Date of Report	14-01-2025
PRISCA	5.2.0.13

Patient Data			
Name	MRS. HARSHITA	Patient ID	12501110161
Birthday	21-06-1998	Sample ID	11484405
Age at Sample date	26.6	Sample Date	11-01-2025
Gestational age	12+2		

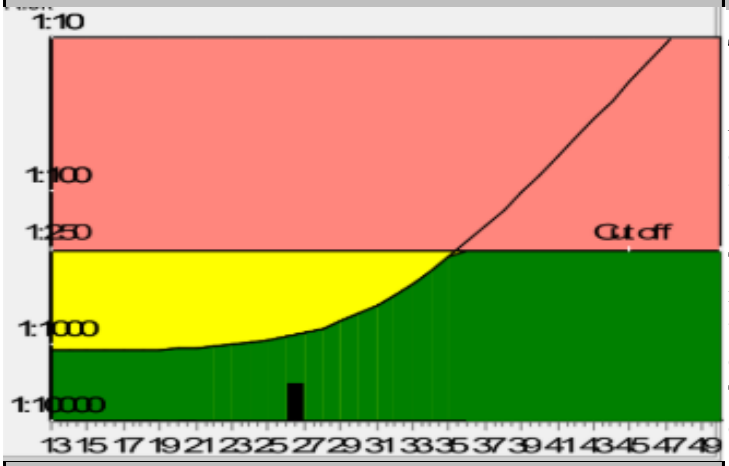
Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	70	Diabetes	NO	Pregnancies	unknown
Smoker	NO	Origin	Asian		

Biochemical Data		
Parameter	**Value**	**Corr Mom**
PAPP-A	4.8 mIU/ml	1.17
fb-hCG	27.4 ng/ml	0.72

Risks at sampling date	
Age Risk	1:877
Biochemical T21 risk	<1:10000
Combined trisomy 21 risk	<1:10000
Trisomy 13/18 + NT	<1:10000

Ultrasound Data	
Gestational age	12+1
Method	CRL (<>Robinson)
Scan date	11-01-2025
Crown rump length in mm	51
Nuchal translucency MoM	0.85
Nasal bone	PRESENT
Sonographer	DR. SHRUTI SANGWAN
Qualifications in measuring NT	MBBS

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 (with NT) it is expected that among more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy.

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 patient combined risk presumes that NT measurement was done according to accepted guidelines (Prenat Diagn 18:511-523; 1998).

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