

Date of Report 19/04/19
PRISCA 5.0.2.37

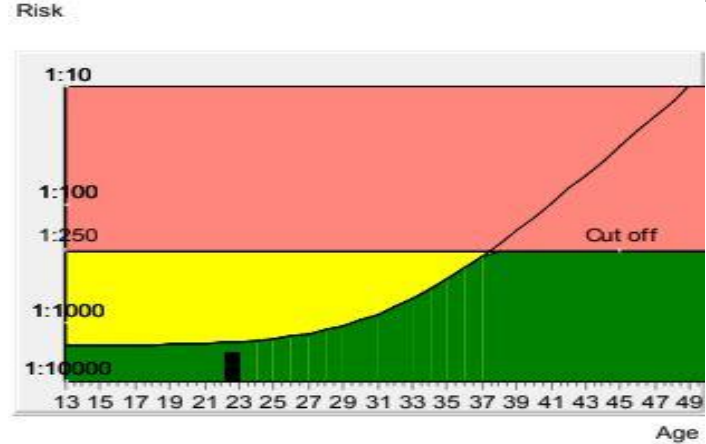
Patient Data			
Name	Mrs Anjali	Patient ID	011904170164
Birth day	12/2/1997	Sample ID	10362338
Age at delivery	22.7	Sample Date	17-04-19
Gestational age	14+4		

Correction factors			
Fetuses	1	IVF	unknown
Weight in kg	61	Diabetes	unknown
Smoker	Unknown	Origin	Asian
		Previous trisomy 21	unknown
		Pregnancies	unknown

Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	12+4
AFP	24.1 ng/ml	0.8	Method	CRL (<>Robinson)
uE3	0.36 mIU/ml	1.16	Scan Date	
hCG	35017.5 ng/ml	0.88	Crown rump length (mm)	

Risks at sampling date			
Age Risk	1:1476	Nuchal translucency	1.2
Biochemical T21 Risk	1:5136	Nuchal translucency MoM	1
Combined Trisomy 21 Risk	<1:10000	Nasal Bone	Present
Trisomy 13/18 + NT	<1:10000	Sonographer	DR.SANJEEV KUMAR SINGHAL
		Qualifications in measuring NT	MBBS,PGDUS,DMRD

Risk	Down's Syndrome Risk (Trisomy 21 Screening)
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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 10000 women with the same data, there is one woman with a trisomy 21 pregnancy.

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