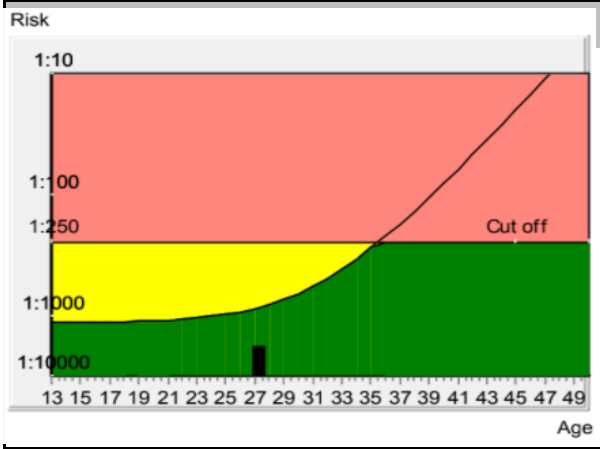
Date of Report	05-01-2021
PRISCA	5.0.2.37

Patient Data			
Name	**MRS RAJNI KUMARI**	Patient ID	012012310097
Birthday	23-09-1993	Sample ID	10855954
Age at delivery	27.3	Sample Date	31/12/2020
Gestational age			

Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	67	Diabetes	unknown	Pregnancies	unknown
Smoker	Unknown	Origin	Asian		

Biochemical Data			Ultrasound Data	
Parameter	**Value**	**Corr Mom**	Gestational age	13+2
PAPP-A	3.62 mIU/ml	0.7	Method	LMP
fb-hCG	42.2 ng/ml	1.02	Scan date	
			Crown rump length in mm	

Risks at sampling date		
Age Risk	1:865	Nuchal translucency MOM
Biochemical T21 risk	1:600	Nasal bone
Combined Trisomy 21 Risk	1:2308	Sonographer
Trisomy 13/18 + NT	<1:10000	Qualification in measuring NT



Down's Syndrome Risk (Trisomy 21 Screening)

The calculated risk for Trisomy 21 (with NT) is below the cut off, which represents a low risk.

After the result of the Trisomy 21 test (with NT) 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. 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 aapproaches and have no diagnostic value!

The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values

Trisomy 13/18 + NT

The calculated risk for Trisomy 13/18 (with NT) is <1:10000, which indicates a low risk

Risk Above Cut Off Risk above Age Risk Risk below Age risk