SAGEPATH LABS PVT LTD

Prisca 5.1.0.17

Date of report: 27-11-2024

NA

Patient data			
Name	Mrs. MANISHA KUMARI	Patient ID	0472411260082
Birthday	18-11-1997	Sample ID	A1678576
Age at sample date	27.0	Sample Date	26-11-2024
Gestational age	11 + 5		

Correction factors					
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	unknown
Weight	69	diabetes	no		
Smoker	no	Origin	Asian		

Biochemical data

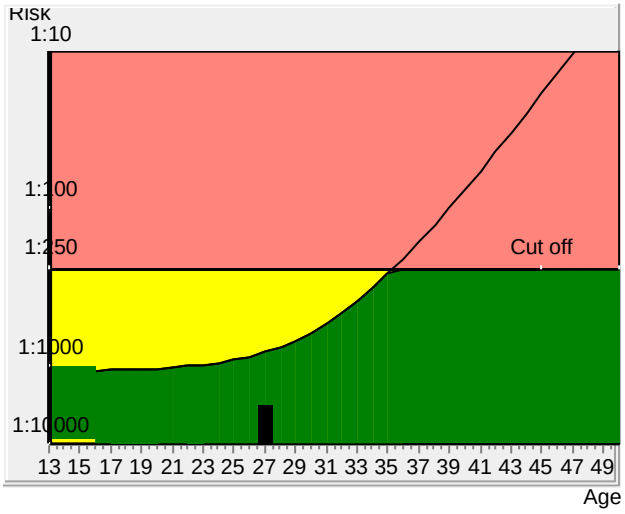
Parameter	Value	Corr. MoM
PAPP-A	2.49 mIU/mL	0.96
fb-hCG	33.92 ng/ml	0.75

Risks at sampling date

Age risk	1:833
Biochemical T21 risk	1:8833
Combined trisomy 21 risk	<1:10000
Trisomy 13/18 + NT	<1:10000

Ultrasound data

Gestational age	11 + 4
Method	CRL Robinson
Scan date	25-11-2024
Crown rump length in mm	51
Nuchal translucency MoM	1.24
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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy.
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