KOS DIAGNOSTIC LAB

6349/1, NICHOLSON ROAD, AMBALA CANTT

Prisca 5.0.2.37

Date of report: 13/04/17

Patient data

Name	MRS. SHURUTI	Patient ID	
Birthday	23/05/91	Sample ID	17040220280/AMB
Age at sample date	25.9	Sample Date	12/04/17
Gestational age	11 + 2		

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	2.5 mIU/ml	1.11
fb-hCG	45.1 ng/ml	1.00

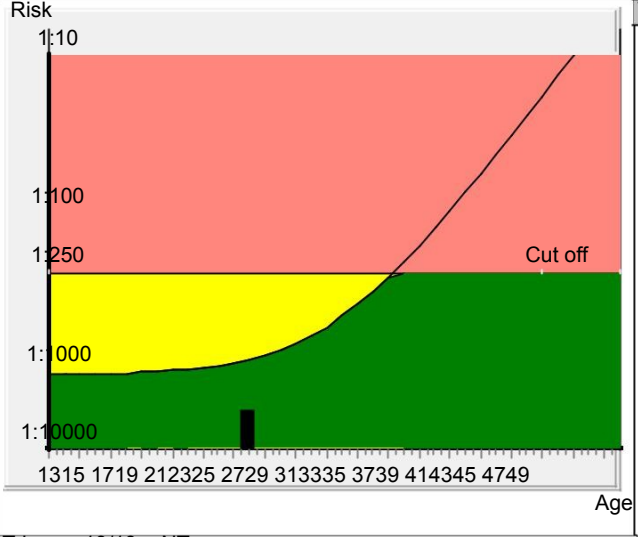
Risks at sampling date

Age risk	1:878
Biochemical T21 risk	1:6873
Combined trisomy 21 risk	<1:10000
Trisomy 13/18 + NT	<1:10000

Ultrasound data

Gestational age	11 + 2
Method	CRL Robinson
Scan date	12/04/17
Crown rump length in mm	47.5
Nuchal translucency MoM	0.75
Nasal bone	present
Sonographer	.
Qualifications in measuring NT	MD

Risk



Trisomy 13/18 + NT

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

Sign of Physician

below cut off | Below Cut-Off, but above Age Risk | above cut off