Prisca 5.1.0.17

Date of report: 02/09/25

NA

Patient data			
Name	Mrs. PRAJAKTA NAGBHIDE	Patient ID	0372508300181
Birthday	28/05/93	Sample ID	B3128616
Age at sample date	32.3	Sample Date	30/08/25
Gestational age	12 + 1		

Correction factors					
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	unknown
Weight	63	diabetes	no		
Smoker	no	Origin	Asian		

Biochemical data

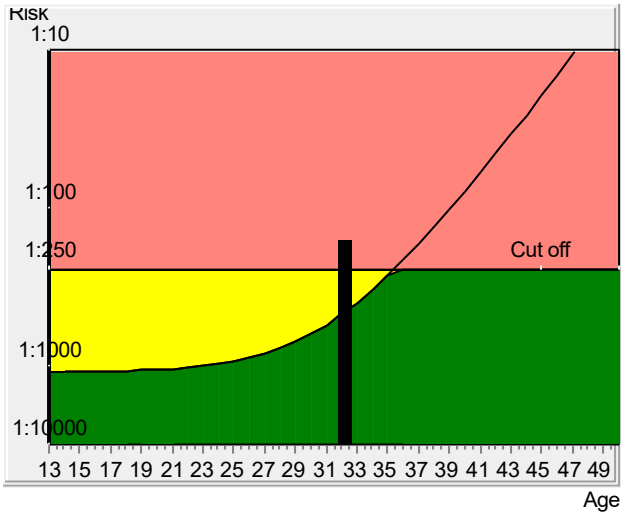
Parameter	Value	Corr. MoM
PAPP-A	0.65 mIU/mL	0.19
fb-hCG	42.41 ng/ml	1.01

Risks at sampling date

Age risk	1:457
Biochemical T21 risk	>1:50
Combined trisomy 21 risk	1:162
Trisomy 13/18 + NT	1:388

Ultrasound data

Gestational age	11 + 6
Method	CRL Robinson
Scan date	28/08/25
Crown rump length in mm	55
Nuchal translucency MoM	1.03
Nasal bone	present
Sonographer	NA
Qualifications in measuring NT	MD



Trisomy 13/18 + NT

The calculated risk for Trisomy 13/18 (with nuchal translucency) is 1:388, which represents a low risk.

Trisomy 21

The calculated risk for Trisomy 21 (with nuchal translucency) is above the cut off, which indicates an increased risk.

After the result of the Trisomy 21 test (with NT) it is expected that among 162 women with the same data, there is one woman with a trisomy 21 pregnancy and 161 women with not affected pregnancies.
The PAPP-A level is low.
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