

Date of Report 2/11/2021
PRISCA 5.1.0.17

Patient Data			
Name	MRS NANDITA	Patient ID	012111010092
Birthday	1/1/1996	Sample ID	11199562
Age at delivery	26.3	Sample Date	01/11/2021

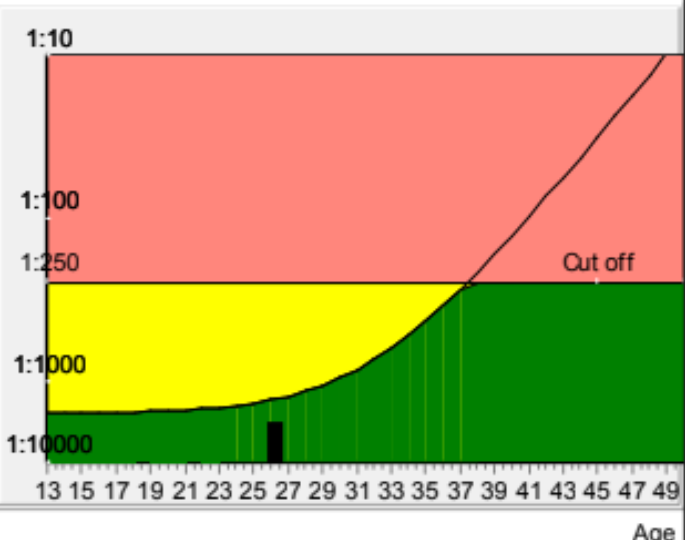
Correction factors			
Fetuses	1	IVF	unknown
Weight in kg	60.6	Diabetes	unknown
Smoker	Unknown	Origin	Asian
		Previous trisomy 21	unknown
		Pregnancies	unknown

Biochemical Data			Risks at sampling date	
Parameter	Value	Corr MoM	Age Risk	1:1302
AFP	52.2 ng/ml	1.12	Trisomy 21 risk	1:7341
uE3	1.48 ng/ml	1.42	Combined trisomy 21 risk	<1:10000
hCG	20607.8 mIU/ml	0.85	Trisomy 18	<1:10000

Ultrasound Data		Down's Syndrome Risk (Trisomy 21 Screening)	
Gestational age	17+5	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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy and 9999 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 (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!

