SagePath Labs Pvt Ltd.,

Prisca 5.1.0.17

Date of report: 03/04/21

SHIRISHA

Patient data			
Name	Mrs. MAMATHA	Patient ID	0012104010191
Birthday	04/01/97	Sample ID	20123792
Age at sample date	24.2	Sample Date	01/04/21
Gestational age	13 + 3		

Correction factors					
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	unknown
Weight	46	diabetes	no		
Smoker	no	Origin	Asian		

Biochemical data

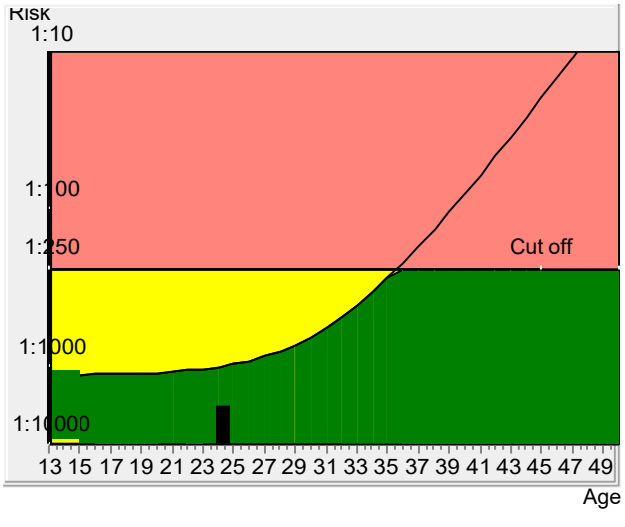
Parameter	Value	Corr. MoM
PAPP-A	5.14 mIU/mL	0.68
fb-hCG	45.32 ng/mL	1.25

Risks at sampling date

Age risk	1:1021
Biochemical T21 risk	1:1590
Combined trisomy 21 risk	1:9025
Trisomy 13/18 + NT	<1:10000

Ultrasound data

Gestational age	13 + 2
Method	CRL Robinson
Scan date	31/03/21
Crown rump length in mm	75
Nuchal translucency MoM	0.70
Nasal bone	present
Sonographer	DR NA
Qualifications in measuring NT	MD



Trisomy 13/18 + NT

The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, 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 9025 women with the same data, there is one woman with a trisomy 21 pregnancy and 9024 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!

Sign of Physician

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