Prisca 5.2.0.13

Date of report: **21/12/2024**

Patient data

Name	MRS. SHEETAL	Patient ID	
Birthday	15/06/1994	Sample ID	2412034205/NOD
Age at sample date	30.5	Sample Date	20/12/2024
Gestational age	12 + 5		

Correction factors

Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	no
Weight	59.5	diabetes	no		
Smoker	no	Origin	Asian		

Biochemical data

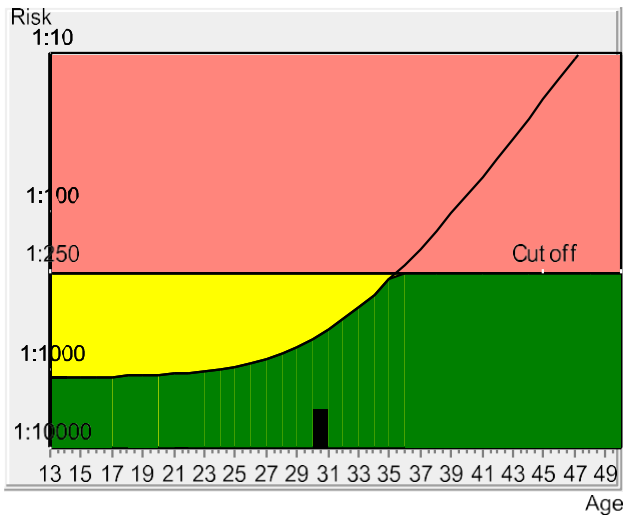
Parameter	Value	Corr. MoM
PAPP-A	3.35 mIU/ml	0.76
fb-hCG	29.5 ng/ml	0.67

Risks at sampling date

Age risk	1:608
Biochemical T21 risk	1:4904
Combined trisomy 21 risk	1:4549
Trisomy 13/18 + NT	<1:10000

Ultrasound data

Gestational age	12 + 3
Method	CRL Robinson
Scan date	18/12/2024
Crown rump length in mm	61
Nuchal translucency MoM	1.45
Nasal bone	present
Sonographer	..
Qualifications in measuring NT	..



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 4549 women with the same data, there is one woman with a trisomy 21 pregnancy and 4548 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