SagePath Labs Pvt Ltd.,

Prisca 5.1.0.17

Date of report: 08-05-2021

Patient data

Name	Mrs. V SRILATHA	Patient ID	0012105081374
Birthday	14-08-1997	Sample ID	19151830
Age at sample date	23.7	Sample Date	08-05-2021
Gestational age	12 + 1		

Correction factors

Fetuses	1	IVF	unknown	Previous trisomy 21 pregnancies	unknown
Weight	65	diabetes	no		
Smoker	no	Origin	Asian		

Biochemical data

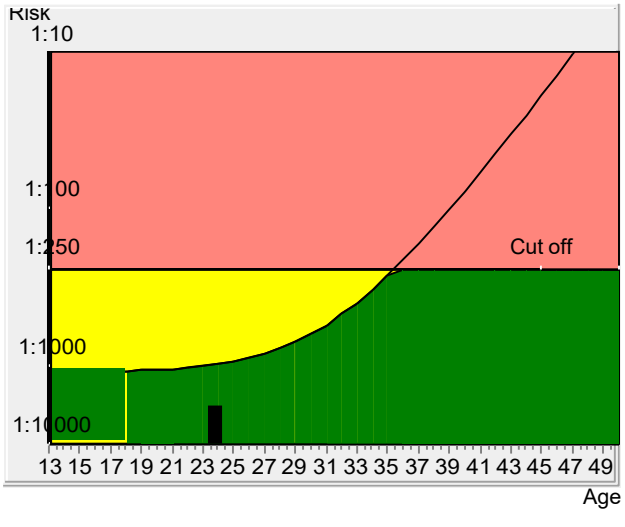
Parameter	Value	Corr. MoM
PAPP-A	1.74 mIU/mL	0.57
fb-hCG	48.69 ng/mL	1.09

Risks at sampling date

Age risk	1:995
Biochemical T21 risk	1:1324
Combined trisomy 21 risk	1:7159
Trisomy 13/18 + NT	<1:10000

Ultrasound data

Gestational age	11 + 1
Method	CRL Robinson
Scan date	01-05-2021
Crown rump length in mm	45
Nuchal translucency MoM	0.89
Nasal bone	present
Sonographer	DR 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 7159 women with the same data, there is one woman with a trisomy 21 pregnancy and 7158 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