PRISCA 5.1.0.17

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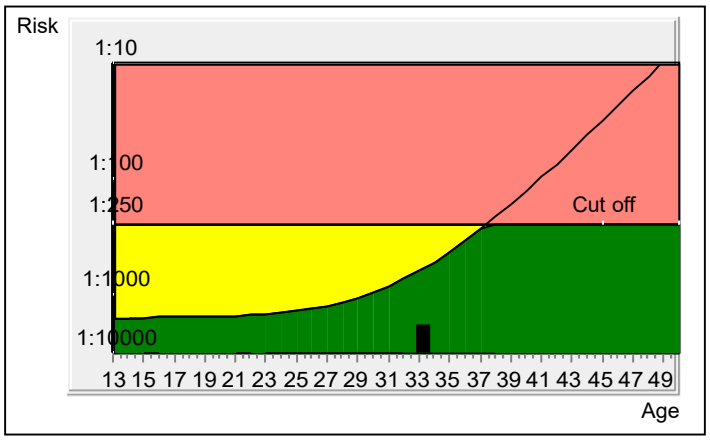
Results for:	Sample no	Date of report:
Mrs. POORNIMA VAIBHAV PATIL	NSK62210	11-03-2022

Referring Doctors	

Summary



Patient data	
Age at delivery	33.3
WOP	19 + 6
Weight	49.9 kg
Patient ID	0352203100092
Ethnic origin	Asian

Risks at term	
Biochemical risk for Tr.21	1:2541
Age risk:	1:598
Neural tube defects risk	<1:10000

For Mrs. POORNIMA VAIBHAV PATIL , born on 20-04-1989, a screening test was performed on the 10-03-2022. Prisca screens for Trisomy 21, Trisomy 18 and Neural tube defects (NTD).

MEASURED SERUM VALUES

	Value		Corr. MoMs
AFP	89.95	ng/mL	1.23
HCG	21179	mIU/mL	1.08
uE3	1.45	ng/mL	0.91

Gestation age 19+ 6
Method BPD Hadlock
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 2541 women with the same data, there is one woman with a trisomy 21 pregnancy and 2540 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 (1.23) is located in the low risk area for neural tube defects.

Risk above Cut off

Risk above Age risk

Risk below Age risk