

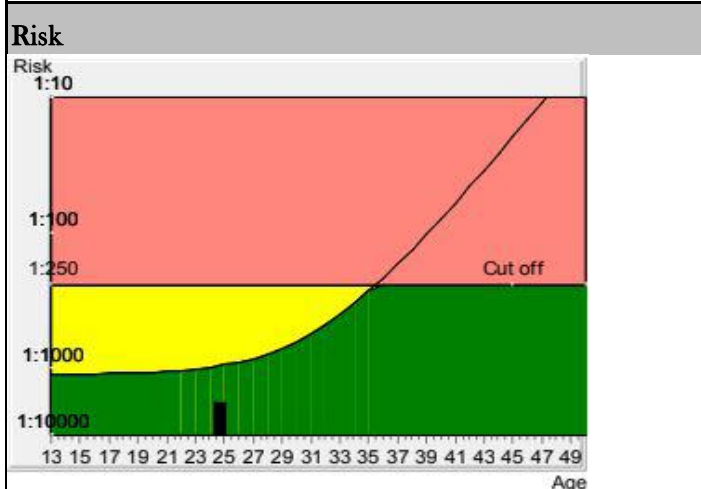
Date of Report
PRISCA

28/06/2019
5.0.2.37

Patient Data			
Name	Mrs. Shilpi	Patient ID	011906270061
Birthday	8/11/1994	Sample ID	10362227
Age at delivery	24.6	Sample Date	27/06/2019
Gestational age	12+4		

Correction factors			
Fetuses	F1 IVF	unknown	Previous trisomy 21
Weight in kg	73 Diabetes	unknown	Pregnancies
Smoker	Unknown Origin	Asian	

Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	12+2
PAPP-A	4.36 mIU/ml	0.65	Method	CRL (<>Robinson)
fb-hCG	141.5 ng/ml	1.5	Scan Date	25/06/2019
Risks at sampling date			Crown rump length in mm	54.9
Age Risk	1:978		Nuchal translucency MoM	1.03
Overall population risk	1:886		Nasal bone	Present
Combined Trisomy 21 Risk	1:3489		Sonographer	DR.SANJEEV KUMAR SINGHAL
Trisomy 13/18 + NT	<1:10000		Qualifications in measuring NT	MBBS,PGDUS,DMRD



Down's Syndrome Risk (Trisomy 21 Screening)

The calculated risk for Trisomy 21 (with NT) is below the cut off, which represents a low risk.

After the result of the Trisomy 21 test (with NT) it is expected that among more than 3489 women with the same data, there is one woman with a trisomy 21 pregnancy and 3488 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 NT) is <1:10000, which indicates 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