SAGEPATH LABS PVT LTD.,

Prisca 5.1.0.17

Date of report: 24/08/22

Patient data

Name	Mrs. SWAPNA	Patient ID	0012208230470
Birthday	21/04/84	Sample ID	23727326
Age at sample date	38.3	Sample Date	23/08/22
Gestational age	11 + 4		

Correction factors

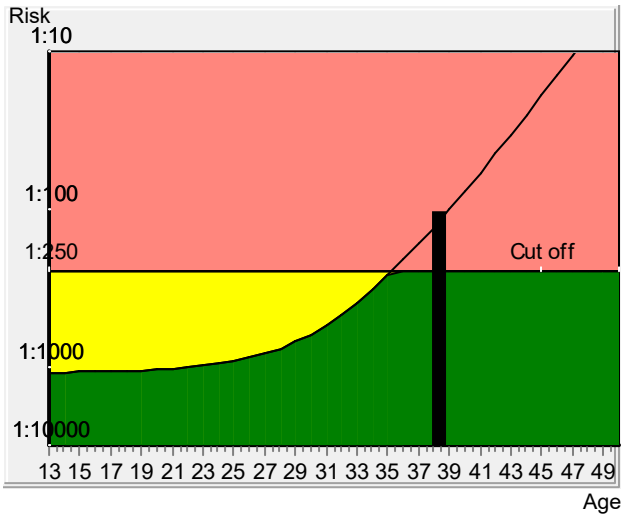
Fetuses	1	IVF	yes	Previous trisomy 21 pregnancies	unknown
Weight	56	diabetes	no		
Smoker	no	Origin	Asian		

Biochemical data

Parameter	Value	Corr. MoM
PAPP-A	0.89 mIU/mL	0.31
fb-hCG	45.66 ng/mL	0.87

Risks at sampling date

Age risk	1:117
Biochemical T21 risk	>1:50
Combined trisomy 21 risk	1:105
Trisomy 13/18 + NT	1:442



Trisomy 13/18 + NT

The calculated risk for Trisomy 13/18 (with nuchal translucency) is 1:442, which represents a low risk.

Ultrasound data

Gestational age	11 + 4
Method	CRL Robinson
Scan date	23/08/22
Crown rump length in mm	50
Nuchal translucency MoM	1.26
Nasal bone	present
Sonographer	NA
Qualifications in measuring NT	NAGA SAI TEJA

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 105 women with the same data, there is one woman with a trisomy 21 pregnancy and 104 women with not affected pregnancies.

The PAPP-A level is low.

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