JITM SKILL PVT LTD

Prisca 5.2.0.13

Date of report: 7/07/2023

Patient data

Name	MRS. REETA	Patient ID	
Birthday	7/07/1989	Sample ID	2307010299/NOD
Age at sample date	34.0	Sample Date	6/07/2023

Gestational age 12 + 4

Correction factors

Fetuses	1	IVF	no	Previous trisomy 21	no
Weight	77	diabetes	no	pregnancies	
Smoker	no	Origin	Asian		

Biochemical data

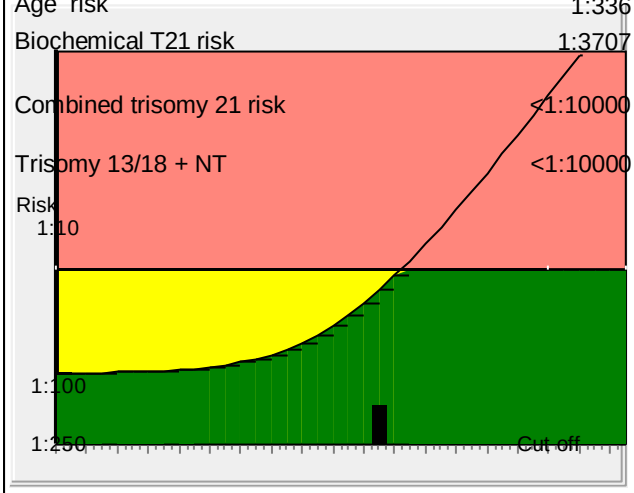
Parameter	Value	Corr. MoM
PAPP-A	2.5 mIU/ml	0.83
fb-hCG	26.1 ng/ml	0.63

Ultrasound data

Gestational age	12 + 4
Method	CRL Robinson
Scan date	6/07/2023
Crown rump length in mm	63
Nuchal translucency MoM	0.62
Nasal bone	present
Sonographer	DR. A.K. SURYAVANSH
Qualifications in measuring NT	MD

Risks at sampling date

Age risk	1:336
Biochemical T21 risk	1:3707
Combined trisomy 21 risk	<1:10000
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 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.