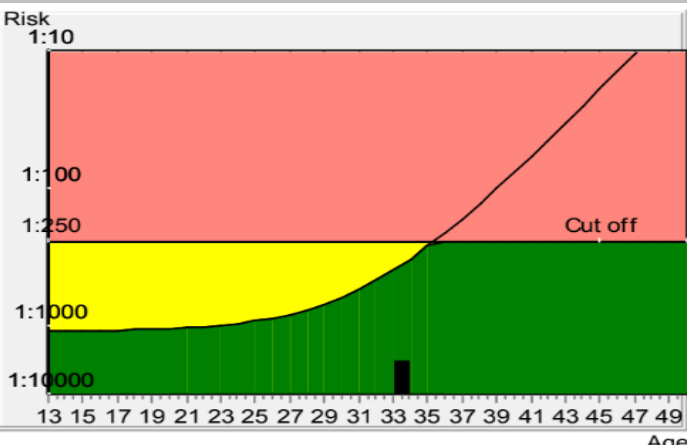


Date of Report 17/02/2020
PRISCA 5.0.2.37

Patient Data				
Name	Mrs Sonali Singh	Patient ID	012002160016	
Birthday	28/07/1986	Sample ID	10520408	
Age at delivery	33.6	Sample Date	16/02/2020	
Gestational age	11+6			
Correction factors				
Fetuses	1	IVF	unknown	Previous trisomy 21
Weight in kg	59.8	Diabetes	no	Pregnancies
Smoker	no	Origin	Asian	
Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	11+4
PAPP-A	1.67 mIU/ml	0.48	Method	CRL (<>Robinson)
fb-hCG	29.8 ng/ml	0.59	Scan Date	14/02/2020
Risks at sampling date			Crown Rump Length (mm)	48.3
Age Risk		1:357	Nuchal translucency MoM	0.91
Biochemical Trisomy 21 Risk		1:1137	Nasal Bone	present
Combined Trisomy 21 Risk		1:5797	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 5797 women with the same data, there is one woman with a trisomy 21 pregnancy and 5796 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