SAGEPATH LABS PVT LTD.,

Prisca 5.1.0.17

Date of report: 05/06/23

Patient data

Name	Mrs. POOJA KUMARI	Patient ID	0532306020007
Birthday	16/03/96	Sample ID	A1235423
Age at sample date	27.2	Sample Date	02/06/23
Gestational age	12 + 2		

Correction factors

Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	unknown
Weight	53.5	diabetes	no		
Smoker	no	Origin	Asian		

Biochemical data

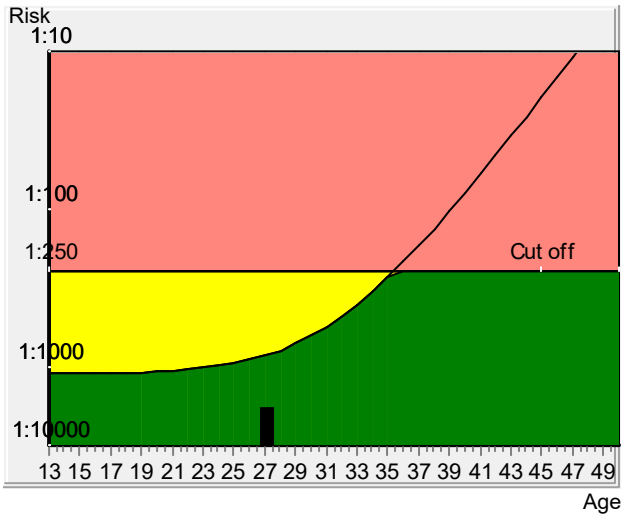
Parameter	Value	Corr. MoM
PAPP-A	2.3 mIU/mL	0.56
fb-hCG	45.65 ng/mL	0.99

Ultrasound data

Gestational age	12 + 2
Method	CRL Robinson
Scan date	02/06/23
Crown rump length in mm	59.5
Nuchal translucency MoM	0.84
Nasal bone	unknown
Sonographer	NA
Qualifications in measuring NT	MD

Risks at sampling date

Age risk	1:839
Biochemical T21 risk	1:1347
Combined trisomy 21 risk	1:7617
Trisomy 13/18 + NT	<1:10000



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 7617 women with the same data, there is one woman with a trisomy 21 pregnancy and 7616 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