SAGEPATH LABS PVT LTD

Prisca 5.1.0.17

Date of report: 24-12-2022

NA

Patient data			
Name	Mrs. SUPRIYA SHINDE	Patient ID	0012212240096
Birthday	05-07-2000	Sample ID	23598442
Age at sample date	22.5	Sample Date	24-12-2022
Gestational age	13 + 2		

Correction factors					
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	unknown
Weight	43.5	diabetes	no		
Smoker	no	Origin	Asian		

Biochemical data

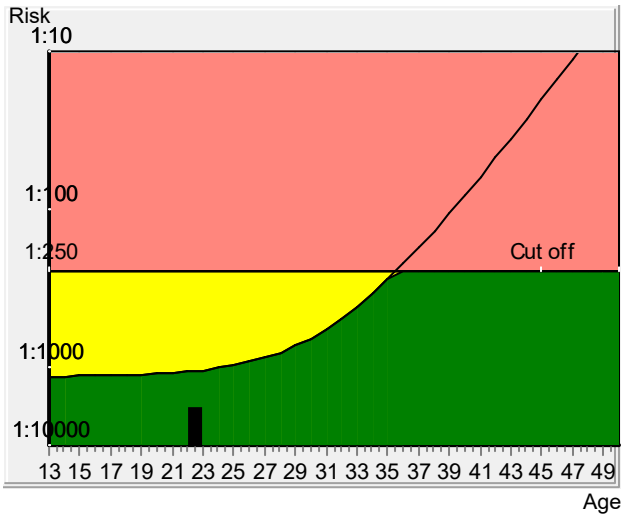
Parameter	Value	Corr. MoM
PAPP-A	3.33 mIU/mL	0.43
fb-hCG	33.1 ng/mL	0.86

Risks at sampling date

Age risk	1:1070
Biochemical T21 risk	1:1205
Combined trisomy 21 risk	1:7298
Trisomy 13/18 + NT	<1:10000

Ultrasound data

Gestational age	12 + 2
Method	CRL Robinson
Scan date	17-12-2022
Crown rump length in mm	60.6
Nuchal translucency MoM	0.76
Nasal bone	present
Sonographer	NA
Qualifications in measuring NT	NA



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 7298 women with the same data, there is one woman with a trisomy 21 pregnancy and 7297 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