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

5.1.0.17

Date of report: 10-03-2021

Prisca

Patient data				
Name	MRS. UPMA			
Birthday	07-02-1985	Sample ID 2103220316/AM		2103220316/AMB
Age at sample date	36.1	Sample Date		08-03-2021
Gestational age	13 + 2			
Correction factors				
Fetuses 1	IVF	no	Previous trisomy 21	no
Weight 71.2	diabetes	no	pregnancies	
Smoker no	Origin	Asian		
Biochemical data		Ultrasound data		
Parameter Value	Corr. MoM	Gestational age 13 + 1		
PAPP-A 9.4 mIU/m	l 2.12	Method CRL Robinson		
fb-hCG 28.1 ng/ml	0.72	Scan date 07-03-2021		
Risks at sampling date	5		Crown rump length in mm 70.87	
Age risk	1:219		Nuchal translucency MoM 0.60	
Biochemical T21 risk	<1:10000		Nasal bone pre	
Combined trisomy 21 risk <1:10000		Sonographer		
Trisomy 13/18 + NT	• NT <1:10000		Qualifications in measuring NT M	
Risk 1:10		Trisomy 21	ated risk for Trisomy 21	(with pucked
1: 00 1:250 Cut off 1:10000 1:100000 1:10000 1:10000 1:10000 1:100000 1:100000 1:100000		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