		Date of Report	18/9/2021
		PRISCA	5.2.0.13

Patient Data			
Name	**MRS.DIVYA**	Patient ID	052109180008
Birthday	26/01/1989	Sample ID	11103345
Age at delivery	33.17	Sample Date	18/9/2021
Gestational age	12+5		

Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	70.5	Diabetes	NO	Pregnancies	unknown
Smoker	NO	Origin	Asian		

Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	12+4
PAPP-A	1.14 mIU/ml	0.3	Method	CRL(<>Robinson
fb-hCG	48.3 ng/ml	1.21	Scan date	17/9/2021

Risks at sampling date		Nuchal translucency in mm	0.8
Age Risk	1:735	Nuchal translucency MOM	0.49
Biochemical T21 risk	1:104	Nasal bone	Present
Combined Trisomy 21 Risk	1:610		
Trisomy 13/18 + NT	<1:10000	Qualification in measuring NT	CR

Risk

Down's Syndrome Risk (Trisomy 21 Screening)
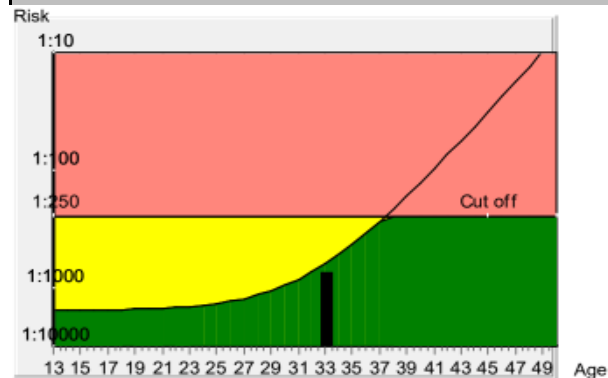
The calculated risk for Trisomy 21 (with NT) is below the cut off, which represents a low risk.

After the result of the Trisomy 21 test (with NT) it is expected that among 610 women with the same data, there is one woman with a trisomy 21 pregnancy and 609 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 aapproaches and have no diagnostic value!

Trisomy 13/18 + NT

The calculated risk for Trisomy 13/18 (with NT) is <1:10000, which indicates a low risk

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