

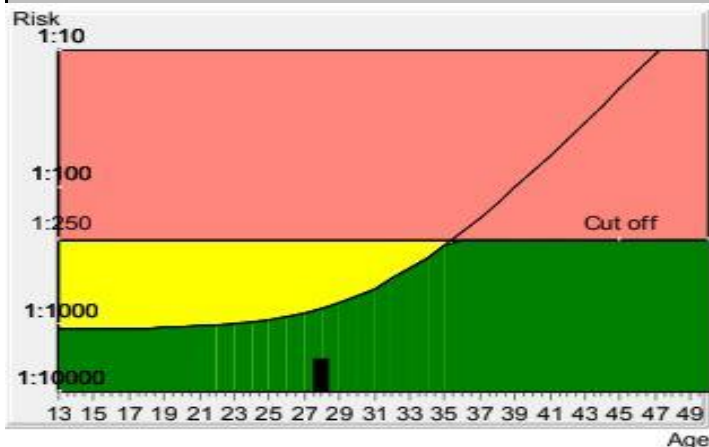
Date of Report 21/11/19
PRISCA 5.0.2.37

Patient Data	
Name	Mrs Rashmi
Birth day	3/1/1992
Age at delivery	27.9
Gestational age	12+0
Patient ID	011911200191
Sample ID	10362045
Sample Date	20/11/19

Correction factors	
Fetuses	1 IVF
Weight in kg	56 Diabetes
Smoker	no Origin
unknown	Previous trisomy 21
no	Pregnancies
Asian	

Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	12+0
PAPP-A	1.73 mIU/ml	0.43	Method	CRL (<>Robinson)
fb-hCG	24.68 ng/ml	0.49	Scan Date	20/11/19
Risks at sampling date			Crown Rump Length (mm)	47.2
Age Risk	1:788		Nuchal translucency MoM	0.93
Biochemical Trisomy 21 Risk	1:2779		Nasal Bone	present
Combined Trisomy 21 Risk	<1:10000		Sonographer	DR. SANJEEV SINGHAL
Trisomy 13/18 + NT	<1:10000		Qualification in measuring NT	MBBS, DMRD

Risk **Down's Syndrome Risk (Trisomy 21 Screening)**



The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates 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 nuchal translucency) is <1:10000, which represents a low risk.

The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values

