KOS DIAGNOSTIC LAB

6349/1, NICHOLSON ROAD, AMBALA CANTT

Prisca 5.0.2.37

Date of report: 26-10-2018

Patient data			
Name	MRS. MANJEET	Patient ID	1806220428/AMB
Birthday	21-04-1985	Sample ID	1806220428/AMB
Age at sample date	32.0	Sample Date	26-10-2018

Gestational age	12 + 5	
Correction factors		
Fetuses 1	IVF no	Previous trisomy 21 no

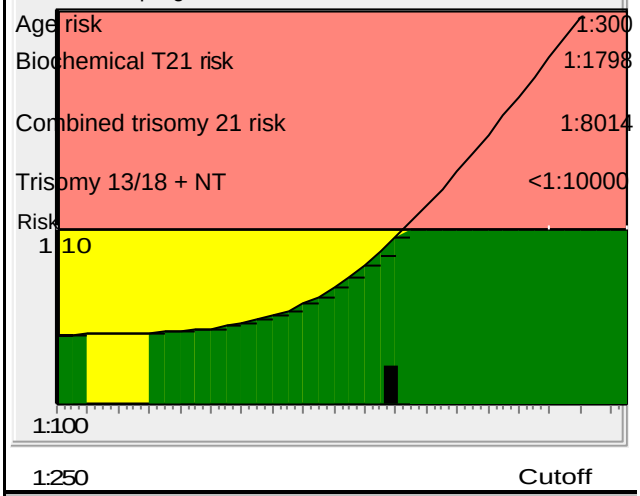
Weight	79	diabetes	no
Smoker	no	Origin	Asian

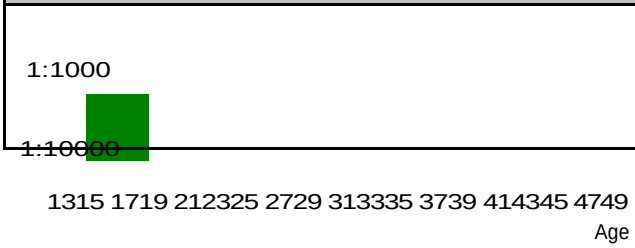
Biochemical data		
Parameter	Value	Corr. MoM
PAPP-A	13.4 mIU/ml	3.78
fb-hCG	50.8 ng/ml	1.75

Ultrasound data	
Gestational age	12 + 5
Method	CRL Robinson
Scan date	26-10-2018
Crown rump length in mm	62.41
Nuchal translucency MoM	0.85
Nasal bone	present
Sonographer	.
Qualifications in measuring NT	MD

Risks at sampling date

Age risk	1:300
Biochemical T21 risk	1:1798
Combined trisomy 21 risk	1:8014
Trisomy 13/18 + NT	<1:10000





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 8014 women with the same data, there is one woman with a trisomy 21 pregnancy and 8013 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 NTmeasurement 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!

below risk of Trisomy 13/18 + NT | Below Cut Off, but above Age Risk | above cut off

The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low

risk.

Sign of Physician