Prisca 5.1.0.17

Date of report: 02-04-2022

Patient data

Name	Mrs. SANGITA CHANDRAPRAKASH SA	Patient ID	0352204010067
Birthday	25-03-1994	Sample ID	23288484
Age at sample date	28.0	Sample Date	01-04-2022
Gestational age	11 + 4		

Correction factors

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

Biochemical data

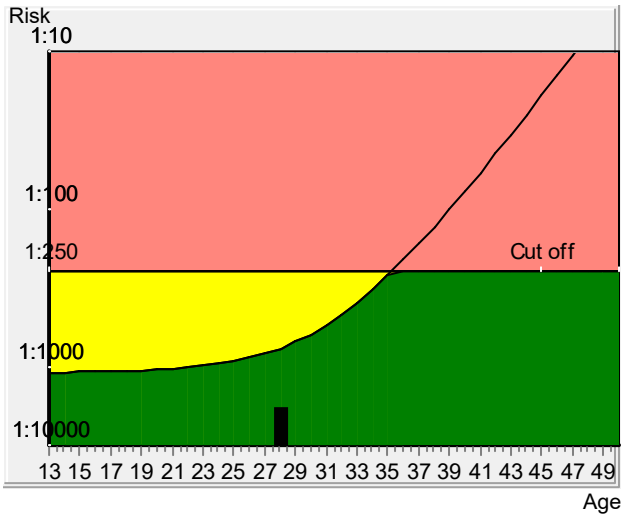
Parameter	Value	Corr. MoM
PAPP-A	3.01 mIU/mL	1.13
fb-hCG	56.87 ng/mL	1.10

Risks at sampling date

Age risk	1:766
Biochemical T21 risk	1:5001
Combined trisomy 21 risk	<1:10000
Trisomy 13/18 + NT	<1:10000

Ultrasound data

Gestational age	11 + 2
Method	CRL Robinson
Scan date	30-03-2022
Crown rump length in mm	46.4
Nuchal translucency MoM	0.94
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 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