

Date of Report 23/04/19
PRISCA 5.0.2.37

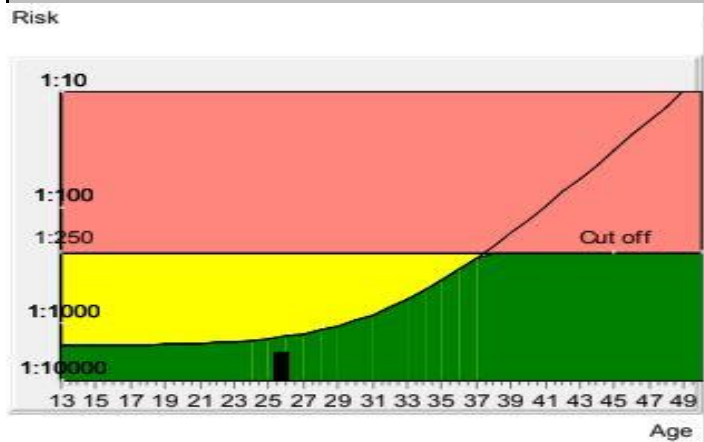
Patient Data			
Name	Mrs Sangeeta	Patient ID	011904210174
Birthday	12/1/1994	Sample ID	10421664
Age at delivery	25.8	Sample Date	21-04-19
Gestational age	14+1		

Correction factors			
Fetuses	1	IVF	unknown
Weight in kg	96	Diabetes	unknown
Smoker	Unknown	Origin	Asian
		Previous trisomy 21	unknown
		Pregnancies	unknown

Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	14+0
AFP	22.01 ng/ml	1.07	Method	CRL (<>Robinson)
uE3	0.37 mIU/ml	1.80	Scan Date	21/04/19
hCG	80171.5 ng/ml	2.41	Crown rump length (mm)	80.0 mm

Risks at sampling date			
Age Risk	1:1334	Nuchal translucency	1.3
Biochemical T21 Risk	1:1198	Nuchal translucency MoM	0.67
Combined Trisomy 21 Risk	1:7326	Nasal Bone	Present
Trisomy 13/18 + NT	<1:10000	Sonographer	Dr Nikhil Sharma
		Qualifications in measuring NT	MD

Risk **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 7326 women with the same data, there is one woman with a trisomy 21 pregnancy and 7325 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