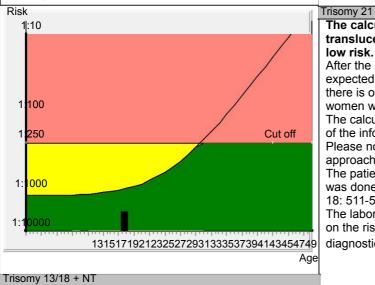
## KOS DIAGNOSTIC LAB 6349/1, NICHOLSON ROAD, AMBALA CANTT

Prisca 5.0.2.37

Date of report: 25-04-2017

Patient data						
Name	MRS. GURJEET			Patient ID		17040220611/AMB
Birthday	05-08-1991			Sample ID		17040220611/AMB
Age at sample date	25.7			Sample Date		24-04-2017
Gestational age			13 + 1			
Correction factors						
Fetuses	1	IVF		no	Previous trisomy 21	no
Weight	48	diabetes		no	pregancies	
Smoker	no	Origin		Asian		
Biochemical data			Ultrasound data			
Parameter	Value Corr. Mo		Corr. MoM	Gestational age		12 + 5
PAPP-A 5	5.4 mIU/mI 0.9		0.96	Method		CRL Robinson
fb-hCG 67	67.1 ng/ml 1.96		Scan date		21-04-2017	
Risks at sampling date				Crown rump length in mm		65.2
Age risk	1:948			Nuchal translucency MoM		0.68
Biochemical T21 risk			1:1091	Nasal bone		present
Combined trisomy 21 risk 1:6024			Sonographer			
Trisomy 13/18 + NT			<1:10000	Qualification	s in measuring NT	MD



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

## 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 6024 women with the same data, there is one woman with a trisomy 21 pregnancy and 6023 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!

## Sign of Physician