KOS DIAGNOSTIC LAB

6349/1, NICHOLSON ROAD,AMBALA CANTT

Date of report: 18-11-2016

Patient data

Name	MRS. JASMINE KAUR	Patient ID	1611220282/AMB
Birthday	29-07-1988	Sample ID	1611220282/AMB
Age at sample date	28.3	Sample Date	17-11-2016
Gestational age	12 + 4		

Correction factors

Fetuses	1	IVF	no	Previous trisomy 21	no
Weight	55	diabetes	no	pregancies	
Smoker	no	Origin	Asian		

Biochemical data

Parameter	Value	Corr. MoM
PAPP-A	9.8 mIU/ml	2.51
fb-hCG	105 ng/ml	2.91

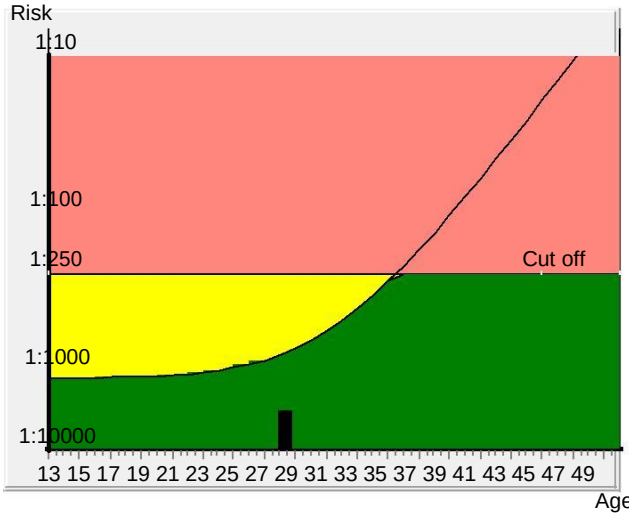
Risks at sampling date

Age risk	1:775
Biochemical T21 risk	1:1233
Combined trisomy 21 risk	1:3785
Trisomy 13/18 + NT	<1:10000

Ultrasound data

Gestational age	12 + 3
Method	CRL Robinson
Scan date	16-11-2016
Crown rump length in mm	62
Nuchal translucency MoM	1.06
Nasal bone	present
Sonographer	DR. NIHA MD
Qualifications in measuring NT	MD

Risk



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 3785 women with the same data, there is one woman with a trisomy 21 pregnancy and 3784 women with not affected pregnancies.
The free beta HCG level is high.
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