SAGEPATH LABS PVT LTD.,

Prisca 5.1.0.17

Date of report: 19-07-2022

Patient data			
Name	Mrs. SHEETAL W/O SHRADHA KANTH	Patient ID	0012207170295
Birthday	25-03-1983	Sample ID	23567325
Age at sample date	39.3	Sample Date	17-07-2022
Gestational age	12 + 4		

Correction factors					
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	unknown
Weight	75	diabetes	no		
Smoker	no	Origin	Asian		

Biochemical data

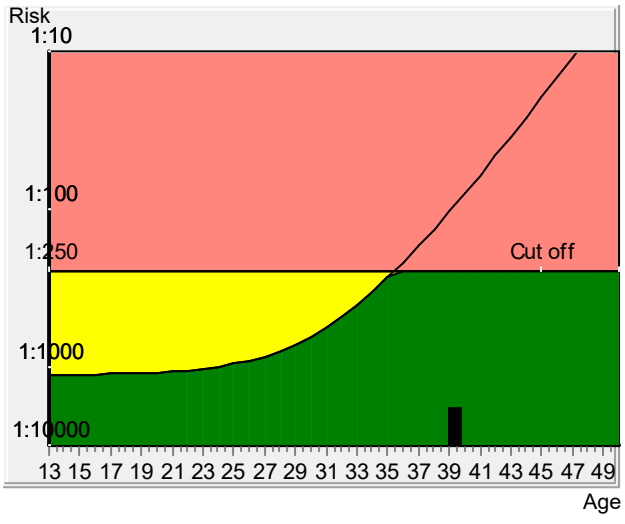
Parameter	Value	Corr. MoM
PAPP-A	3.08 mIU/mL	1.00
fb-hCG	41.22 ng/mL	1.07

Risks at sampling date

Age risk	1:95
Biochemical T21 risk	1:510
Combined trisomy 21 risk	1:2562
Trisomy 13/18 + NT	<1:10000

Ultrasound data

Gestational age	12 + 3
Method	CRL Robinson
Scan date	16-07-2022
Crown rump length in mm	61.7
Nuchal translucency MoM	0.81
Nasal bone	unknown
Sonographer	NA
Qualifications in measuring NT	NAGA SAI TEJA



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 2562 women with the same data, there is one woman with a trisomy 21 pregnancy and 2561 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