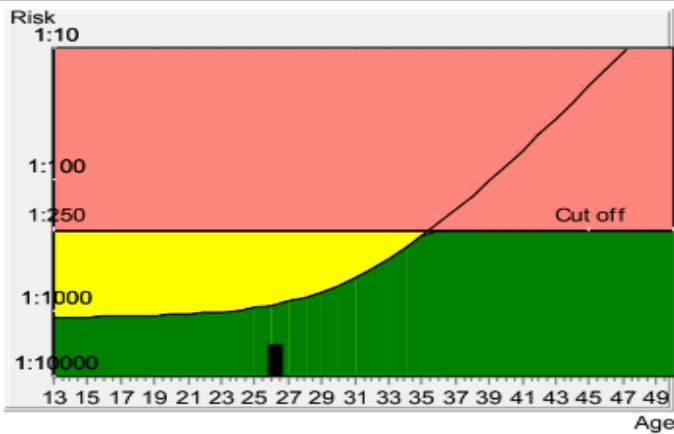


Date of Report 28/10/19  
PRISCA 5.0.2.37

Patient Data				
Name	Mrs Priyanka	Patient ID	011910260024	
Birthday	12/8/1993	Sample ID	10551494	
Age at delivery	26.2	Sample Date	26/10/19	
Gestational age	12+3			
Correction factors				
Fetuses	1 IVF	unknown	Previous trisomy 21	unknown
Weight in kg	57 Diabetes	no	Pregnancies	
Smoker	no Origin	Asian		
Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	11+5
PAPP-A	5.12 mIU/ml	0.6	Method	CRL (<>Robinson)
fb-hCG	131.2 ng/ml	1.26	Scan Date	21/10/19
Risks at sampling date			Crown Rump Length (mm)	50.6
Age Risk	1:901		Nuchal translucency MoM	0.37
Biochemical T21 Risk	1:989		Nasal Bone	Present
Combined Trisomy 21 Risk	1:5815		Sonographer	Dr. Parul
Trisomy 13/18 + NT	<1:10000		Qualification in measuring NT	C/R
Risk			Down's Syndrome Risk (Trisomy 21 Screening)	
			<p>The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.</p> <p>After the result of the Trisomy 21 test (with NT) it is expected that among 5815 women with the same data, there is one woman with a trisomy 21 pregnancy and 5814 women with not affected pregnancies.</p> <p>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!</p>	
Trisomy 13/18 + NT			The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values	
The calculated risk for trisomy 13/18 is <1:10000, which represents a low risk.				

 Risk Above Cut Off

 Risk above Age Risk

 Risk below Age risk

