Date of Report	13/11/2022
PRISCA	5.1.0.17

Patient Data			
Name	MRS. KOMAL	Patient ID	012211120126
Birthday	12/6/1996	Sample ID	11510107
Age at term	26.1	Sample Date	12/11/2022
Gestational age	13+1		

Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	38	Diabetes	NO	Pregnancies	unknown
Smoker	NO	Origin	Asian		

Biochemical Data

Parameter	Value	Corr Mom
PAPP-A	5.6 mIU/ml	0.49
fb-hCG	51.4 ng/ml	1.38

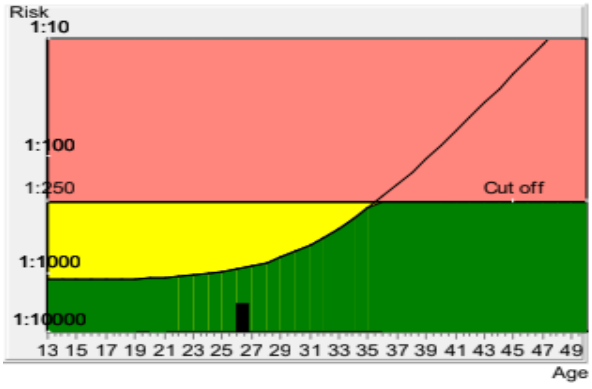
Risks at sampling date

Age Risk	1:912
Biochemical T21 risk	1:482
Combined trisomy 21 risk	1:3031
Trisomy 13/18 + NT	<1:10000

Ultrasound Data

Gestational age	13+1
Method	CRL (<>Robinson)
Scan date	12/11/2022
Nuchal translucency	1.2
Nuchal translucency MoM	0.68
Nasal bone	Present

Risk



Trisomy 13/18+NT

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 3031 women with the same data, there is one woman with a trisomy 21 pregnancy and 3030 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