SAGEPATH LABS PVT LTD.,

Prisca 5.1.0.17

Date of report: 17/02/22

Patient data			
Name	Mrs. YESHASHWINI	Patient ID	0012202160468
Birthday	01/10/95	Sample ID	23133901
Age at sample date	26.4	Sample Date	16/02/22
Gestational age	13 + 0		

Correction factors					
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	unknown
Weight	52.7	diabetes	no		
Smoker	no	Origin	Asian		

Biochemical data

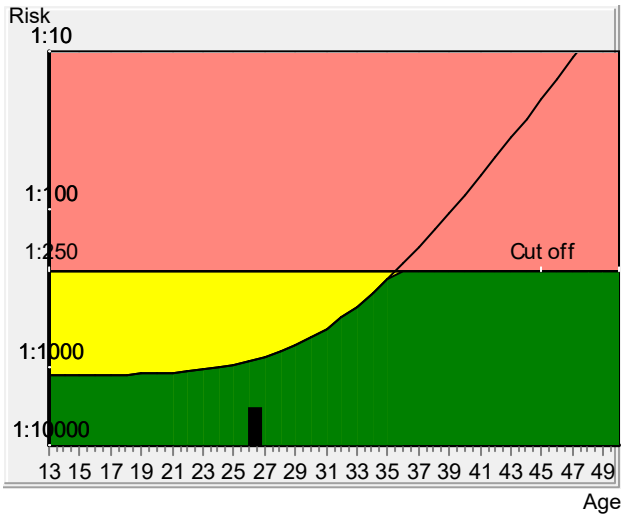
Parameter	Value	Corr. MoM
PAPP-A	4.21 mIU/mL	0.76
fb-hCG	49.21 ng/mL	1.27

Risks at sampling date

Age risk	1:910
Biochemical T21 risk	1:1777
Combined trisomy 21 risk	1:8795
Trisomy 13/18 + NT	<1:10000

Ultrasound data

Gestational age	13 + 0
Method	CRL Robinson
Scan date	16/02/22
Crown rump length in mm	70
Nuchal translucency MoM	0.91
Nasal bone	present
Sonographer	DR C VENU GOPAL
Qualifications in measuring NT	MBBS DMRD



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 8795 women with the same data, there is one woman with a trisomy 21 pregnancy and 8794 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