SAGEPATH LABS PVT LTD.,

Prisca 5.1.0.17

Date of report: 30/12/21

Patient data			
Name	Mrs. YARA.MANASA	Patient ID	0012112300056
Birthday	10/05/99	Sample ID	23087043
Age at sample date	22.6	Sample Date	30/12/21
Gestational age	13 + 0		

Correction factors					
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	unknown
Weight	45	diabetes	no		
Smoker	no	Origin	Asian		

Biochemical data

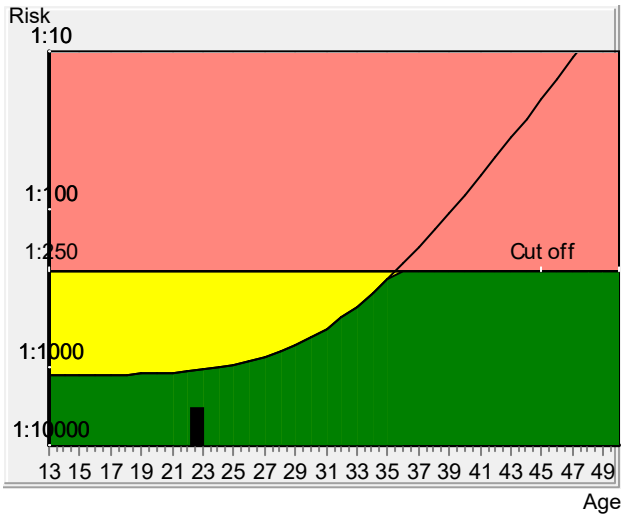
Parameter	Value	Corr. MoM
PAPP-A	2.7 mIU/mL	0.41
fb-hCG	33.22 ng/mL	0.81

Risks at sampling date

Age risk	1:1056
Biochemical T21 risk	1:1118
Combined trisomy 21 risk	1:6833
Trisomy 13/18 + NT	<1:10000

Ultrasound data

Gestational age	12 + 1
Method	CRL Robinson
Scan date	24/12/21
Crown rump length in mm	58.5
Nuchal translucency MoM	0.72
Nasal bone	present
Sonographer	DR K MANDAKINI
Qualifications in measuring NT	MDRD



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 6833 women with the same data, there is one woman with a trisomy 21 pregnancy and 6832 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