

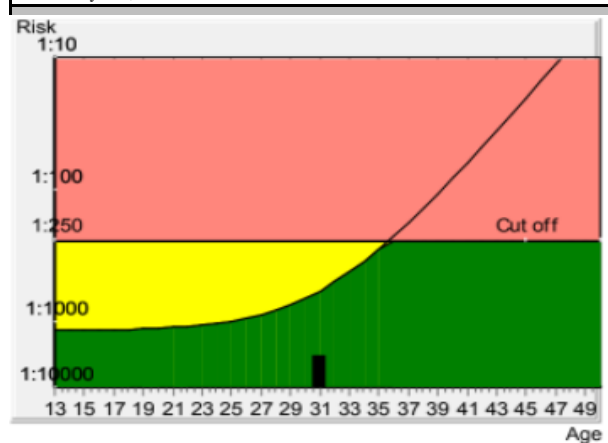
Date of Report 11-01-2021
PRISCA 5.0.2.37

Patient Data			
Name	MRS MALA KUMARI	Patient ID	012101090048
Birthday	02-02-1990	Sample ID	10845835
Age at delivery	30.9	Sample Date	09/01/2021
Gestational age	13+6		

Correction factors			
Fetuses	1	IVF	unknown
Weight in kg	57	Diabetes	unknown
Smoker	Unknown	Origin	Asian
		Previous trisomy 21	unknown
		Pregnancies	unknown

Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	13+3
PAPP-A	5.3 mIU/ml	0.71	Method	CRL(<>Robinson
fb-hCG	62.4 ng/ml	1.53	Scan date	06-01-2021

Risks at sampling date			Ultrasound Data	
Age Risk	1:597		Crown rump length in mm	72.8
Biochemical T21 risk	1:635		Nuchal translucency MOM	0.83
Combined Trisomy 21 Risk	1:3587		Nasal bone	Present
Trisomy 13/18 + NT	<1:10000		Sonographer	DR.NIDHI
			Qualification in measuring NT	MBBS



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 3587 women with the same data, there is one woman with a trisomy 21 pregnancy and 3586 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 13/18 + NT

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

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

Risk Above Cut Off

Risk above Age Risk

Risk below Age risk