KOS DIAGNOSTIC LAB

6349/1, NICHOLSON ROAD, AMBALA CANTT

Prisca 5.0.2.37

Date of report: 05-12-2018

Patient data

Name	MRS. PARMINDER KAUR	Patient ID	
Birthday	12-04-1993	Sample ID	1804220693/AMB
Age at sample date	25	Sample Date	04-12-2018
Gestational age	13 + 1		

Correction factors

Fetuses	1	IVF	unknown	Previous trisomy 21 pregancies	unknown
Weight	89	diabetes	unknown		
Smoker	unknown	Origin	Asian		

Biochemical data

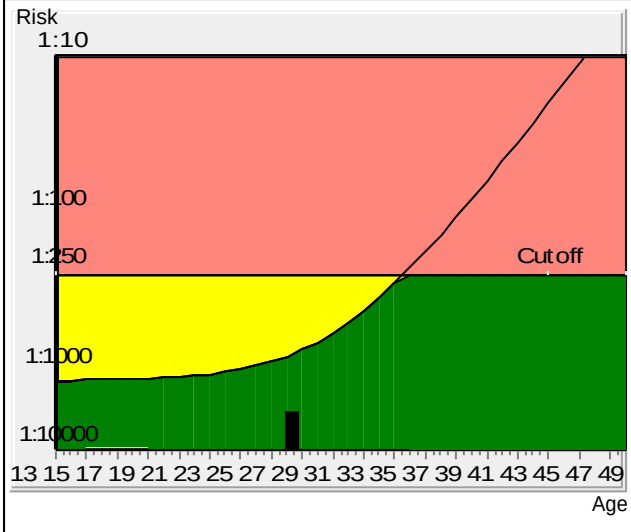
Parameter	Value	Corr. MoM
PAPP-A	2.8 mIU/ml	0.69
fb-hCG	35.1 ng/ml	1.17

Risks at sampling date

Age risk	1:790
Biochemical T21 risk	1:1460
Combined trisomy 21 risk	1:2641
Trisomy 13/18 + NT	<1:10000

Ultrasound data

Gestational age	13 + 1
Method	CRL Robinson
Scan date	04-12-2018
Crown rump length in mm	74
Nuchal translucency MoM	1.27
Nasal bone	
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 2641 women with the same data, there is one woman with a trisomy 21 pregnancy and 2640 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