KOS DIAGNOSTIC LAB…

6349/1, NICHOLSON ROAD, AMBALA CANTT

Date of report: 15-10-2016

Patient data

Name	MRS. MANISHA	Patient ID	1610220215/AMB
Birthday	17-10-1989	Sample ID	1610220215/AMB
Age at sample date	27.0	Sample Date	13-10-2016
Gestational age	13 + 2		

Correction factors

Fetuses	1	IVF	no	Previous trisomy 21	no
Weight	52	diabetes	no	pregancies	
Smoker	no	Origin	Asian		

Biochemical data

Parameter	Value	Corr. MoM
PAPP-A	4.6 mIU/ml	0.85
fb-hCG	19.8 ng/ml	0.61

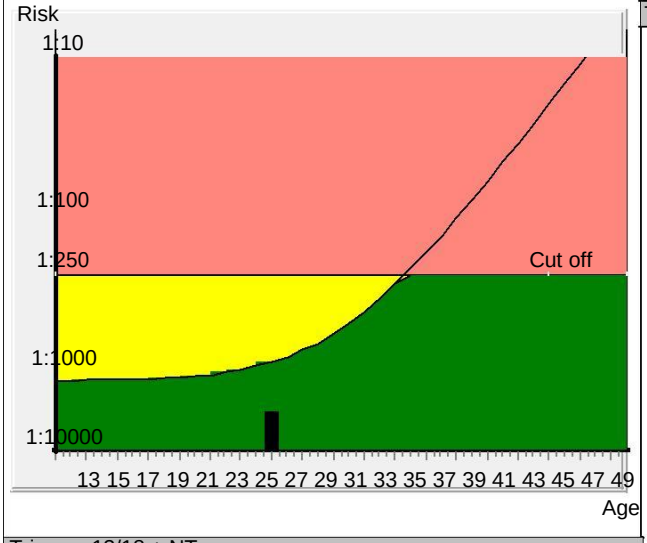
Risks at sampling date

Age risk	1:882
Biochemical T21 risk	<1:10000
Combined trisomy 21 risk	<1:10000
Trisomy 13/18 + NT	<1:10000

Ultrasound data

Gestational age	13 + 2
Method	CRL Robinson
Scan date	13-10-2016
Crown rump length in mm	74.8
Nuchal translucency MoM	0.62
Nasal bone	present
Sonographer	DR. POONAM LOOMBA
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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy.

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