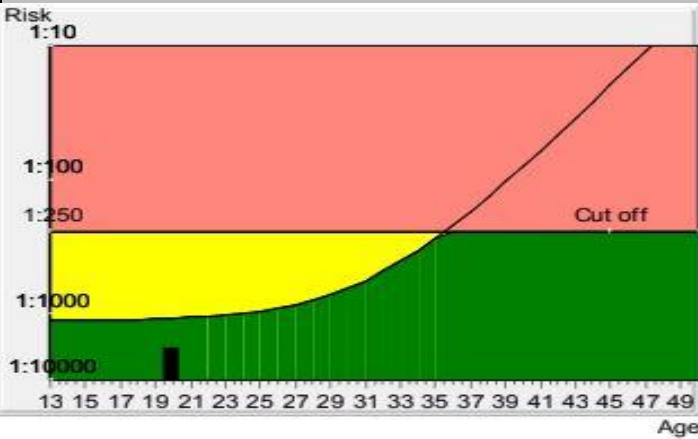


Date of Report 6/8/2019
PRISCA 5.0.2.37

Patient Data				
Name	Mrs Pinki	Patient ID	011908050143	
Birthday	1/9/1999	Sample ID	10593875	
Age at delivery	19.9	Sample Date	05/08/19	
Gestational age	12+6			
Correction factors				
Fetuses	1	IVF	unknown	Previous trisomy 21
Weight in kg	45	Diabetes	no	Pregnancies
Smoker	no	Origin	Asian	
Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	12+6
PAPP-A	3.14 mIU/ml	0.44	Method	CRL (<>Robinson)
fb-hCG	25.98 ng/ml	0.52	Scan Date	5/8/2019
Risks at sampling date			Crown Rump Length (mm)	64.8
Age Risk	1:1102		Nuchal translucency MoM	0.66
Biochemical Trisomy 21 Risk	1:3626		Nasal Bone	present
Combined Trisomy 21 Risk	<1:10000		Sonographer	DR.RAHUL
Trisomy 13/18 + NT	<1:10000		Qualification in measuring NT	DNB
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

