KOS DIAGNOSTIC LAB

6349/1, NICHOLSON ROAD, AMBALA CANTT

Prisca 5.0.2.37

Date of report: 24-09-2020

Patient data

Name	MRS. ARPANA	Patient ID	2009220845/AMB
Birthday	03-05-1995	Sample ID	2009220845/AMB
Age at delivery	25.9	Sample Date	23-09-2020
Gestational age	11 + 5		

Correction factors

Fetuses	1	IVF	yes	Previous trisomy 21 pregancies	no
Weight	51.5	diabetes	no		
Smoker	no	Origin	Asian		

Biochemical data

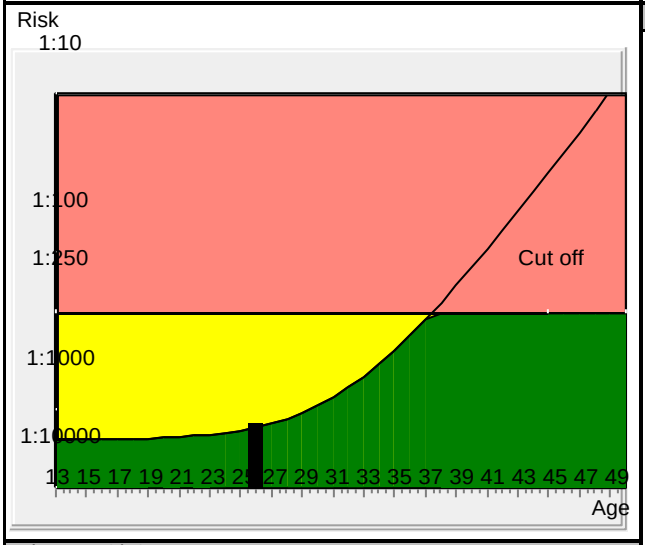
Parameter	Value	Corr. MoM
PAPP-A	1.78 mIU/ml	0.55
fb-hCG	92.2 ng/ml	1.83

Risks at term

Age risk	1:1324
Biochemical T21 risk	1:478
Combined trisomy 21 risk	1:1240
Trisomy 13/18 + NT	<1:10000

Ultrasound data

Gestational age	11 + 2
Method	CRL Robinson
Scan date	20-09-2020
Crown rump length in mm	46.8
Nuchal translucency MoM	1.17
Nasal bone	present
Sonographer	.
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 1240 women with the same data, there is one woman with a trisomy 21 pregnancy and 1239 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