SAGEPATH LABS PVT LTD.

Prisca 5.1.0.17

Date of report: 09/09/25

NA

Patient data			
Name	Mrs. P.POOJITHA	Patient ID	0942509030002
Birthday	22/06/99	Sample ID	B3002598
Age at sample date	26.2	Sample Date	03/09/25
Gestational age	13 + 0		

Correction factors					
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	unknown
Weight	48	diabetes	no		
Smoker	no	Origin	Asian		

Biochemical data

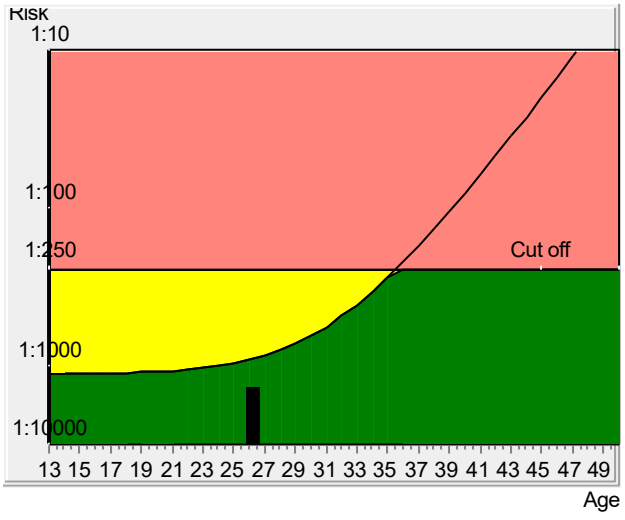
Parameter	Value	Corr. MoM
PAPP-A	2.02 mIU/mL	0.29
fb-hCG	36.62 ng/ml	0.95

Risks at sampling date

Age risk	1:919
Biochemical T21 risk	1:241
Combined trisomy 21 risk	1:1387
Trisomy 13/18 + NT	1:9407

Ultrasound data

Gestational age	12 + 6
Method	CRL Robinson
Scan date	02/09/25
Crown rump length in mm	68
Nuchal translucency MoM	0.93
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 1:9407, 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 1387 women with the same data, there is one woman with a trisomy 21 pregnancy and 1386 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