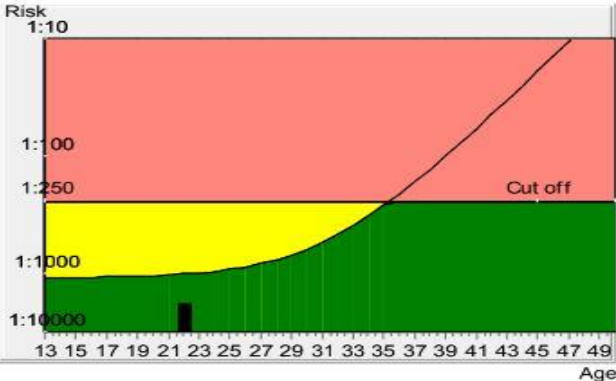




Date of Report 23-01-20
PRISCA 5.0.2.37

Patient Data				
Name	MRS POONAM	Patient ID	012001220176	
Birthday	25-12-97	Sample ID	10370286	
Age at delivery	22.1	Sample Date	22/01/2020	
Gestational age	11+5			
Correction factors				
Fetuses	1	IVF	unknown	Previous trisomy 21
Weight in kg	60.9	Diabetes	unknown	Pregnancies
Smoker	Unknown	Origin	Asian	
Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	11+4
PAPP-A	1.62 mIU/ml	0.51	Method	CRL(<>Robinson
fb-hCG	21.69 ng/ml	0.43	Scan date	21-01-20
Risks at sampling date			Crown rump length in mm	51.2
Age Risk	1:1023		Nuchal translucency MOM	0.87
Biochemical T21 risk	1:7038		Nasal bone	Present
Combined Trisomy 21 Risk	<1:10000		Sonographer	DR.DEEPAK BANSAL
Trisomy 13/18 + NT	<1:10000		Qualification in measuring NT	MBBS,MD
Risk			Down's Syndrome Risk (Trisomy 21 Screening)	
			<p>The calculated risk for Trisomy 21 (with NT) is below the cut off, which represents a low risk.</p> <p>After the result of the Trisomy 21 test (with NT) it is expected that among more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy and 9999 women with not affected pregnancies. 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 NT) is <1:10000, which indicates a low risk				

 Risk Above Cut Off

 Risk above Age Risk

 Risk below Age risk