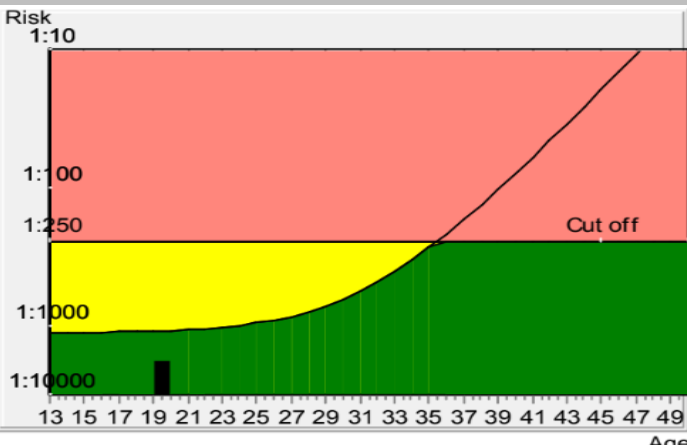


Date of Report 24/02/2020
PRISCA 5.0.2.37

Patient Data				
Name	Mrs Anju	Patient ID	012002220183	
Birthday	20/08/2000	Sample ID	10658857	
Age at delivery	19.5	Sample Date	22/02/2020	
Gestational age	12+4			
Correction factors				
Fetuses	1	IVF	unknown	Previous trisomy 21
Weight in kg	38	Diabetes	no	Pregnancies
Smoker	no	Origin	Asian	
Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	11+2
PAPP-A	4.36 mIU/ml	0.57	Method	CRL (<>Robinson)
fb-hCG	64.2 ng/ml	1.16	Scan Date	14/02/2020
Risks at sampling date			Crown Rump Length (mm)	45.7
Age Risk		1:1096	Nuchal translucency MoM	0.88
Biochemical Trisomy 21 Risk		1:1267	Nasal Bone	present
Combined Trisomy 21 Risk		1:6989	Sonographer	DR SHELJA JAIN
Trisomy 13/18 + NT		<1:10000	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 6989 women with the same data, there is one woman with a trisomy 21 pregnancy and 6988 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