

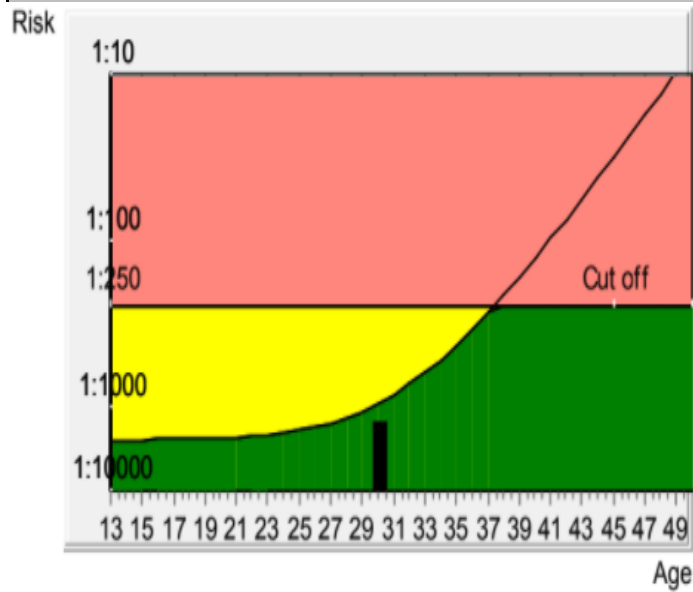
Date of Report 14-12-2020
PRISCA 5.0.2.37

Patient Data			
Name	MRS PARMEET KAUR	Patient ID	012012120163
Birthday	11-03-1991	Sample ID	10788735
Age at delivery	30.2	Sample Date	12/12/2020

Correction factors			
Fetuses	1	IVF	unknown
Weight in kg	87.3	Diabetes	unknown
Smoker	Unknown	Origin	Asian
		Previous trisomy 21	unknown
		Pregnancies	unknown

Biochemical Data			Risks at sampling date	
Parameter	Value	Corr MoM		
AFP	39.2 ng/ml	1.02	Age Risk	1:945
uE3	1.5 ng/ml	1.43	Neural Tube Defect (NTD) Screening	1:8776
hCG	36199.2 mIU/ml	1.97	Combined Trisomy 21 Risk	1:1218
			Trisomy 18	<1:10000

Ultrasound Data		Down's Syndrome Risk (Trisomy 21 Screening)	
Gestational age	18+1	The calculated risk for Trisomy 21 is below the cut off which indicates a low risk.	
Method	BPD(<>Hadlock)	After the result of the Trisomy 21 test it is expected that among 1218 women with the same data, there is one woman with a trisomy 21 pregnancy and 1217 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 (1.02) 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!

