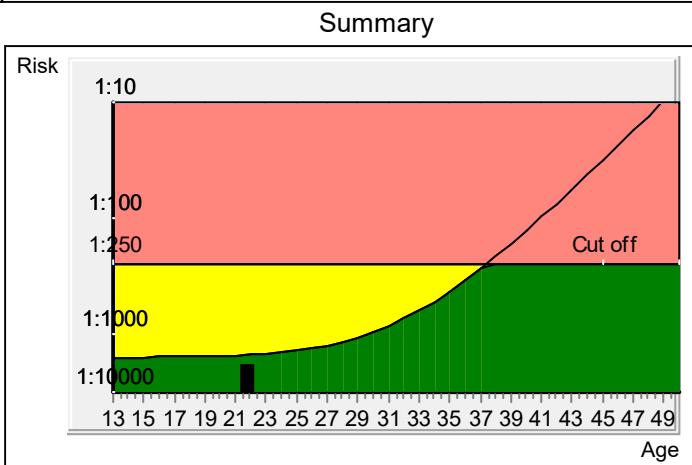


Results for:
Mrs. MANISHA RAHUL KOMBADESample no
NSK62745Date of report:
19/02/22

Referring Doctors



Patient data

Age at delivery	21.8
WOP	16 + 6
Weight	49.6 kg
Patient ID	0352202180138
Ethnic origin	Asian

Risks at term

Biochemical risk for Tr.21	1:4942
Age risk:	1:1500
Neural tube defects risk	<1:10000

For Mrs. MANISHA RAHUL KOMBADE, born on 09-10-2000, a screening test was performed on the 17-02-2022. Prisca screens for Trisomy 21, Trisomy 18 and Neural tube defects (NTD).

MEASURED SERUM VALUES

	Value	Corr. MoMs
AFP	45.22 ng/mL	0.99
HCG	31011.3 mIU/mL	1.07
uE3	1.25 ng/mL	1.68
Gestation age	16+ 6	
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 4942 women with the same data, there is one woman with a trisomy 21 pregnancy and 4941 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 (0.99) is located in the low risk area for neural tube defects.

Risk above
Cut off

Risk above
Age risk

Risk below
Age risk