Prisca 5.1.0.17

Date of report: 20-11-2023

SHILPAKALA

Patient data			
Name	Mrs. KEERTHANA	Patient ID	0012311190411
Birthday	14-03-1999	Sample ID	24668762
Age at sample date	24.7	Sample Date	19-11-2023
Gestational age	13 + 5		

Correction factors					
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	unknown
Weight	38	diabetes	no		
Smoker	no	Origin	Asian		

Biochemical data

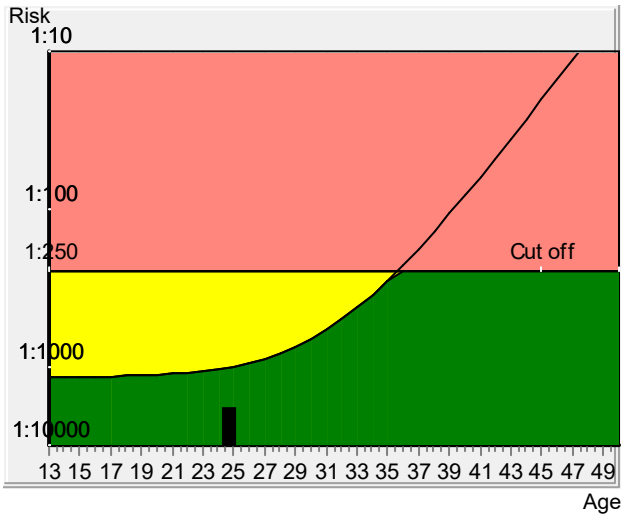
Parameter	Value	Corr. MoM
PAPP-A	4.98 mIU/mL	0.48
fb-hCG	29.22 ng/mL	0.81

Risks at sampling date

Age risk	1:1013
Biochemical T21 risk	1:1710
Combined trisomy 21 risk	1:4012
Trisomy 13/18 + NT	<1:10000

Ultrasound data

Gestational age	13 + 4
Method	CRL Robinson
Scan date	18-11-2023
Crown rump length in mm	78
Nuchal translucency MoM	1.21
Nasal bone	unknown
Sonographer	SHILPAKALA
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 4012 women with the same data, there is one woman with a trisomy 21 pregnancy and 4011 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

below cut off | Below Cut Off, but above Age Risk | above cut off