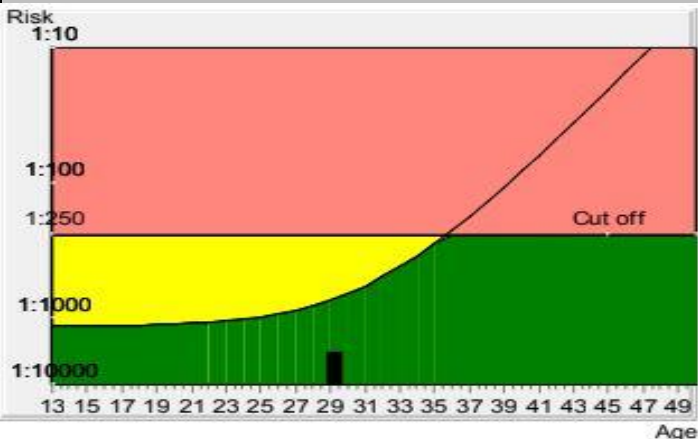


Date of Report 9/9/2019  
PRISCA 5.0.2.37

Patient Data				
Name	Mrs Amrita	Patient ID	011909070322	
Birthday	9/7/1990	Sample ID	10566672	
Age at delivery	29.2	Sample Date	07/09/19	
Gestational age	13+6			
Correction factors				
Fetuses	1	IVF	unknown	Previous trisomy 21 unknown
Weight in kg	53	Diabetes	no	Pregnancies
Smoker	no	Origin	Asian	
Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	13+5
PAPP-A	5.4 mIU/ml	0.66	Method	CRL (<>Robinson)
fb-hCG	39.2 ng/ml	0.94	Scan Date	7/9/2019
Risks at sampling date			Crown Rump Length (mm)	77.7
Age Risk		1:743	Nuchal translucency MoM	0.69
Biochemical Trisomy 21 Risk		1:2084	Nasal Bone	present
Combined Trisomy 21 Risk		<1:10000	Sonographer	Dr. Subhansish Mohapatra
Trisomy 13/18 + NT		<1:10000	Qualification in measuring NT	MBBS, MD
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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy.</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 (with nuchal translucency) is < 1:10000, which represents a low risk.				

 Risk Above Cut Off

 Risk above Age Risk

 Risk below Age risk

