

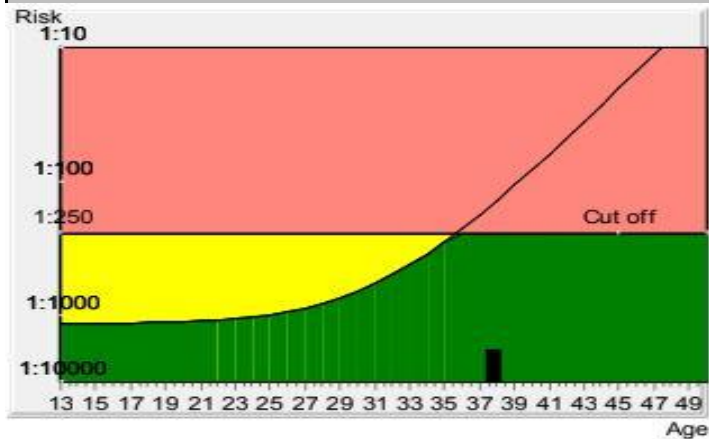
Date of Report 24/07/2019
PRISCA 5.0.2.37

Patient Data			
Name	Mrs Arti Gaur	Patient ID	011907230055
Birthday	20/10/1981	Sample ID	10598171
Age at delivery	37.8	Sample Date	23/07/19
Gestational age	13+5		

Correction factors			
Fetuses	1	IVF	unknown
Weight in kg	61	Diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21	unknown
		Pregnancies	

Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	13+4
PAPP-A	4.61 mIU/ml	0.7	Method	CRL (<>Robinson)
fb-hCG	42.36 ng/ml	1.05	Scan Date	22/07/19
Risks at sampling date			Crown Rump Length (mm)	74.9
Age Risk		1:148	Nuchal translucency MoM	0.81
Biochemical Trisomy 21 Risk		1:368	Nasal Bone	present
Combined Trisomy 21 Risk		1:2015	Sonographer	DR.RUBY RAHUL
Trisomy 13/18 + NT		<1:10000	Qualification in measuring NT	CON. RADIOLOGIT

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 2015 women with the same data, there is one woman with a trisomy 21 pregnancy and 2014 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

