

# Dr. Sonal Phadtare

Fetal Medicine Consultant  
PCPNDT Reg. No. PH/CSP/SDH-Baramati/746/2021  
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Reg. No.: 2013/05/1597  
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## BARAMATI FETAL MEDICINE CENTRE



### First Trimester Screening Report

#### Snehal Malgunde

Date of birth: 15 January 2002  
Referring doctor:

Examination date: 08 June 2024  
Patient id: 1141

#### History

Ethnic origin: South Asian (Indian, Pakistani, Bangladeshi).  
Parity: 0; Spontaneous deliveries between 16-30 weeks: 0.  
Maternal weight: 55.0 kg; Height: 160.0 cm.  
Smoking in this pregnancy: no; Diabetes Mellitus: no; Chronic hypertension: no; Systemic lupus erythematosus: no; Antiphospholipid syndrome: no.  
Conception: spontaneous;  
Last period: 06 March 2024

EDD by dates: 11 December 2024

#### First Trimester Ultrasound

US system: Voluson E 10 (BT 20). View: good.  
Gestational age: 13 weeks + 3 days from dates

EDD by scan: 11 December 2024

Findings	alive fetus	
Fetal heart activity	visualised	
FHR	156 bpm	
Crown-rump length (CRL)	72.5 mm	
Nuchal translucency (NT)	1.90 mm	
Biparietal diameter (BPD)	25.4 mm	
Head circumference (HC)	84.0 mm	
Abdominal circumference (AC)	70.4 mm	
Femur length (FL)	12.2 mm	
Intracranial translucency	present, 2.5 mm	
Ductus Venosus PI	0.85	
Placenta	appears normal	
Amniotic fluid	normal	

#### Chromosomal markers:

Nasal bone: present; Tricuspid Doppler: normal.

#### Fetal anatomy:

Skull/brain: appears normal; Spine: appears normal; Heart: four chambers three vessels seen; Abdomen: appears normal; Stomach visible; Bladder / Kidneys: visible; Hands: both visible; Feet: both visible.  
Comments: BS:BSOB ratio- 0.56.

Uterine artery mean PI:	1.255	equivalent to 0.780 MoM
Mean Arterial Pressure:	83.333 mmHg	equivalent to 1.0074 MoM
Endocervical length:	30.0 mm	

#### Risk calculation

Patient counselled and consent given.

FMF Operator: Sonal Phadtare, FMF Id: 198408

Condition	Background risk	Adjusted risk
Trisomy 21	1 in 1076	1 in 8090

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The nasal bone is present. The Tricuspid valve and Ductus venosus Doppler are normal.

I have offered the option of early risk reassessment for aneuploidies (free bHCG, PAPP-A, PLGF and AFP), i.e. first trimester Quadruple marker test. This also assesses the risk for fetal Neural Tube Defects (NTDs), Fetal Growth Restrictions (FGR) and maternal pre-eclampsia (PE).

If the risk for maternal PE is more than 1:100, scientific evidence based on large study (the ASPRE study) from the Fetal Medicine Foundation, UK, has shown that, administration of Aspirin 150mg to the mother at bedtime, would reduce the effects of PE for the mother by 50-80%.

The cervix measures 30 mm with no evidence of funneling. The uterine artery Doppler is normal for this gestation.

The detection rate for chromosomal abnormalities with various screening tests are as follows-

First trimester NT scan only- 75%

First trimester Combined (NT + maternal blood test)- 80-85%

First trimester Combined test with 1T Quad marker- early screening for aneuploidies + fetal NTDs + FGR + maternal PE- 90% detection rate

Sequential screening (Combined + 2nd trimester Quadruple at 15-19w + Genetic sonogram at 18-20w) - 95%

Maternal blood test for cell free fetal DNA- 99%

Invasive testing (CVS/Amniocentesis), which is the definitive test has a procedure related risk of miscarriage about 1:300.

After counselling, the couple decided against invasive testing.

Please note for all the future visits to ANY SCAN CENTRE, it is mandatory by the GOI to produce prior to the scan

1. Photocopy (xerox) of Government approved photo ID card of the prospective mother (Aadhaar card, passport, voter ID card, driving licences etc)
2. Referral letter from your Doctor with indication for the scan with her/ his SEAL and SIGNATURE.

Also, please keep extra time to fill GOI mandated "F- Form" before the scan which needs to be submitted online prior to the scan.

