		Date of Report	27-10-18
		PRISCA	5.0.2.37

Patient Data			
Name	**Mrs Suman W/o Jai**	Patient ID	011810260005
Birthday	01-06-84	Sample ID	10377474
Age at delivery	34.4	Sample Date	26/10/18
Gestational age	13+2		

Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	74.5	Diabetes	unknown	Pregnancies	unknown
Smoker	Unknown	Origin	Asian		

Biochemical Data			**Ultrasound Data**	
Parameter	Value	Corr Mom	Gestational age	13+1
PAPP-A	3.41 mIU/ml	0.88	Method	CRL (<>Robinson)
fb-hCG	57.49 ng/ml	1.49	Scan Date	25-10-18
Risks at sampling date			Crown rump length (mm)	67
Age Risk		1:318	Nuchal translucency MoM	1.53
Biochemical T21 Risk		1:594	Nasal Bone	Present
Combined Trisomy 21 Risk		1:364	Sonographer	DR.ANJALI GUPTA
Trisomy 13/18 + NT		<1:10000	Qualifications in measuring NT	MD

Risk	**Down's Syndrome Risk (Trisomy 21 Screening)**

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.
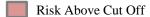
After the result of the Trisomy 21 test (with NT) it is expected that among 364 with the same data, there is one women with a trisomy 21 pregnancy and 363 women with not affected pregnancies.

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!

The patient combined risk presumes the NT measurement was done according to accepted guidelines.

Trisomy 13/18 + NT

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

The laboratory can not be held responsible for their impact on the risk assessment! Calculated value has no diagnostic value!

🟥 Risk Above Cut Off	🟨 Risk above Age Risk	🟩 Risk below Age risk