Prisca 5.1.0.17

Date of report: 13/08/24

KOLTE SIR

Patient data

Name	Mrs. KOMAL MAHAJAN	Patient ID	0662408120169
Birthday	05/04/97	Sample ID	A0504919
Age at sample date	27.4	Sample Date	12/08/24
Gestational age	12 + 0		

Correction factors

Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	unknown
Weight	83	diabetes	no		
Smoker	no	Origin	Asian		

Biochemical data

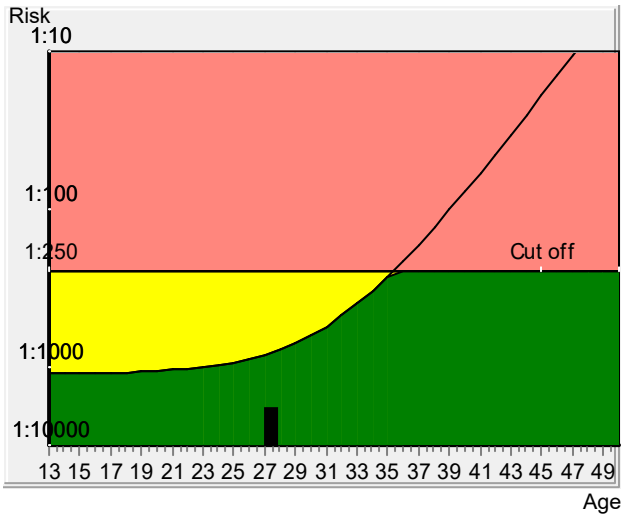
Parameter	Value	Corr. MoM
PAPP-A	7.95 mIU/mL	3.72
fb-hCG	49.32 ng/mL	1.15

Risks at sampling date

Age risk	1:822
Biochemical T21 risk	<1:10000
Combined trisomy 21 risk	1:4225
Trisomy 13/18 + NT	<1:10000

Ultrasound data

Gestational age	11 + 5
Method	CRL Robinson
Scan date	10/08/24
Crown rump length in mm	53
Nuchal translucency MoM	1.70
Nasal bone	present
Sonographer	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 4225 women with the same data, there is one woman with a trisomy 21 pregnancy and 4224 women with not affected pregnancies.
The PAPP-A level is high.
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