Please note:

1. All anomalies cannot be ruled out on ultrasound due to technical limitations, maternal factors like amount of liquor, maternal previous scar, advanced gestational age etc. and fetal conditions like multiple pregnancies, fetal position, late appearance of anomalies etc.
2. Absence of fetal anomalies on ultrasound scan does not absolutely rule out the possibility of having one.
3. The opinion reported is based on data generated by the computer, clinical correlation is required for deciding a treatment.

I Dr. Sonal Phadtare (M.S OBGY; Fetal Medicine Consultant), declare that while conducting ultrasound/ image scanning of Mrs. Snehal Malgunde, I have neither detected nor disclosed the sex of her fetus to anybody in any manner.

**Recommendations** Rescan at 18- 20 weeks to assess the fetal anatomy in detail (15th to 25th July)

DR SONAL PHADTARE  
MMC 2013/05/1597  
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Trisomy 18 1 in 2733 1 in 10898  
Trisomy 13 1 in 8545 <1 in 20000

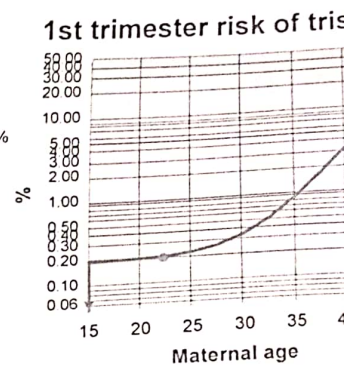
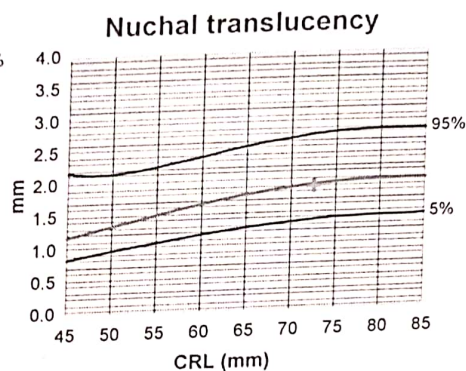
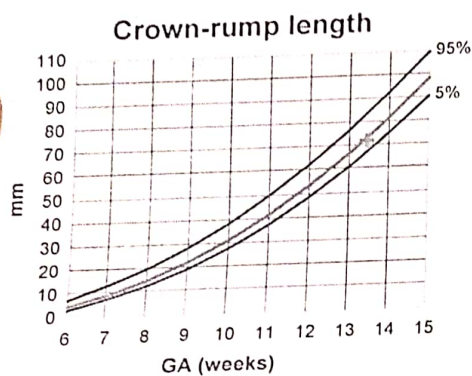
Preeclampsia before 37 weeks 1 in 408  
Fetal growth restriction before 37 weeks 1 in 232  
Spontaneous delivery before 34 weeks 1 in 71

The background risk for aneuploidies is based on maternal age (22 years). The adjusted risk is the risk at the time of screening, calculated from the background risk and ultrasound factors (fetal nuchal translucency thickness, nasal bone, fetal heart rate).

Risks for preeclampsia and fetal growth restriction are based on maternal demographic characteristics, medical and obstetric history, blood pressure (MAP) and uterine artery Doppler. The risk of spontaneous delivery before 34 weeks is based on maternal characteristics, obstetric history and cervical length.

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

The estimated risk is calculated by the FMF-2018 software (version 4.4) 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 is accredited by the Fetal Medicine Foundation and has submitted results for regular audit (see [www.fetalmedicine.org](http://www.fetalmedicine.org)).



#### Doppler ultrasound

Uterine artery 1.320  
PI left 1.190  
PI right 1.255  
Mean PI  
RI left 0.67  
RI right 0.62  
Notch no notch



#### Ductus Venosus

A-wave positive  
PIV 0.85

#### Cervical assessment

Cervix length 30.0 mm  
Funnelling no  
Comment normal cervical length

#### Conclusions

Diagnosis No obvious fetal defects  
normal uterine artery Doppler and cervical length

Conclusion routine scan adequate assessment

Recommendations Rescan at 18- 20 weeks to assess the fetal anatomy in detail (15th to 25th week)

#### Comments

There is single viable intrauterine pregnancy corresponding to dates. The nuchal translucency measurements, nasal bone and fetal heart rate has reduced the maternal age related risk for chromosomal abnormalities from 1 in 1076 to 1 in 8090.