

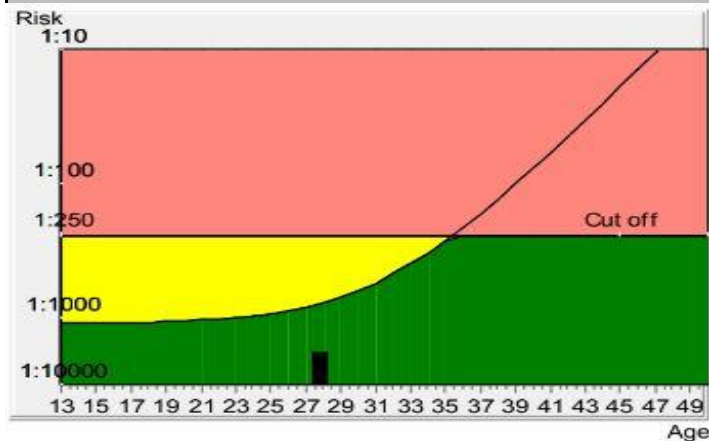
Date of Report 25/12/19
PRISCA 5.0.2.37

Patient Data			
Name	Mrs Shweta Maheshwari	Patient ID	051912240034
Birth day	4/3/1992	Sample ID	10501627
Age at delivery	27.8	Sample Date	24/12/19
Gestational age	12+0		

Correction factors			
Fetuses	1	IVF	unknown
Weight in kg	40.5	Diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21	unknown
		Pregnancies	

Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	12+0
PAPP-A	3.77 mIU/ml	0.65	Method	CRL (<>Robinson)
fb-hCG	102.5 ng/ml	1.79	Scan Date	24/12/19
Risks at sampling date			Crown Rump Length (mm)	52.9
Age Risk		1:793	Nuchal translucency MoM	0.54
Biochemical Trisomy 21 Risk		1:476	Nasal Bone	present
Combined Trisomy 21 Risk		1:2866	Sonographer	DR APRAJITA MEHTA
Trisomy 13/18 + NT		<1:10000	Qualification in measuring NT	C/R

Risk **Down's Syndrome Risk (Trisomy 21 Screening)**



The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.

After the result of the Trisomy 21 test (with NT) it is expected that among 2866 women with the same data, there is one woman with a trisomy 21 pregnancy and 2865 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 nuchal translucency) is <1:10000, which represents 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

