

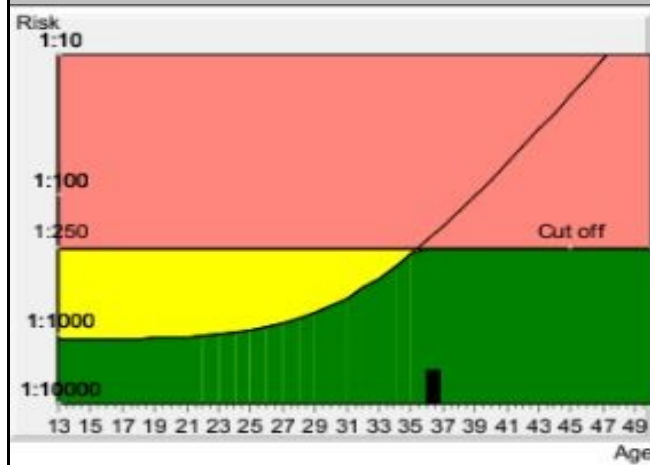
Date of Report 07-01-19
PRISCA 5.0.2.37

Patient Data			
Name	Dr. Sheenam	Patient ID	051901060001
Birthday	20-08-82	Sample ID	10344101
Age at delivery	36.4	Sample Date	06/01/2019
Gestational age	12+1		

Correction factors			
Fetuses	1	IVF	unknown
Weight in kg	76.1	Diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21	unknown
		Pregnancies	

Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom		
PAPP-A	1.43 mIU/ml	0.57	Gestational age	12+0
fb-hCG	23.7 ng/ml	0.54	Method	CRL (\leqHadlock)
			Scan Date	05-01-19

Risks at sampling date			CRL	
Age Risk		1:195		55.5
Biochemical T21 Risk		1:1182	Nuchal translucency MoM	1.09
Combined Trisomy 21 Risk		1:3984	Nasal Bone	Present
Trisomy 13/18 + NT		1:8658	Sonographer	DR.SHREYASI SHARMA
			Qualifications in measuring NT	MBBS,MD

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 3984 women with the same data, there is one woman with a trisomy 21 pregnancy and 3983 women with not affected pregnancies.</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.