Prisca 5.2.0.13

Date of report: 17/03/25

JITM Diagnostics

Patient data			
Name	MRS. SUMAN	Patient ID	
Birthday	28/01/86	Sample ID	2503032398/NOD
Age at sample date	39.1	Sample Date	17/03/25
Gestational age	12 + 6		

Correction factors					
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	no
Weight	72	diabetes	no		
Smoker	no	Origin	Asian		

Biochemical data

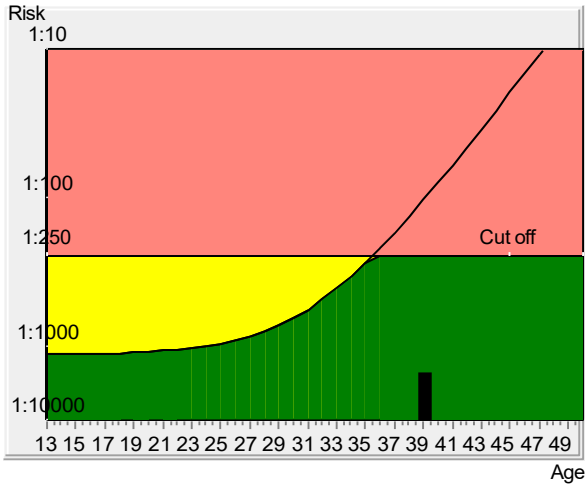
Parameter	Value	Corr. MoM
PAPP-A	3.73 mIU/ml	1.01
fb-hCG	57.1 ng/ml	1.40

Risks at sampling date

Age risk	1:101
Biochemical T21 risk	1:294
Combined trisomy 21 risk	1:1516
Trisomy 13/18 + NT	<1:10000

Ultrasound data

Gestational age	12 + 6
Method	CRL Robinson
Scan date	17/03/25
Crown rump length in mm	67
Nuchal translucency MoM	0.82
Nasal bone	present
Sonographer	DR. AMIT RAI
Qualifications in measuring NT	MD



Trisomy 13/18 + NT

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

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 1516 women with the same data, there is one woman with a trisomy 21 pregnancy and 1515 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