KOS DIAGNOSTIC LAB

6349/1, NICHOLSON ROAD, AMBALA CANTT

Prisca 5.0.2.37

Date of report: 14-05-2019

Patient data			
Name	MANMEET KAUR	Patient ID	
Birthday	26-03-1991	Sample ID	211905110001
Age at sample date	28.0	Sample Date	11-05-2019
Gestational age	13 + 3		

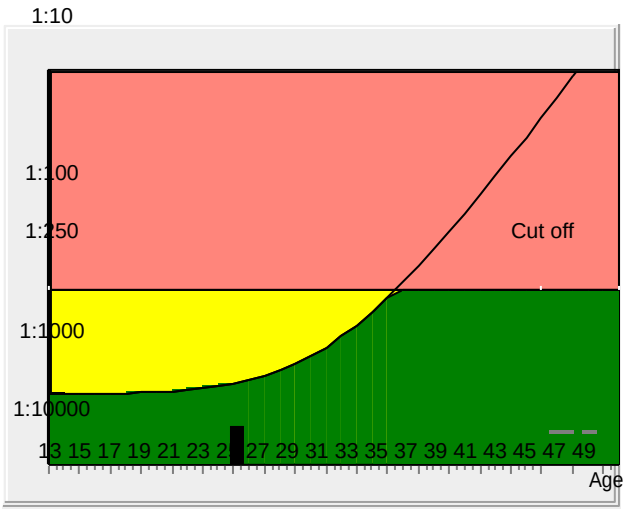
Correction factors					
Fetuses	1	IVF	no	Previous trisomy 21 pregancies	no
Weight	54	diabetes	no		
Smoker	no	Origin	Asian		

Biochemical data		
Parameter	Value	Corr. MoM
PAPP-A	8.60 mIU/ml	1.57
fb-hCG	47.0 ng/ml	1.33

Risks at sampling date	
Age risk	1:960
Biochemical T21 risk	1:7570
Combined trisomy 21 risk	<1:10000
Trisomy 13/18 + NT	<1:10000

Ultrasound data	
Gestational age	13 + 3
Method	CRL Robinson
Scan date	07-05-2019
Crown rump length in mm	67.52
Nuchal translucency MoM	0.66
Nasal bone	present
Sonographer	.
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