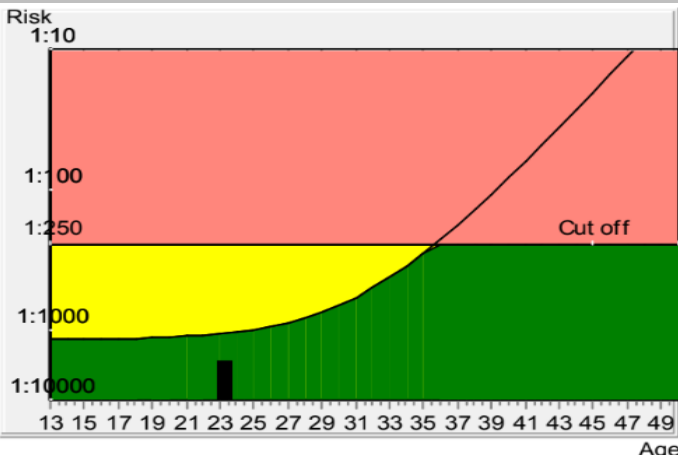


Date of Report 18/02/2020
PRISCA 5.0.2.37

Patient Data				
Name	Mrs Priya w/o Krishan	Patient ID	012002160190	
Birthday	24/10/1996	Sample ID	10370311	
Age at delivery	23.3	Sample Date	16/02/2020	
Gestational age	13+6			
Correction factors				
Fetuses	1 IVF	unknown	Previous trisomy 21	unknown
Weight in kg	46.9	Diabetes	no	Pregnancies
Smoker	no	Origin	Asian	
Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	13+4
PAPP-A	5.12 mIU/ml	0.55	Method	CRL (<>Robinson)
fb-hCG	53.8 ng/ml	1.23	Scan Date	15/02/2020
Risks at sampling date			Crown Rump Length (mm)	75.4
Age Risk	1:1066		Nuchal translucency MoM	1.30
Biochemical Trisomy 21 Risk	1:995		Nasal Bone	present
Combined Trisomy 21 Risk	1:1667		Sonographer	DR. RAKHI BANSAL
Trisomy 13/18 + NT	<1:10000		Qualification in measuring NT	MBBS, DMRD
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 1667 women with the same data, there is one woman with a trisomy 21 pregnancy and 1666 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