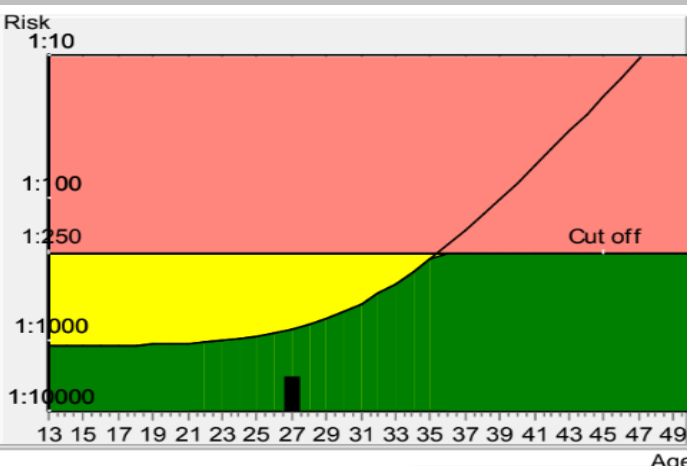


Date of Report 19/01/2020  
PRISCA 5.0.2.37

Patient Data				
Name	Mrs Bindu	Patient ID	012001180122	
Birthday	2/1/1993	Sample ID	10518972	
Age at delivery	27	Sample Date	18/01/2020	
Gestational age	12+1			
Correction factors				
Fetuses	1	IVF	unknown	Previous trisomy 21
Weight in kg	51.6	Diabetes	no	Pregnancies
Smoker	no	Origin	Asian	
Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	12+1
PAPP-A	4.3 mIU/ml	0.93	Method	CRL (<>Robinson)
fb-hCG	89.7 ng/ml	1.75	Scan Date	18/01/2020
Risks at sampling date			Crown Rump Length (mm)	55.9
Age Risk		1:845	Nuchal translucency MoM	0.68
Biochemical Trisomy 21 Risk		1:1199	Nasal Bone	present
Combined Trisomy 21 Risk		1:6579	Sonographer	DR. ARSHBIR SINGH
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 6579 women with the same data, there is one woman with a trisomy 21 pregnancy and 6578 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