Date of Report	1/5/2022
PRISCA	5.1.0.17

Patient Data			
Name	MRS. MANISHA	Patient ID	012204300001
Birthday	14/08/1998	Sample ID	11242497
Age at term	24.01	Sample Date	30/4/2022
Gestational age	13+2		

Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	59	Diabetes	NO	Pregnancies	unknown
Smoker	NO	Origin	Asian		

Biochemical Data

Parameter	Value	Corr Mom
PAPP-A	5.6 mIU/ml	1.27
fb-hCG	58.4 ng/ml	1.23

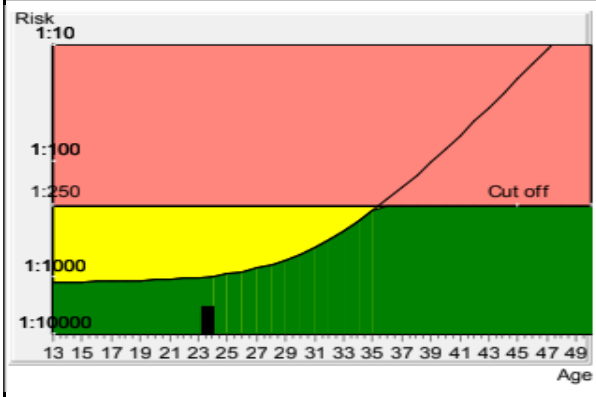
Risks at sampling date

Age Risk	1:1006
Biochemical T21 risk	1:6295
Combined T21 risk	<1:10000
Trisomy 13/18+NT	<1:10000

Ultrasound Data

Gestational age	12+2
Method	CRL (<>Robinson)
Scan date	29/4/2022
Nuchal Translucency	1.2
Nuchal Translucency MoM	0.79
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 21 test it is expected that among more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy 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