

Date of Report 8/12/2022
PRISCA 5.1.0.17

Patient Data			
Name	MRS. KOMAL KUMARI	Patient ID	012212060199
Birthday	1/1/2000	Sample ID	11515857
Age at delivery	23.03	Sample Date	6/12/2022

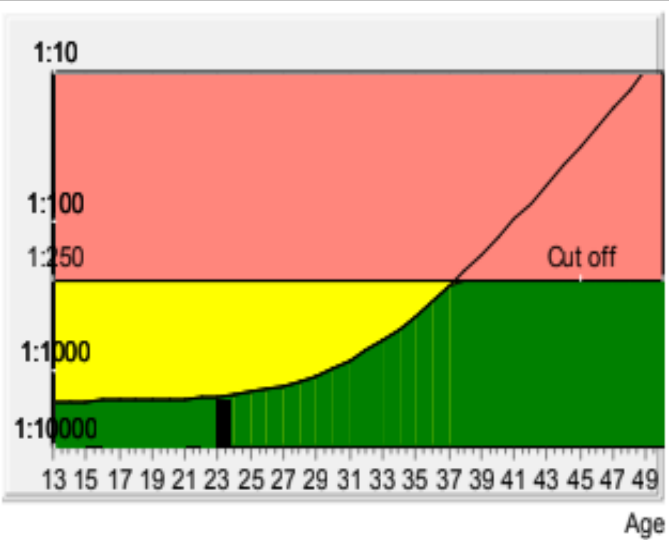
Correction factors			
Fetuses	1	IVF	unknown
Weight in kg	53	Diabetes	unknown
Smoker	Unknown	Origin	Asian
		Previous trisomy 21	unknown
		Pregnancies	unknown

Biochemical Data			Risks at sampling date	
Parameter	Value	Corr MoM		
AFP	38.5 ng/ml	0.5	Age Risk	1:1454
uE3	1.97 ng/ml	1.06	Trisomy 21 risk	1:1570
hCG	19667.2 mIU/ml	1.04	Neural tube defects risk	1:6502
			Trisomy 18	<1:10000

Ultrasound Data		Down's Syndrome Risk (Trisomy 21 Screening)	
WOP	20+4	The calculated risk for Trisomy 21 is below the cut off which indicates a low risk.	
Method	CRL (<>Robinson)	After the result of the Trisomy 21 test it is expected that among 1570 women with the same data, there is one woman with a trisomy 21 pregnancy and 1569 women with not affected pregnancies.	

Risk

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 (0.50) 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!

