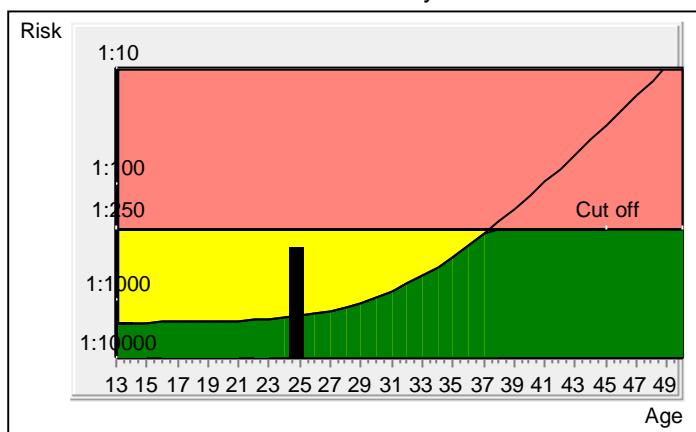


Results for:
Mrs. SNEHAL PAHILWANSample no
A0916923Date of report:
15-08-2024

Referring Doctors

Summary



Patient data

Age at delivery	24.8
WOP	19 + 4
Weight	59 kg
Patient ID	0662408110213
Ethnic origin	Asian

Risks at term

Biochemical risk for Tr.21	1:353
Age risk:	1:1390
Neural tube defects risk	<1:10000

For Mrs. SNEHAL PAHILWAN, born on 24-03-2000, a screening test was performed on the 11-08-2024. Prisca screens for Trisomy 21, Trisomy 18 and Neural tube defects (NTD).

MEASURED SERUM VALUES

	Value	Corr. MoMs
AFP	25.29 ng/mL	0.41
HCG	28211 mIU/mL	1.58
uE3	1.31 ng/mL	0.92

Gestation age 19+4

Method Scan

The MoMs have been corrected according to:
maternal weight
ethnic origin

TRISOMY 21 SCREENING

The calculated risk for Trisomy 21 is below the cut off which represents a low risk.

After the result of the Trisomy 21 test it is expected that among 353 women with the same data, there is one woman with a trisomy 21 pregnancy and 352 women with not affected pregnancies.

The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician.

Please note that risk calculations are statistical approaches and have no diagnostic value!

TRISOMY 18 SCREENING

The calculated risk for trisomy 18 is < 1:10000, which indicates a low risk.

NEURAL TUBE DEFECTS (NTD) SCREENING

The corrected MoM AFP (0.41) is located in the low risk area for neural tube defects.

Risk above
Cut off

Risk above
Age risk

Risk below
Age risk