SAGEPATH LABS PVT LTD.,

Prisca 5.1.0.17

Date of report: 11-07-2024

Patient data

Name	Mrs. PRANITA PAWAR	Patient ID	0662407080270
Birthday	19-07-1999	Sample ID	A0500475
Age at sample date	25.0	Sample Date	08-07-2024
Gestational age	13 + 3		

Correction factors

Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	unknown
Weight	49	diabetes	no		
Smoker	no	Origin	Asian		

Biochemical data

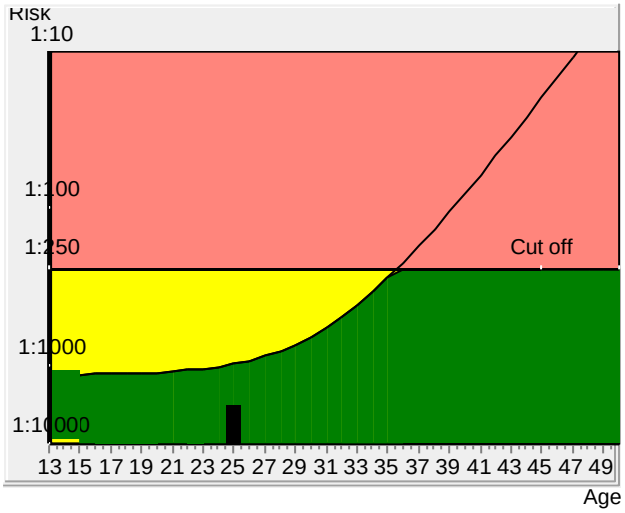
Parameter	Value	Corr. MoM
PAPP-A	5.97 mIU/mL	0.85
fb-hCG	52.66 ng/mL	1.48

Risks at sampling date

Age risk	1:992
Biochemical T21 risk	1:1720
Combined trisomy 21 risk	1:8381
Trisomy 13/18 + NT	<1:10000

Ultrasound data

Gestational age	13 + 0
Method	CRL Robinson
Scan date	05-07-2024
Crown rump length in mm	69.7
Nuchal translucency MoM	0.91
Nasal bone	present
Sonographer	NA
Qualifications in measuring NT	MD



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 8381 women with the same data, there is one woman with a trisomy 21 pregnancy and 8380 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!

Trisomy 13/18 + NT

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

Sign of Physician

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