Date of Report	13/11/2022
PRISCA	5.1.0.17

Patient Data			
Name	MRS. MANPREET KAUR	Patient ID	012211120016
Birthday	26/1/1995	Sample ID	11468259
Age at term	28.02	Sample Date	12/11/2022
Gestational age	12+1		

Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	71.3	Diabetes	NO	Pregnancies	unknown
Smoker	NO	Origin	Asian		

Biochemical Data

Parameter	Value	Corr Mom
PAPP-A	5.1 mIU/ml	1.33
fb-hCG	21.8 ng/ml	0.56

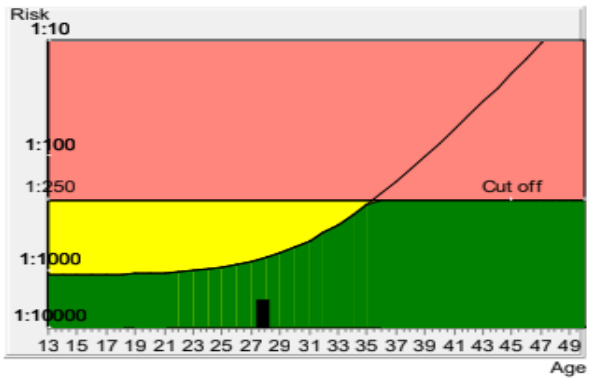
Ultrasound Data

Gestational age	11+6
Method	CRL (<>Robinson)
Scan date	10/11/2022
Nuchal translucency	1.2
Nuchal translucency MoM	0.86
Nasal bone	Present

Risks at sampling date

Age Risk	1:798
Biochemical T21 risk	<1:10000
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

Trisomy 13/18+NT

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