SAGEPATH LABS PVT LTD.

Prisca 5.1.0.17

Date of report: 11-02-2024

Patient data

Name	Mrs. MANISHA W/O RAMESH	Patient ID	0312402100037
Birthday	22-04-1999	Sample ID	a0013788
Age at sample date	24.8	Sample Date	10-02-2024
Gestational age	12 + 0		

Correction factors

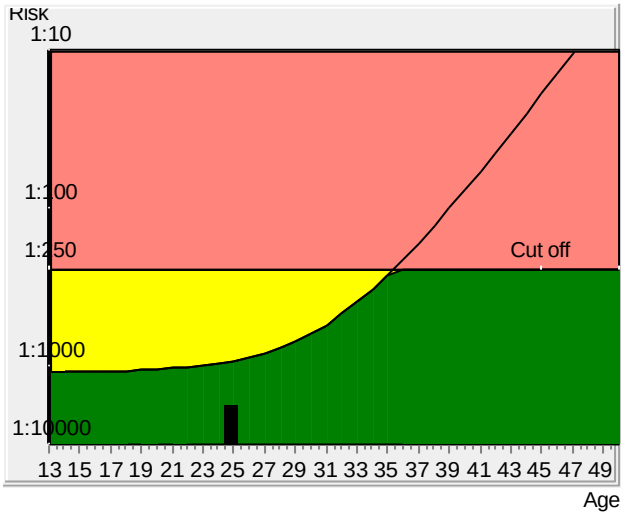
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	unknown
Weight	64	diabetes	no		
Smoker	no	Origin	Asian		

Biochemical data

Parameter	Value	Corr. MoM
PAPP-A	3.99 mIU/mL	1.36
fb-hCG	45.27 ng/mL	0.98

Risks at sampling date

Age risk	1:951
Biochemical T21 risk	<1:10000
Combined trisomy 21 risk	1:8648
Trisomy 13/18 + NT	<1:10000



Ultrasound data

Gestational age	12 + 0
Method	CRL Robinson
Scan date	10-02-2024
Crown rump length in mm	56
Nuchal translucency MoM	1.49
Nasal bone	present
Sonographer	N A
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 8648 women with the same data, there is one woman with a trisomy 21 pregnancy and 8647 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