

Date of Report 5/5/2019
 PRISCA 5.0.2.37

Patient Data	
Name Mrs Renu	Patient ID 011905030219
Birthday 20/08/1991	Sample ID 10303265
Age at delivery 27.7	Sample Date 03/05/2019
Gestational age 13+0	

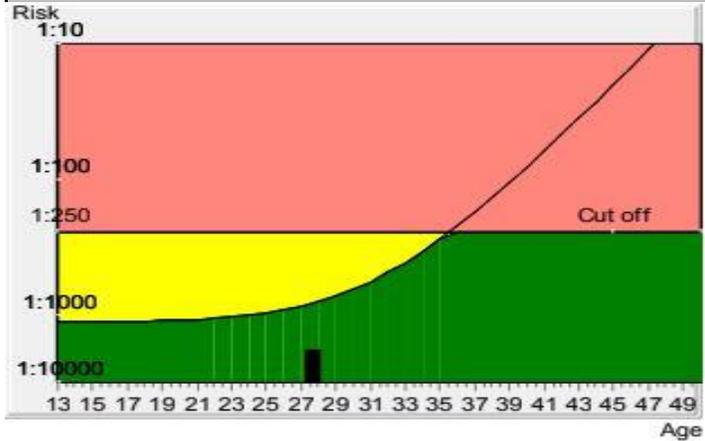
Correction factors			
Fetuses	1	IVF	unknown
Weight in kg	57	Diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21	unknown
		Pregnancies	unknown

Biochemical Data	Ultrasound Data
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Parameter	Value	Corr Mom	Ultrasound Data
PAPP-A	4.26 mIU/ml	0.75	Gestational age 12+0
fb-hCG	58.1 ng/ml	1.29	Method CRL (<>Robinson)
			Scan Date 27/04/2019

Risks at sampling date	Ultrasound Data
Age Risk 1:829	Crown Rump Length (mm) 52
Biochemical Trisomy 21 Risk 1:1513	Nuchal translucency MoM 0.93
Combined Trisomy 21 Risk 1:7217	Nasal Bone present
Trisomy 13/18 + NT <1:10000	Sonographer DR.KRITI RAJ
	Qualification in measuring NT MBBS,MD

Risk	Down's Syndrome Risk (Trisomy 21 Screening)
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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 7217 women with the same data, there is one woman with a trisomy 21 pregnancy and 7216 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	Down's Syndrome Risk (Trisomy 21 Screening)
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

