

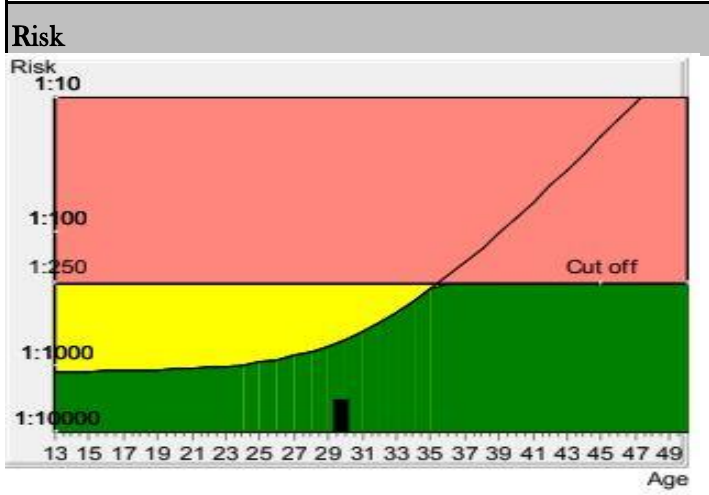
Date of Report 18/09/19
 PRISCA 5.0.2.37

Patient Data	
Name	Mrs Heena
Birth day	15/12/89
Age at delivery	29.8
Gestational age	12+3
Patient ID	011909170080
Sample ID	10556833
Sample Date	18/09/19

Correction factors	
Fetuses	1 IVF
Weight in kg	73.5 Diabetes
Smoker	no Origin
unknown	Previous trisomy 21
no	Pregnancies
Asian	

Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	11+1
PAPP-A	4.25 mIU/ml	1.25	Method	CRL (<>Robinson)
fb-hCG	22.39 ng/ml	0.51	Scan Date	8/9/2019

Risks at sampling date		Down's Syndrome Risk (Trisomy 21 Screening)	
Age Risk	1:663	Crown Rump Length (mm)	42.2
Biochemical Trisomy 21 Risk	<1:10000	Nuchal translucency MoM	0.85
Combined Trisomy 21 Risk	<1:10000	Nasal Bone	present
Trisomy 13/18 + NT	<1:10000	Sonographer	Dr. J.P. Sharma
		Qualification in measuring NT	C/R



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 held responsible for their impact on the risk assessment! Calculated risks have no diagnostic values

