Prisca 5.2.0.13

Date of report: 3/12/2024

JITM Diagnostics

Patient data			
Name	MRS. HIMANSHI SINGHLA	Patient ID	
Birthday	6/08/1994	Sample ID	2412001879/NOD
Age at sample date	30.3	Sample Date	2/12/2024
Gestational age	13 + 3		

Correction factors					
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	no
Weight	79.4	diabetes	no		
Smoker	no	Origin	Asian		

Biochemical data

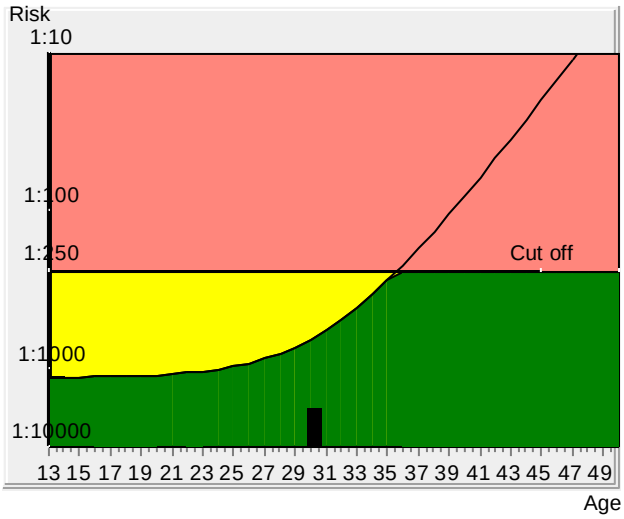
Parameter	Value	Corr. MoM
PAPP-A	2.85 mIU/ml	0.70
fb-hCG	12.3 ng/ml	0.33

Risks at sampling date

Age risk	1:640
Biochemical T21 risk	<1:10000
Combined trisomy 21 risk	<1:10000
Trisomy 13/18 + NT	<1:10000

Ultrasound data

Gestational age	13 + 3
Method	CRL Robinson
Scan date	2/12/2024
Crown rump length in mm	75.7
Nuchal translucency MoM	0.70
Nasal bone	present
Sonographer	DR. NEERJA CHOPRA
Qualifications in measuring NT	M.D



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 free beta HCG level is low.

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