SagePath Labs Pvt Ltd.,

Prisca 5.1.0.17

Date of report: 14/09/21

DEEPTHI REDDY

Patient data			
Name	Mrs. SHARADHA	Patient ID	0012109140365
Birthday	07/07/77	Sample ID	22223046
Age at sample date	44.2	Sample Date	13/09/21
Gestational age	13 + 6		

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

Biochemical data

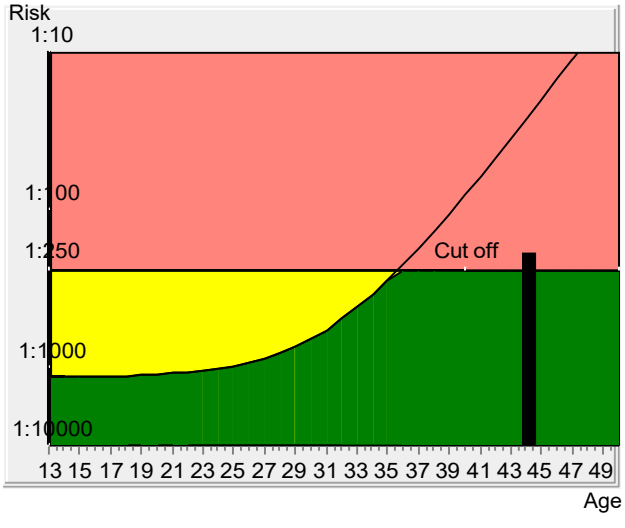
Parameter	Value	Corr. MoM
PAPP-A	5.05 mIU/mL	0.56
fb-hCG	35.87 ng/mL	1.11

Risks at sampling date

Age risk	1:26
Biochemical T21 risk	>1:50
Combined trisomy 21 risk	1:190
Trisomy 13/18 + NT	1:9526

Ultrasound data

Gestational age	12 + 6
Method	CRL Robinson
Scan date	06/09/21
Crown rump length in mm	67.5
Nuchal translucency MoM	0.70
Nasal bone	present
Sonographer	DR NA
Qualifications in measuring NT	MD



Trisomy 21

The calculated risk for Trisomy 21 (with nuchal translucency) is above the cut off, which indicates an increased risk.

After the result of the Trisomy 21 test (with NT) it is expected that among 190 women with the same data, there is one woman with a trisomy 21 pregnancy and 189 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 1:9526, which represents a low risk.

Sign of Physician

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