

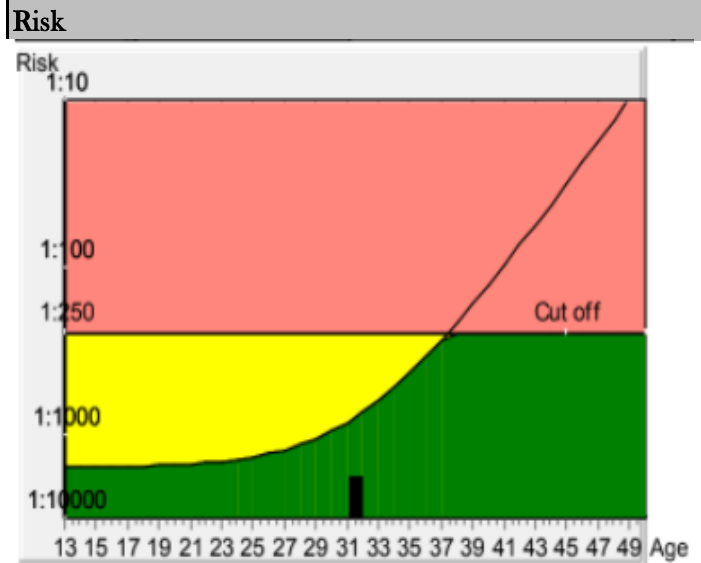
Date of Report 30-10-2020
PRISCA 5.0.2.37

Patient Data	Value		
Name	MRS MANPREET KAUR	Patient ID	012010290059
Birthday	16-08-1989	Sample ID	10787418
Age at delivery	31.6	Sample Date	26/12/2019

Correction factors			
Fetuses	1	IVF	unknown
Weight in kg	63.4	Diabetes	unknown
Smoker	Unknown	Origin	Asian
		Previous trisomy 21	unknown
		Pregnancies	unknown

Biochemical Data			Risks at sampling date	
Parameter	Value	Corr MoM		
AFP	70.2 ng/ml	1.12	Age Risk	1:786
uE3	1.85 ng/ml	1.15	Biochemical Trisomy 21 Risk	1:9041
hCG	5138.8 mIU/ml	0.29	Neural Tube Defect Risk	Below the cut off
Inhibin	274.3 IU/ml	1.2	Trisomy 18	<1:10000

Ultrasound Data		Down's Syndrome Risk (Trisomy 21 Screening)	
Gestational age	20+0	The calculated risk for Trisomy 21 is below the cut off which represents a low risk.	
Method	BPD(<>Hadlock)	After the result of the Trisomy 21 test it is expected that among more than 9041 women with the same data, there is one woman with a trisomy 21 pregnancy and 9040 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 the risk calculations are statistical approaches and have no diagnostic value!

Trisomy 18
The calculated risk for Trisomy 18 is <1:10000, which indicates a low risk
Neural Tube Defect (NTD) Screening
The corrected MoM for AFP (1.12) is located in the low risk area for neural tube defects.

The laboratory can not be held responsible for their impact on the risk assessment! Calculated value has no diagnostic value!