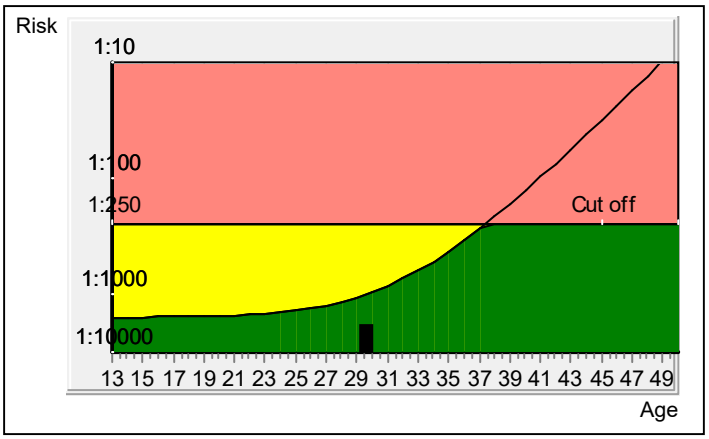
PRISCA 5.1.0.17
Licenced for: SAGEPATH LABS PVT LTD.

Results for:	Sample no	Date of report:
Mrs. FARHIN MANIYAR	A0724676	14-06-2024

Referring Doctors	

Summary



Patient data	
Age at delivery	29.6
WOP	18 + 3
Weight	63 kg
Patient ID	0662406110254
Ethnic origin	Asian

Risks at term	
Biochemical risk for Tr.21	1:2994
Age risk:	1:1011
Neural tube defects risk	<1:10000

For Mrs. FARHIN MANIYAR , born on 16-04-1995, a screening test was performed on the 11-06-2024. Prisca screens for Trisomy 21, Trisomy 18 and Neural tube defects (NTD).

MEASURED SERUM VALUES

	Value		Corr. MoMs
AFP	39.7	ng/mL	0.82
HCG	17447.83	mIU/mL	0.91
uE3	0.98	ng/mL	0.92

Gestation age 18+ 3
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 2994 women with the same data, there is one woman with a trisomy 21 pregnancy and 2993 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 (0.82) is located in the low risk area for neural tube defects.

Risk above Cut off
Risk above Age risk
Risk below Age risk