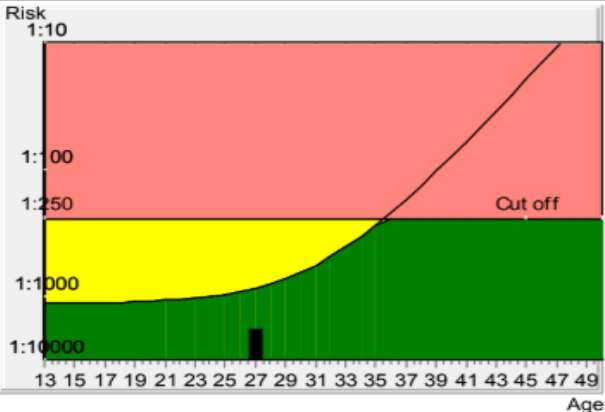


Date of Report 29-01-2020
PRISCA 5.0.2.37

Patient Data				
Name	MRS PRIYANKA	Patient ID	012001280114	
Birthday	01-01-1993	Sample ID	10492624	
Age at delivery	27.1	Sample Date	28/1/2020	
Gestational age	12+6			
Correction factors				
Fetuses	1	IVF	unknown	Previous trisomy 21
Weight in kg	70	Diabetes	unknown	Pregnancies
Smoker	Unknown	Origin	Asian	
Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom		
PAPP-A	3.18 mIU/ml	0.75	Gestational age	12+5
fb-hCG	37.3 ng/ml	0.87	Method	CRL(<>Robinson
Risks at sampling date			Scan date	27-01-2020
Age Risk	1:865		Crown rump length in mm	62.7
Biochemical T21 risk	1:3873		Nuchal translucency MOM	0.93
Combined Trisomy 21 Risk	<1:10000		Nasal bone	Present
Trisomy 13/18 + NT	<1:10000		Sonographer	DR.RUBY RAHUL
Risk			Qualification in measuring NT	CON. RADIOLOGIT
			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	
<p>The calculated risk for Trisomy 13/18 (with NT) is <1:10000, which indicates a low risk</p>				

 Risk Above Cut Off

 Risk above Age Risk

 Risk below Age risk