAGING - ULTRA SONOGRAPHY

Mrs. SNEHA SHOKER / 2025-1135 / 17/11/2025 / Visit No 1

Abdomen

Abdominal situs appeared normal.
Stomach and bowel appeared normal.
Normal bowel pattern appropriate for the gestation seen.
No evidence of ascites.
Abdominal wall intact.

KUB

Right renal pelvis measured 3.5 mm
Left renal pelvis measured 3.1 mm
Right and Left kidneys appeared normal.
Bladder appeared normal

Extremities

All fetal long bones visualized and appear normal for the period of gestation.
Both feet appeared normal

Impression

Single gestation corresponding to a gestational age of 21 weeks 4 days.
Detailed study of fetus does not reveal any obvious anomaly detectable by ultrasound during this period of gestation

Note is made of mild increased NF measuring 6 mm.

Counselling: It is a soft marker for trisomy 21 with a likelihood ratio of 3.7. Patient has undergone double marker test which is low risk with a sensitivity of 85%,. Couple understand that 100% chromosomal normalcy can be obtained by amniocentesis which is an invasive test with 1 in 1000 risk of pregnancy loss in this pregnancy. NIPT can be offered as it has better sensitivity for common aneuploidies.

Disclaimer:

Patient has been explained about limitations of scan due suboptimal visualization because of maternal habitus. Not all congenital anomalies (especially of the limbs, palate, heart and skin) can be detected by ultrasound. Overall detection rate of major congenital abnormalities is about 70%. Minor abnormality like polydactyly, soft palate abnormality are not looked up at routine level 2 scan.

Couple understand that ultrasound cannot rule out functional, chromosomal and genetic abnormalities. Normal means structurally normal for the gestation within reasonable expectations for available technology and patient body habitus. Certain diseases may evolve during pregnancy and may not be seen at the time of level II/ anomaly scan. The reported prevalence of fetal anomalies diagnosed in third trimester is 3.7 per 1000 women, these are typically genitourinary tract, brain and heart.

Conditions like Trisomy 21 may have normal ultrasound. 100% confirmation is obtained by amniocentesis which has a risk of abortion 1:1000. Serum screening like nips also has 98% sensitivity for trisomy 21. Please do not hesitate to contact us if you have any further questions.





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Patient Name : **Mrs. Sneha Shoker** Sex / Age : Female/33 Yrs
 Date / Time : 18/11/2025 12:57PM No : 1 UHID No : **147599**
 Sponsor : SANTEVITA TARIFF Address : MAIN RD RANCHI Contact No. :
 Visit Type : First Visit

Consultant Name

Dr.SnigdhaChatterjee
Gynaecology & Obstetric
 Med. Reg.No.: JMC-756/2006

Vital Signs

BP : 111/64 ^{mmHg} Pulse : 90 b^m RR : 18 b^m
 Temp : 98 F Weight : 55.6 kg Height : 154 cm
 BMI : 23 SpO2 : 100 %

Presenting Complaints :

G, 21 wks.

LMP - 25/6/25

EDD - 14/12/26.

Anomaly free →

SLF : 21 weeks

PL - 2.8 cm post pl.

W - ord. +

CD +
Cx - 3.4 cm

Relevant Past Medical / Family history :

Local / Systemic Examination :

Diet: As Advised

Wt 20 weeks

↑ cd NF - 6 mm.

Investigations :

NIPT

TT, ✓

Next Follow- up :

This is an important document, Please keep it safe and carry it in your next visit.

Contact No: +91 7360033390
 (0651) 7111555

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e-mail : info@santevitahospital.com Website : www.santevitahospital.com

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Provisional Diagnosis :

G, 21 weeks.

Medication :

Rx

- ① Livogen Z - 1 daily 9 AM
→ 30 + 30 +
- ② Calcium forte - 1 daily 9 AM.
③ 30
- ③ Potassium bromate - 2 kg C
milk. 4 PM.

Follow - up advice :

SV - 25 weeks (11/12 - 20/12)
, F. Echo
(CBC
, KFT.
, APGE review

Advice for Admission : Yes / No

If yes, to Wards / Day Care / ICU / NICU
(OGTT (75)
F/1hr/2hr)


Consultant



भारत सरकार

GOVERNMENT OF INDIA



स्नेहा शोकर

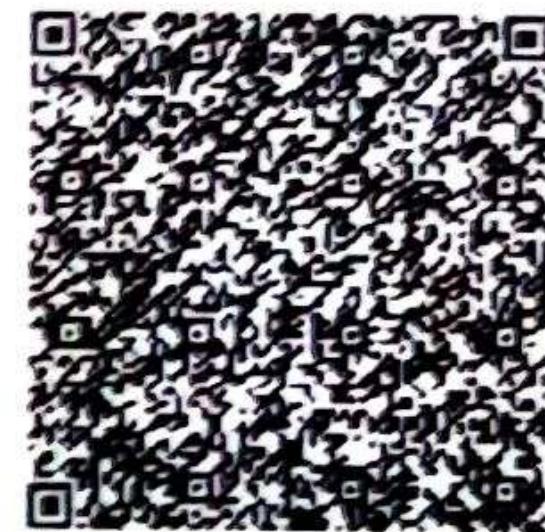
Sneha Shoker

जन्म तिथि/DOB: 19/09/1991

महिला/ FEMALE

Mobile No: 7488753196

2946 5108 6945



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भारतीय विशिष्ट पहचान प्राधिकरण
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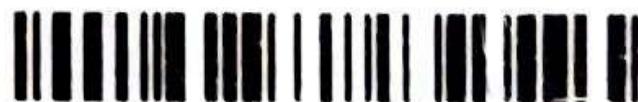
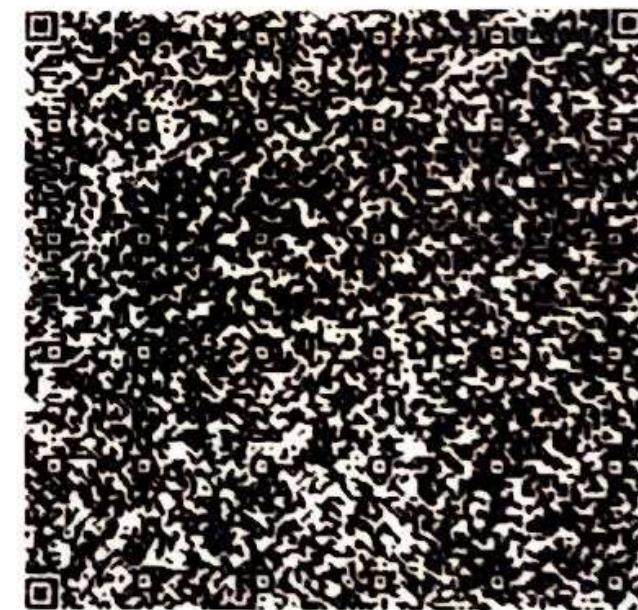
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पता:

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झारखण्ड - 834001

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