

Date of Report 22/08/19
PRISCA 5.0.2.37

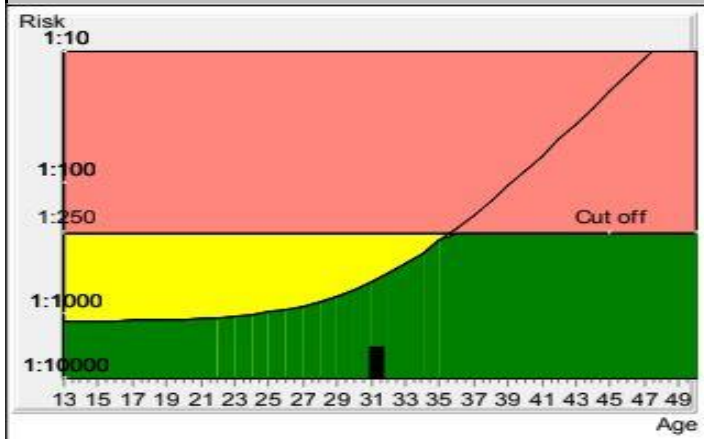
Patient Data			
Name	Mrs Swati Agarwal	Patient ID	011908210068
Birth day	12/5/1988	Sample ID	10550970
Age at delivery	31.3	Sample Date	21/08/19
Gestational age	13+4		

Correction factors			
Fetuses	1	IVF	unknown
Weight in kg	87	Diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21	unknown
		Pregnancies	

Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	13+0
PAPP-A	3.85 mIU/ml	0.95	Method	CRL (<>Robinson)
fb-hCG	48.21 ng/ml	1.30	Scan Date	17/08/2019

Risks at sampling date		Ultrasound Data	
Age Risk	1:563	Crown Rump Length (mm)	63.6
Biochemical Trisomy 21 Risk	1:1708	Nuchal translucency MoM	0.73
Combined Trisomy 21 Risk	1:8949	Nasal Bone	present
Trisomy 13/18 + NT	<1:10000	Sonographer	
		Qualification in measuring NT	

Risk	Down's Syndrome Risk (Trisomy 21 Screening)
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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 8949 women with the same data, there is one woman with a trisomy 21 pregnancy and 8948 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

