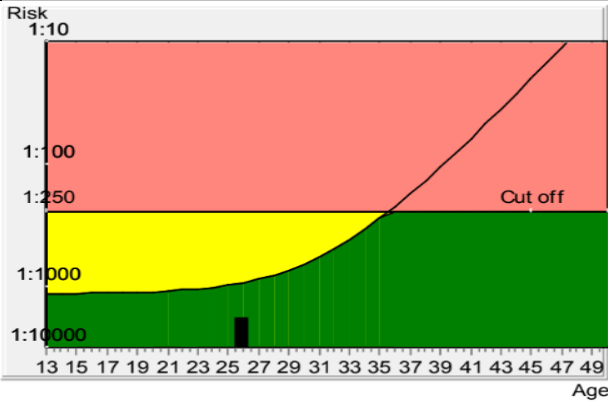


Date of Report 25-01-2020
PRISCA 5.0.2.37

Patient Data				
Name	MRS AMRITA	Patient ID	012001240142	
Birthday	06-03-1994	Sample ID	10567081	
Age at delivery	25.9	Sample Date	24/1/2020	
Gestational age	13+3			
Correction factors				
Fetuses	1 IVF	unknown	Previous trisomy 21	unknown
Weight in kg	60	Diabetes	unknown	Pregnancies
Smoker	Unknown	Origin	Asian	
Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom		
PAPP-A	2.78 mIU/ml	0.45	Gestational age	13+3
fb-hCG	20.5 ng/ml	0.49	Method	CRL(<>Robinson
Risks at sampling date			Scan date	24-01-2020
Age Risk	1:949		Crown rump length in mm	81
Biochemical T21 risk	1:3742		Nuchal translucency MOM	0.67
Combined Trisomy 21 Risk	<1:10000		Nasal bone	Present
Trisomy 13/18 + NT	<1:10000		Sonographer	DR.SONIA B DHANADIA
Risk			Qualification in measuring NT	DMRD,DNB
			Down's Syndrome Risk (Trisomy 21 Screening)	
Trisomy 13/18 + NT The calculated risk for Trisomy 13/18 (with NT) is <1:10000, which indicates a low risk			The calculated risk for Trisomy 21 (with NT) is below the cut off, which represents a low risk. 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!	
The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values				

Risk Above Cut Off
 Risk above Age Risk
 Risk below Age risk