PRISCA 5.1.0.17

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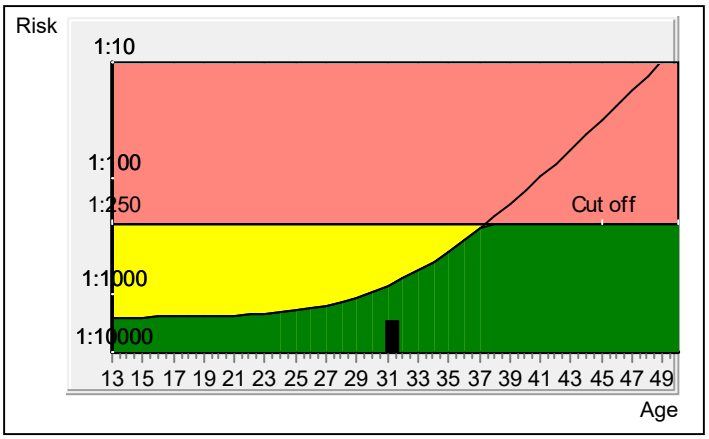
Results for:	Sample no	Date of report:
Mrs. PAYAL KEER	A0525082	16/05/24

Referring Doctors	

Summary



Patient data	
Age at delivery	31.3
WOP	15 + 1
Weight	60 kg
Patient ID	0382405140164
Ethnic origin	Asian

Risks at term	
Biochemical risk for Tr.21	1:1676
Age risk:	1:822
Neural tube defects risk	<1:10000

For Mrs. PAYAL KEER , born on 28-07-1993, a screening test was performed on the 14-05-2024. Prisca screens for Trisomy 21, Trisomy 18 and Neural tube defects (NTD).

MEASURED SERUM VALUES

	Value		Corr. MoMs
AFP	17.18	ng/mL	0.59
HCG	32644.5	mIU/mL	0.91
uE3	0.52	ng/mL	1.34
Gestation age		15+ 1	
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 1676 women with the same data, there is one woman with a trisomy 21 pregnancy and 1675 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 (0.59) is located in the low risk area for neural tube defects.

Risk above Cut off

Risk above Age risk

Risk below Age risk