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

5.1.0.17

Date of report: 16-03-2022

Prisca

Patient data				
Mame MRS. AMANPREET KAUR		Patient ID		2203220443/AMB
Birthday	26-05-1997	Sample ID		2203220443/AMB
Age at sample date	24.8	Sample Date	)	15-03-2022
Gestational age	13 + 1			
Correction factors				
Fetuses 1	IVF	no	Previous trisomy 21	no
Weight 67	diabetes	no	pregnancies	
Smoker no	Origin	Asian		
Biochemical data		Ultrasound data		
Parameter Value	Corr. MoM	Gestational age 12 + 2		
PAPP-A 1.8 mIU/m	I 0.40	Method CRL Robinson		
fb-hCG 10.3 ng/ml	0.26	Scan date 09-03-2022		
Risks at sampling date		Crown rump	length in mm	59.68
Age risk	1:990	Nuchal translucency MoM		1.09
Biochemical T21 risk	1:5560	Nasal bone present		
Combined trisomy 21 risk	<1:10000	Sonographe	r	
Trisomy 13/18 + NT	1:1829	Qualifications in measuring NT MD		
Risk 1:10	Trisomy 21 The calculated risk for Trisomy 21 (with nuchal			
1: 00 1:250 1:1000 1:1000 1:100000 1:100000 1:100000 1:1000	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 free beta HCG level is low. The PAPP-A level is low. 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