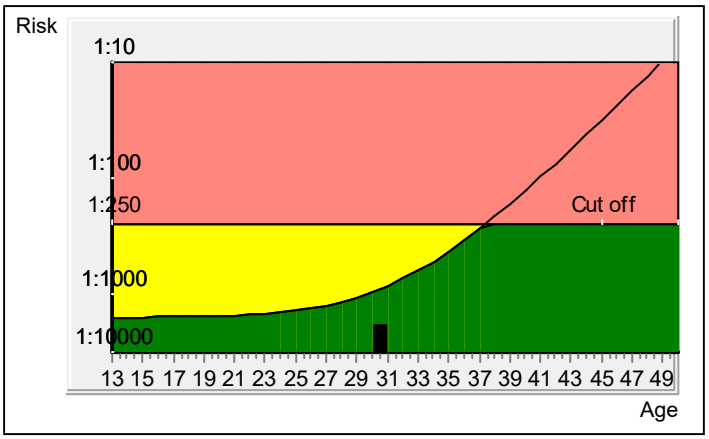
PRISCA 5.1.0.17
Licenced for: SAGEPATH LABS PVT LTD.

Results for:	Sample no	Date of report:
Mrs. FARHEEN ARFIN	A0810128	20-06-2024

Referring Doctors	

Summary



Patient data	
Age at delivery	30.6
WOP	14 + 6
Weight	57 kg
Patient ID	0482406160151
Ethnic origin	Asian

Risks at term	
Biochemical risk for Tr.21	1:3102
Age risk:	1:903
Neural tube defects risk	<1:10000

For Mrs. FARHEEN ARFIN , born on 19-05-1994, a screening test was performed on the 16-06-2024. Prisca screens for Trisomy 21, Trisomy 18 and Neural tube defects (NTD).

MEASURED SERUM VALUES

	Value		Corr. MoMs
AFP	23.3	ng/mL	0.80
HCG	33632.54	mIU/mL	0.85
uE3	0.57	ng/mL	1.62

Gestation age 14+ 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 3102 women with the same data, there is one woman with a trisomy 21 pregnancy and 3101 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 (0.80) is located in the low risk area for neural tube defects.

Risk above Cut off

Risk above Age risk

Risk below Age risk