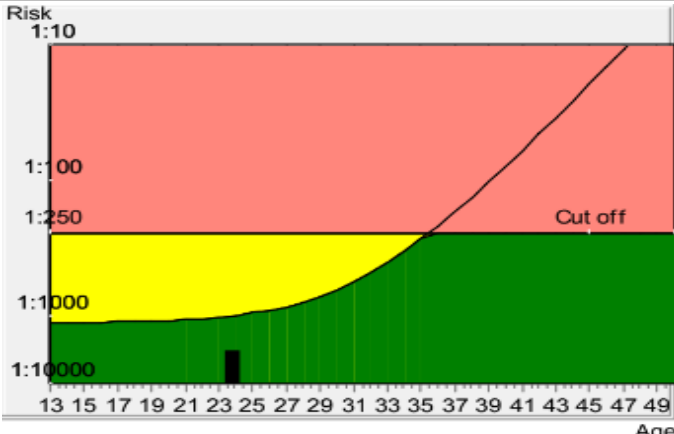


Date of Report 1/11/2019  
 PRISCA 5.0.2.37

Patient Data				
Name	Mrs Kavita	Patient ID	011910310179	
Birthday	1/1/1996	Sample ID	10579907	
Age at delivery	23.8	Sample Date	31/10/19	
Gestational age	12+4			
Correction factors				
Fetuses	1 IVF	unknown	Previous trisomy 21	unknown
Weight in kg	66 Diabetes	no	Pregnancies	
Smoker	no Origin	Asian		
Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	12+4
PAPP-A	3.5 mIU/ml	0.86	Method	CRL (<>Robinson)
fb-hCG	49.62 ng/ml	1.10	Scan Date	31/10/19
Risks at sampling date			Crown Rump Length (mm)	
Age Risk	1:1007		60.2	
Biochemical T21 Risk	1:3582		Nuchal translucency MoM	
Combined Trisomy 21 Risk	<1:10000		0.82	
Trisomy 13/18 + NT	<1:10000		Nasal Bone	
			Present	
			Sonographer	
			Dr. ANKIT BHARGAVA	
			Qualification in measuring NT	
			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. 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

