Date of Report		20/4/2023	
PRISCA		5.2.0.13	

Patient Data			
Name	MRS. VANDANA	Patient ID	012304190082
Birthday	16/10/1992	Sample ID	11671159
Age at term	30.1	Sample Date	19/4/2023
Gestational age	13+3		

Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	84	Diabetes	NO	Pregnancies	unknown
Smoker	NO	Origin	Asian		

Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom		13+2
PAPP-A	3.45 mIU/ml	0.69	Method	CRL (<>Robinson)
fb-hCG	11.8 ng/ml	0.45	Scan date	18/04/2023

Risks at sampling date		Nuchal translucency	1.4
Age Risk	1:624	Nuchal translucency MoM	0.78
Biochemical T21 risk	1:8619	Nasal bone	Present
Combined trisomy 21 risk	<1:10000		
Trisomy 13/18 + NT	<1:10000		
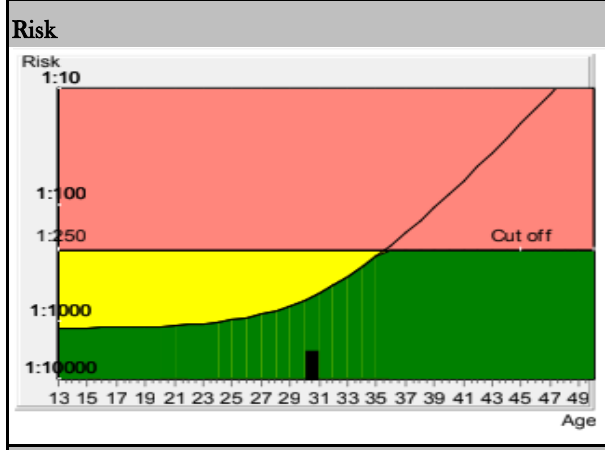
Risk

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

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