SAGEPATH LABS PVT LTD.,

Prisca 5.1.0.17

Date of report: 02-05-2024

Patient data

Name	Mrs. SARITHA W/O. JANARDHAN	Patient ID	0312404300035
Birthday	01-01-1994	Sample ID	A0015467
Age at sample date	30.3	Sample Date	30-04-2024
Gestational age	12 + 3		

Correction factors

Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	unknown
Weight	52	diabetes	no		
Smoker	no	Origin	Asian		

Biochemical data

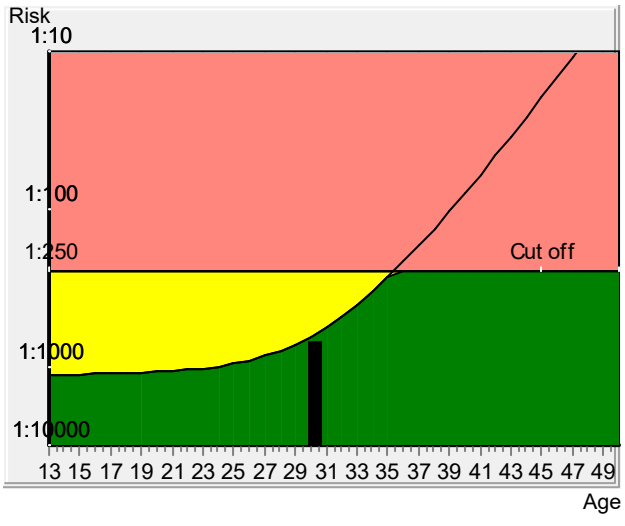
Parameter	Value	Corr. MoM
PAPP-A	3.22 mIU/mL	0.72
fb-hCG	65.12 ng/mL	1.45

Risks at sampling date

Age risk	1:617
Biochemical T21 risk	1:762
Combined trisomy 21 risk	1:699
Trisomy 13/18 + NT	<1:10000

Ultrasound data

Gestational age	12 + 3
Method	CRL Robinson
Scan date	30-04-2024
Crown rump length in mm	62
Nuchal translucency MoM	1.44
Nasal bone	present
Sonographer	NA
Qualifications in measuring NT	MD



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 699 women with the same data, there is one woman with a trisomy 21 pregnancy and 698 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