KOS DIAGNOSTIC LAB

6349/1, NICHOLSON ROAD, AMBALA CANTT

Prisca 5.0.2.37

Date of report: 22-04-2019

Patient data

Name	MRS. HARINDER	Patient ID	1904220769/AMB
Birthday	23-08-1982	Sample ID	1904220769/AMB
Age at sample date	36.7	Sample Date	20-04-2019
Gestational age	11 + 4		

Correction factors

Fetuses	1	IVF	no	Previous trisomy 21 pregancies	unknown
Weight	56	diabetes	no		
Smoker	no	Origin	Asian		

Biochemical data

Parameter	Value	Corr. MoM
PAPP-A	2.4 mIU/ml	0.95
fb-hCG	65.1 ng/ml	1.37

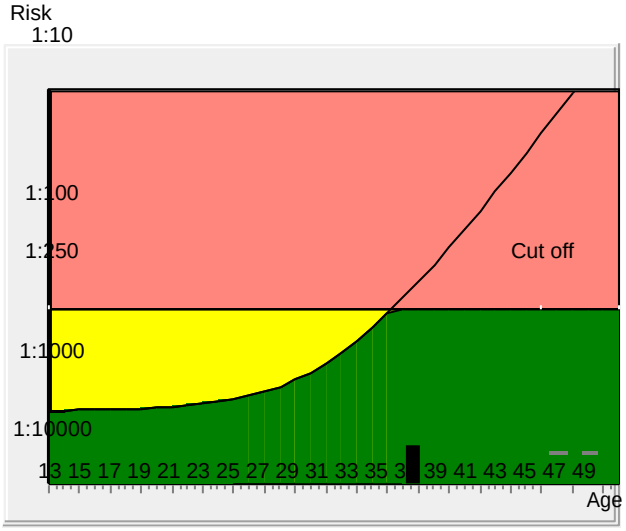
Ultrasound data

Gestational age	11 + 1
Method	CRL Robinson
Scan date	17-04-2019
Crown rump length in mm	46
Nuchal translucency MoM	0.95
Nasal bone	unknown
Sonographer	.
Qualifications in measuring NT	MD

Risks at sampling date

Age risk	1:179
Biochemical T21 risk	1:489
Combined trisomy 21 risk	1:2167
Trisomy 13/18 + NT	<1:10000

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 2167 women with the same data, there is one woman with a trisomy 21 pregnancy and 2166 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