

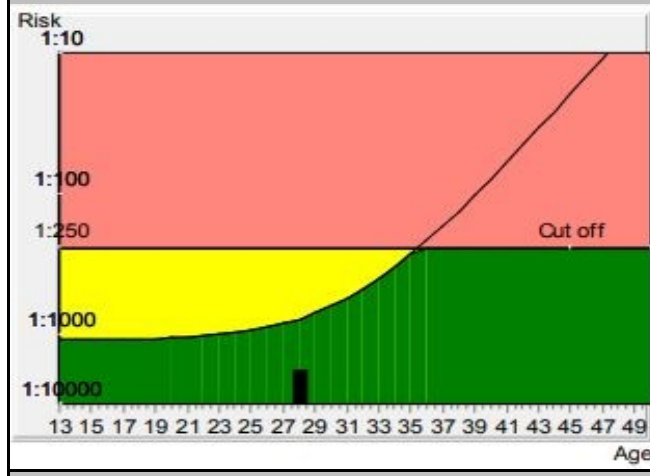
Date of Report 11-01-19
PRISCA 5.0.2.37

Patient Data			
Name	Mrs Binni	Patient ID	011901090194
Birthday	30-11-90	Sample ID	10352151
Age at delivery	28.1	Sample Date	09/01/2019
Gestational age	12+2		

Correction factors			
Fetuses	1	IVF	unknown
Weight in kg	57.1	Diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21	unknown
		Pregnancies	

Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	12+1
PAPP-A	1.77 mIU/ml	0.48	Method	CRL (< Hadlock)
fb-hCG	28.87 ng/ml	0.62	Scan Date	09-01-19

Risks at sampling date			CRL	
Age Risk		1:781	CRL	54.8
Biochemical T21 Risk		1:2227	Nuchal translucency MoM	0.48
Combined Trisomy 21 Risk		<1:10000	Nasal Bone	Present
Trisomy 13/18 + NT		<1:10000	Sonographer	DR.SAHIL LOOMBA
			Qualifications in measuring NT	MBBS,DNB

Risk	Down's Syndrome Risk (Trisomy 21 Screening)
	<p>The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.</p> <p>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</p> <p>The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician. Please note that risk calculations are statistical approach and have no diagnostic values!</p> <p>The patient combined risk presumes the NT measurement was done according to accepted guidelines</p>

Trisomy 13/18 + NT
The calculated risk for trisomy 13/18 (with nuchal translucency) is 1:8658, which represents a low risk.