Prisca 5.2.0.13

Date of report: 28/03/2025

JITM Diagnostics

Patient data			
Name	MRS. PRIYAKA	Patient ID	
Birthday	26/02/1992	Sample ID	2503056188/NOD
Age at sample date	33.1	Sample Date	27/03/2025
Gestational age	13 + 1		

Correction factors					
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	no
Weight	64.6	diabetes	no		
Smoker	no	Origin	Asian		

Biochemical data

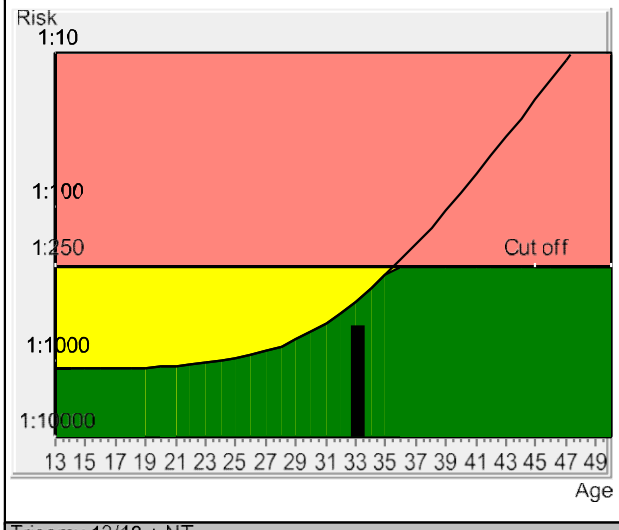
Parameter	Value	Corr. MoM
PAPP-A	3.19 mIU/ml	0.68
fb-hCG	107 ng/ml	2.62

Risks at sampling date

Age risk	1:410
Biochemical T21 risk	1:100
Combined trisomy 21 risk	1:594
Trisomy 13/18 + NT	<1:10000

Ultrasound data

Gestational age	13 + 0
Method	CRL Robinson
Scan date	26/03/2025
Crown rump length in mm	70.3
Nuchal translucency MoM	0.85
Nasal bone	present
Sonographer	..
Qualifications in measuring NT	..



Trisomy 13/18 + NT

The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.

Trisomy 21

The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.

After the result of the Trisomy 21 test (with NT) it is expected that among 594 women with the same data, there is one woman with a trisomy 21 pregnancy and 593 women with not affected pregnancies.
The free beta HCG level is high.
The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician.
Please note that risk calculations are statistical approaches and have no diagnostic value!
The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).
The laboratory can not be hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!

Sign of Physician