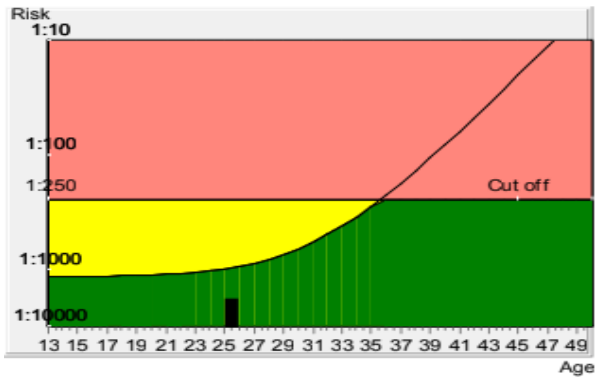
Date of Report	10/4/2023
PRISCA	5.2.0.13

Patient Data			
Name	MRS. SIMRANJIT KAUR	Patient ID	012304090128
Birthday	9/11/1997	Sample ID	11586189
Age at term	25.1	Sample Date	9/4/2023
Gestational age	13+5		

Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	48	Diabetes	NO	Pregnancies	unknown
Smoker	NO	Origin	Asian		

Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom		13+4
PAPP-A	4.87 mIU/ml	0.46	Method	CRL (<>Robinson)
fb-hCG	18.8 ng/ml	0.67	Scan date	8/4/2023
			Nuchal translucency	1.46
Risks at sampling date			Nuchal translucency MoM	0.79
Age Risk		1:981	Nasal bone	Present
Biochemical T21 risk		1:2110		
Combined trisomy 21 risk		<1:10000		
Trisomy 13/18 + NT		<1:10000		

Risk



The calculated risk for Trisomy 13/18 (with NT) is <1:10000 , which indicates a low 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 21with NT test 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 aapproaches 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