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

5.0.2.37

Date of report: 01-07-2019

Prisca

Patient data				
Name	MRS. SANDEEP	Patient ID		101906290012
Birthday	01-11-1990			101906290012
Age at sample date	28.0	Sample Date		29-06-2019
Gestational age	13 + 4			
Correction factors				
Fetuses 1	IVF	no	Previous trisomy 21	no
Weight 65	diabetes	no	pregancies	
Smoker no	Origin	Asian		
Biochemical data		Ultrasound da	ata	
Parameter Value	Corr. MoM	Gestational a	age	13 + 4
PAPP-A 2.36 mIU/m	nl 0.63	Method LMP		
fb-hCG 63.2 ng/ml	1.96	Scan date		
Risks at sampling date		Crown rump length in mm		
Age risk	1:900	Nuchal translucency MoM		
Biochemical T21 risk	1:410	Nasal bone		
Combined trisomy 21 risk	1:1545			
Trisomy 13/18 + NT	<1:10000	Qualifications in measuring NT ME Trisomy 21		
1: 000 1: 000 1: 000 1: 100 1: 000 1: 1: 000 1: 000 1: 000			The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicatesa low risk. After the result of the Trisomy 21 test (with NT) it is expected that among 1545 women with the same data, there is one woman with a trisomy 21 pregnancy and 1544 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