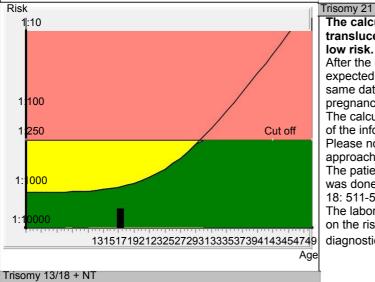
KOS DIAGNOSTIC LAB 6349/1, NICHOLSON ROAD, AMBALA CANTT

Prisca 5.0.2.37

Date of report: 29-04-2017

Patient data						
Name	MRS. NANCY GARG			Patient ID		1704220747/AMB
Birthday	11-02-1992			Sample ID		1704220747/AMB
Age at sample date	25.2			Sample Date		28-04-2017
Gestational age			13 + 0			
Correction factors						
Fetuses	1	IVF		no	Previous trisomy 21	no
Weight	56	diabetes		no	pregancies	
Smoker	no	Origin		Asian		
Biochemical data				Ultrasound da	ata	
Parameter Va	Value Corr		Corr. MoM	Gestational age		13 + 0
PAPP-A 9.5	9.5 mIU/ml		2.12	Method		CRL Robinson
fb-hCG 42.2	42.2 ng/ml		1.27	Scan date		28-04-2017
Risks at sampling date				Crown rump length in mm		69.3
Age risk 1:968			Nuchal translucency MoM		0.45	
Biochemical T21 risk <1:10000			Nasal bone		present	
Combined trisomy 21 risk <1:10000				Sonographer		
Trisomy 13/18 + NT			<1:10000	Qualification	s in measuring NT	MD



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

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

risk.