Prisca 5.2.0.13

Date of report: 23/01/2025

Patient data

Name	MRS. SANDHYA GHARANA	Patient ID	
Birthday	14/01/1989	Sample ID	2501035367/NOD
Age at sample date	36.0	Sample Date	22/01/2025
Gestational age	12 + 3		

Correction factors

Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	no
Weight	67	diabetes	no		
Smoker	no	Origin	Asian		

Biochemical data

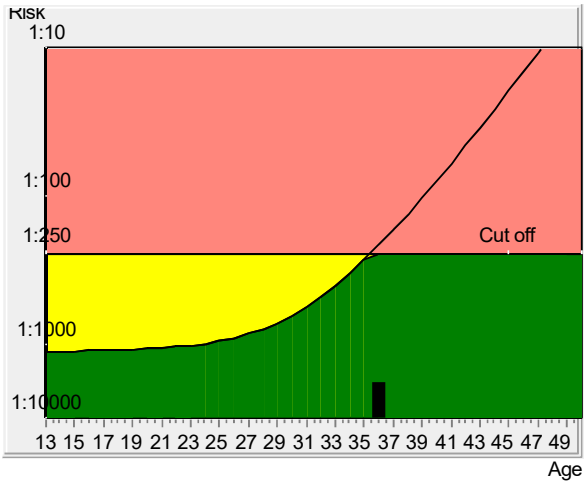
Parameter	Value	Corr. MoM
PAPP-A	2.61 mIU/ml	0.78
fb-hCG	41.6 ng/ml	0.95

Risks at sampling date

Age risk	1:215
Biochemical T21 risk	1:850
Combined trisomy 21 risk	1:2441
Trisomy 13/18 + NT	<1:10000

Ultrasound data

Gestational age	12 + 3
Method	CRL Robinson
Scan date	22/01/2025
Crown rump length in mm	61.5
Nuchal translucency MoM	1.13
Nasal bone	present
Sonographer	DR. SHAILEN SINGHA
Qualifications in measuring NT	M.D



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 2441 women with the same data, there is one woman with a trisomy 21 pregnancy and 2440 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

below cut off | Below Cut Off, but above Age Risk | above cut off