SAGEPATH LABS PVT LTD.,

Prisca 5.1.0.17

Date of report: 27/09/23

Patient data

Name	Mrs. MANISHA KUL	Patient ID	0662309250374
Birthday	05/10/96	Sample ID	24745332
Age at sample date	27.0	Sample Date	24/09/23
Gestational age	13 + 6		

Correction factors

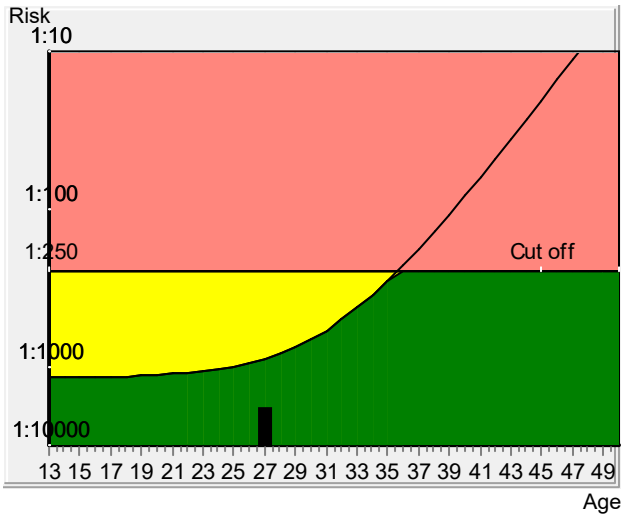
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	unknown
Weight	59	diabetes	no		
Smoker	no	Origin	Asian		

Biochemical data

Parameter	Value	Corr. MoM
PAPP-A	4.64 mIU/mL	0.71
fb-hCG	35.65 ng/mL	1.22

Risks at sampling date

Age risk	1:900
Biochemical T21 risk	1:1623
Combined trisomy 21 risk	1:9087
Trisomy 13/18 + NT	<1:10000



Trisomy 13/18 + NT

The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.

Ultrasound data

Gestational age	13 + 1
Method	CRL Robinson
Scan date	19/09/23
Crown rump length in mm	71
Nuchal translucency MoM	0.68
Nasal bone	unknown
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 9087 women with the same data, there is one woman with a trisomy 21 pregnancy and 9086 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