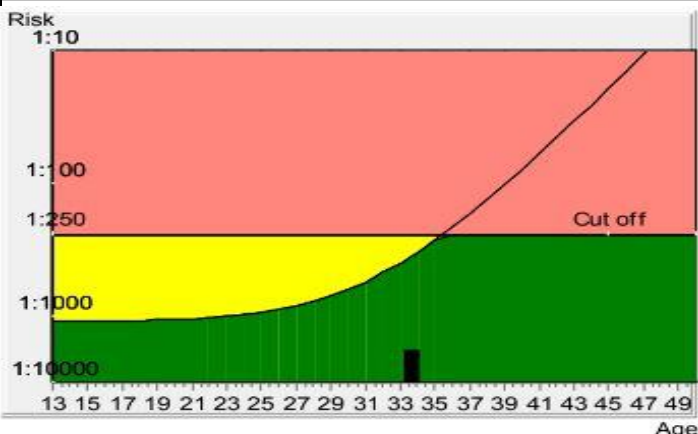


Date of Report 23/12/19  
PRISCA 5.0.2.37

Patient Data				
Name	Mrs Deepali	Patient ID	011912210212	
Birthday	19/04/1986	Sample ID	10303055	
Age at delivery	33.7	Sample Date	21/12/19	
Gestational age	12+1			
Correction factors				
Fetuses	1	IVF	unknown	Previous trisomy 21
Weight in kg	60	Diabetes	no	Pregnancies
Smoker	no	Origin	Asian	
Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	12+0
PAPP-A	2.16 mIU/ml	0.56	Method	CRL (<>Robinson)
fb-hCG	53.4 ng/ml	1.10	Scan Date	21/12/19
Risks at sampling date			Crown Rump Length (mm)	53.7
Age Risk		1:353	Nuchal translucency MoM	0.81
Biochemical Trisomy 21 Risk		1:445	Nasal Bone	present
Combined Trisomy 21 Risk		1:2593	Sonographer	DR RAJESH ARORA
Trisomy 13/18 + NT		<1:10000	Qualification in measuring NT	HMC
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 2593 women with the same data, there is one woman with a trisomy 21 pregnancy and 2592 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

