Date of Report	22/4/2023
PRISCA	5.2.0.13

Patient Data			
Name	MRS. FARHEEN	Patient ID	012304210155
Birthday	9/10/1995	Sample ID	11372088
Age at term	28	Sample Date	21/4/2023
Gestational age	13+6		

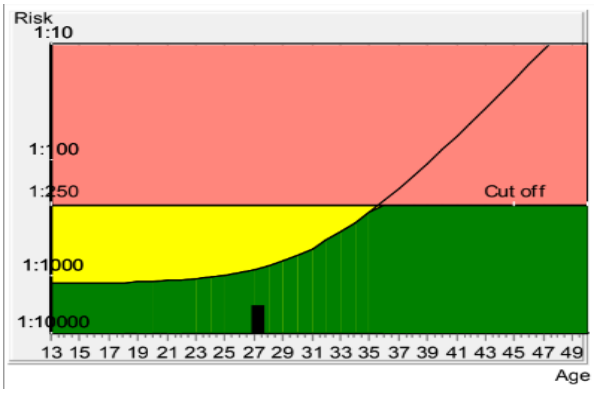
Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	73	Diabetes	NO	Pregnancies	unknown
Smoker	NO	Origin	Asian		

Biochemical Data		
Parameter	**Value**	**Corr Mom**
PAPP-A	4.27 mIU/ml	0.45
fb-hCG	18.5 ng/ml	0.65

Risks at sampling date	
Age Risk	1:879
Biochemical T21 risk	1:1921
Combined trisomy 21 risk	1:8706
Trisomy 13/18 + NT	<1:10000

Ultrasound Data	
	13+3
Method	CRL (<>Robinson)
Scan date	18/04/2023
Nuchal translucency	1.8
Nuchal translucency MoM	0.99
Nasal bone	Present

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 8076 women with the same data, there is one woman with a trisomy 21 pregnancy and 8075 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