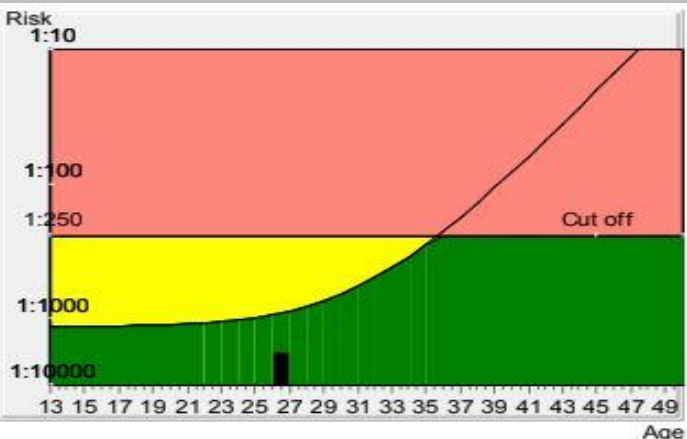


Date of Report 22/10/19
PRISCA 5.0.2.37

Patient Data				
Name	Mrs Shilpa Guria	Patient ID	011910200290	
Birthday	3/4/1993	Sample ID	10494660	
Age at delivery	26.5	Sample Date	20/10/19	
Gestational age	13+5			
Correction factors				
Fetuses	1	IVF	unknown	Previous trisomy 21
Weight in kg	56	Diabetes	no	Pregnancies
Smoker	no	Origin	Asian	
Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	12+6
PAPP-A	4.68 mIU/ml	0.64	Method	CRL (<>Robinson)
fb-hCG	92.1 ng/ml	2.21	Scan Date	15/10/19
Risks at sampling date			Crown Rump Length (mm)	64.4
Age Risk		1:921	Nuchal translucency MoM	0.73
Biochemical Trisomy 21 Risk		1:310	Nasal Bone	present
Combined Trisomy 21 Risk		1:1930	Sonographer	DR. ELLORA
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 1930 women with the same data, there is one woman with a trisomy 21 pregnancy and 1929 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 (with nuchal translucency) is <1:10000, which represents a low risk.				

 Risk Above Cut Off

 Risk above Age Risk

 Risk below Age risk

