SAGEPATH LABS PVT LTD.,

Prisca 5.1.0.17

Date of report: 13-01-2023

Patient data

Name	Mrs. P SANDHYA	Patient ID	0352301120057
Birthday	28-11-2001	Sample ID	24106055
Age at sample date	21.1	Sample Date	12-01-2023
Gestational age	11 + 3		

Correction factors

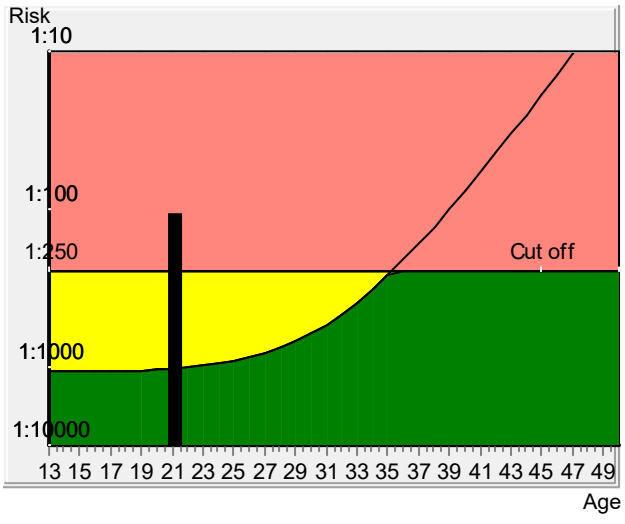
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	unknown
Weight	52.5	diabetes	no		
Smoker	no	Origin	Asian		

Biochemical data

Parameter	Value	Corr. MoM
PAPP-A	2.01 mIU/mL	0.71
fb-hCG	54.22 ng/mL	0.98

Risks at sampling date

Age risk	1:1029
Biochemical T21 risk	1:3061
Combined trisomy 21 risk	1:107
Trisomy 13/18 + NT	1:1200



Trisomy 13/18 + NT

The calculated risk for Trisomy 13/18 (with nuchal translucency) is 1:1200, which represents a low risk.

Ultrasound data

Gestational age	11 + 2
Method	CRL Robinson
Scan date	11-01-2023
Crown rump length in mm	46.7
Nuchal translucency MoM	2.19
Nasal bone	unknown
Sonographer	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 107 women with the same data, there is one woman with a trisomy 21 pregnancy and 106 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