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

5.0.2.37

Date of report: 10-04-2019

Prisca

Patient data				
Name	MRS. MANPREET			211904090003
Birthday	10-01-1997	Sample ID		211904090003
Age at sample date	22.0	Sample Date	e	09-04-2019
Gestational age	11 + 2			
Correction factors				
Fetuses 1	IVF	no	Previous trisomy 21	no
Weight 76	diabetes	no	pregancies	
Smoker no	Origin	Asian		
Biochemical data		Ultrasound da	ata	
Parameter Value	Corr. MoM	Gestational a	age	11 + 2
PAPP-A 2.92 mIU/m	l 1.24	Method LMP		
fb-hCG 36.6 ng/ml	0.84	Scan date		
Risks at sampling date	ate		Crown rump length in mm	
Age risk	1:900	Nuchal translucency MoM		
Biochemical T21 risk	<1:10000	Nasal bone present		
Combined trisomy 21 risk			er	
Trisomy 13/18 + NT			ns in measuring NT	MD
Risk 1:10	Trisomy 21 The calculated risk for Trisomy 21 (with nuchal			
1:100 1:250 1:1000 1:1000 1:1000 1:1000 1:1000 Trisomy 13/18 + NT The calculated risk for trisomy 13 translucency) is < 1:10000, which risk.	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