Prisca 5.2.0.13

Date of report: 29/03/2025

JITM Diagnostics

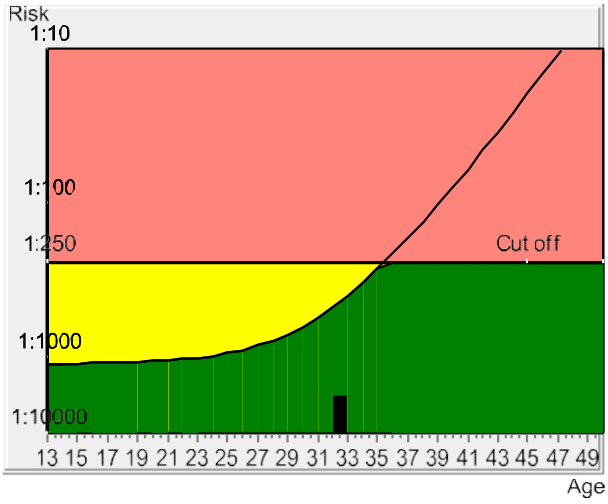
Patient data			
Name	MRS. BHAWANA	Patient ID	
Birthday	21/10/1992	Sample ID	2503059597/NOD
Age at sample date	32.4	Sample Date	29/03/2025
Gestational age	12 + 3		

Correction factors					
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	no
Weight	53.7	diabetes	no		
Smoker	no	Origin	Asian		

Biochemical data		
Parameter	Value	Corr. MoM
PAPP-A	2.88 mIU/ml	0.66
fb-hCG	37.1 ng/ml	0.79

Risks at sampling date	
Age risk	1:448
Biochemical T21 risk	1:1796
Combined trisomy 21 risk	1:9625
Trisomy 13/18 + NT	<1:10000

Ultrasound data	
Gestational age	12 + 2
Method	CRL Robinson
Scan date	28/03/2025
Crown rump length in mm	60.5
Nuchal translucency MoM	0.70
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 9625 women with the same data, there is one woman with a trisomy 21 pregnancy and 9624 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 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