

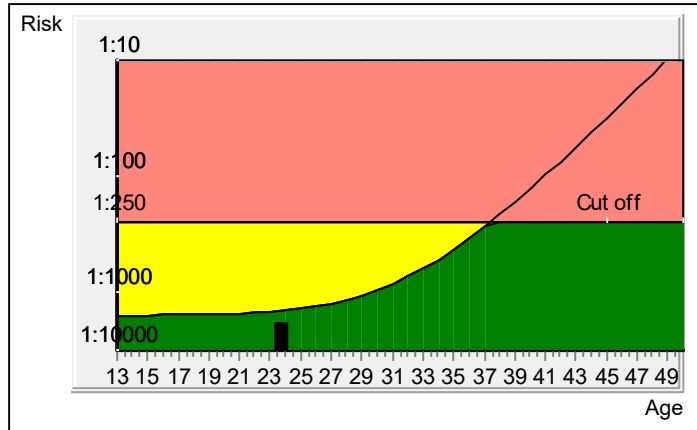
PRISCA 5.1.0.17

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Results for: Mrs. PAYAL KISHOR HOOD	Sample no 23611285	Date of report: 16/05/23
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Referring Doctors	NA
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Summary



Patient data		Risks at term	
Age at delivery	23.8	Biochemical risk for Tr.21	1:1925
WOP	18 + 0	Age risk:	1:1436
Weight	66 kg	Neural tube defects risk	<1:10000
Patient ID	0372305100097		
Ethnic origin	Asian		

For Mrs. PAYAL KISHOR HOOD, born on 02-01-2000, a screening test was performed on the 10-05-2023. Prisca screens for Trisomy 21, Trisomy 18 and Neural tube defects (NTD).

MEASURED SERUM VALUES

	Value	Corr. MoMs
AFP	23.66 ng/mL	0.54
HCG	18569.91 mIU/mL	0.93
uE3	0.88 ng/mL	0.93

Gestation age 18+ 0
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 1925 women with the same data, there is one woman with a trisomy 21 pregnancy and 1924 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 (0.54) is located in the low risk area for neural tube defects.

Risk above
Cut off

Risk above
Age risk

Risk below
Age risk