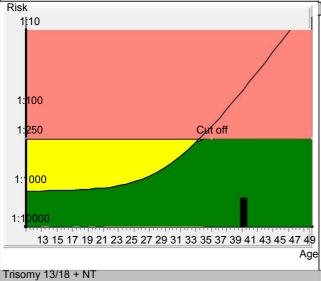
KOS DIAGNOSTIC LAB 6349/1, NICHOLSON ROAD, AMBALA CANTT

Prisca 5.0.2.37

Date of report: 09-03-2017

Patient data						
Name	DR. DEEPTI MANOCHA			Patient ID		1703220202/AMB
Birthday	13-12-1975			Sample ID		1703220202/AMB
Age at sample date	41.2			Sample Date		08-03-2017
Gestational age	12 + 3					
Correction factors	Î					
Fetuses	1	IVF		no	Previous trisomy 21	no
Weight	87	diabetes		no	pregancies	
Smoker	no	Origin		Asian		
Biochemical data				Ultrasound data		
Parameter	Value Corr. MoM		Gestational age		12 + 3	
PAPP-A	2.2 mIU/ml 1.04		Method		CRL Robinson	
fb-hCG	37.5 ng/ml 1.1		1.16	Scan date		08-03-2017
Risks at sampling date				Crown rump length in mm		61.9
Age risk	ge risk 1:56			Nuchal translucency MoM		0.71
Biochemical T21 risk 1:270			Nasal bone		present	
Combined trisomy 21 risk 1:1361			Sonographer		DR. POONAM LOOMBA	
Trisomy 13/18 + NT <1:10000			Qualifications in measuring NT		MD	



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

After the result of the Trisomy 21 test (with NT) it is expected that among 1361 women with the same data, there is one woman with a trisomy 21 pregnancy and 1360 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