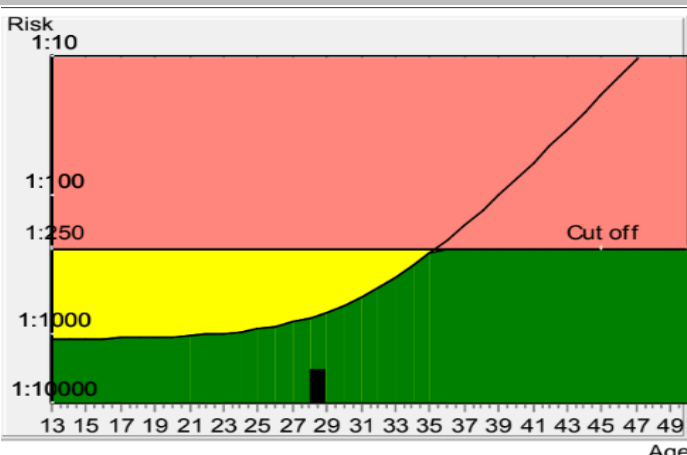


Date of Report 26/02/2020  
PRISCA 5.0.2.37

Patient Data				
Name	Mrs Seema Devi	Patient ID	012002250007	
Birthday	23/08/1991	Sample ID	10684289	
Age at delivery	28.5	Sample Date	25/02/2020	
Gestational age	11+5			
Correction factors				
Fetuses	1 IVF	unknown	Previous trisomy 21	unknown
Weight in kg	46	Diabetes	no	Pregnancies
Smoker	no	Origin	Asian	
Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	11+4
PAPP-A	4.18 mIU/ml	0.94	Method	CRL (<>Robinson)
fb-hCG	89.6 ng/ml	1.59	Scan Date	24/02/2020
Risks at sampling date			Crown Rump Length (mm)	48.3
Age Risk	1:737		Nuchal translucency MoM	0.76
Biochemical Trisomy 21 Risk	1:1357		Nasal Bone	present
Combined Trisomy 21 Risk	1:7320		Sonographer	DR. NITIN GARG
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 7320 women with the same data, there is one woman with a trisomy 21 pregnancy and 7319 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