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		Date of Report	5/9/2019
		PRISCA	5.0.2.37

Patient Data			
Name	Mrs Purnima Dua	Patient ID	011909040084
Birthday	19/06/1987	Sample ID	10573671
Age at delivery	32.2	Sample Date	04/09/19
Gestational age	13+6		

Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	84	Diabetes	no	Pregnancies	
Smoker	no	Origin	Asian		

Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	13+5
PAPP-A	3.56 mIU/ml	0.76	Method	CRL (<>Robinson)
fb-hCG	29.8 ng/ml	0.82	Scan Date	3/9/2019
Risks at sampling date			Crown Rump Length (mm)	78
Age Risk		1:490	Nuchal translucency MoM	0.75
Biochemical Trisomy 21 Risk		1:2575	Nasal Bone	present
Combined Trisomy 21 Risk		<1:10000	Sonographer	DR. PAULDEEP KAUR
Trisomy 13/18 + NT		<1:10000	Qualification in measuring NT	MBBS, MD

Risk	Down's Syndrome Risk (Trisomy 21 Screening)
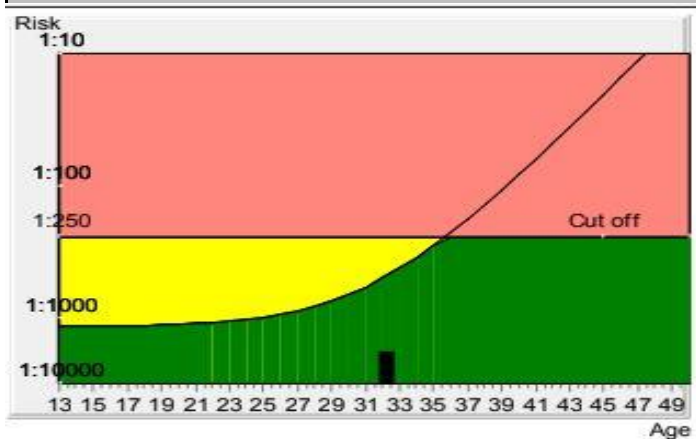
TThe calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.

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 aapproaches 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

	Risk Above Cut Off		Risk above Age Risk		Risk below Age risk