Date of Report	15-07-2024
PRISCA	5.2.0.13

Patient Data			
Name	**MRS. POOJA YADAV**	Patient ID	12407130122
Birthday	01-07-1993	Sample ID	11893892
Age at Sample date	31.0	Sample Date	14-07-2024
Gestational age	13+5		

Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	64.1	Diabetes	NO	Pregnancies	unknown
Smoker	NO	Origin	Asian		

Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	13+5
PAPP-A	6.8 mIU/ml	0.89	Method	CRL(<>Robinson)
fb-hCG	22.1 ng/ml	0.87	Scan date	13-07-2024

Risks at sampling date		Ultrasound Data (cont.)	
		Crown rump length in mm	76.6
Age Risk	1:586	Nuchal translucency MoM	0.80
Biochemical T21 risk	1:3885	Nasal bone	present
Combined trisomy 21 risk	<1:10000	Sonographer	DR. JAG MOHAN
Trisomy 13/18 + NT	<1:10000	Qualifications in measuring NT	MD

Risk

Down's Syndrome Risk (Trisomy 21 Screening)

The calculated risk for Trisomy 21 test (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 amog more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy.

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 that NT measurement was done according to accepted guidelines (Prenat Diagn 18:511-523; 1998).

The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values

Trisomy 13/18 + NT

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