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

Prisca

5.1.0.17

Date of report: 22-02-2022

Patient data		•		
Name MRS. POONAM		Patient ID		
Birthday	31-03-1989	Sample ID		2202220456/AMB
Age at sample date	32.9	Sample Date		18-02-2022
Gestational age	11 + 3			
Correction factors				
Fetuses 1	IVF	yes	Previous trisomy 21	no
Weight 63.9	diabetes	no	pregnancies	
Smoker no	Origin	Asian		
Biochemical data		Ultrasound da	ita	
Parameter Value	Corr. MoM	Gestational age 11 + 3		
PAPP-A 3.37 mIU/m	l 1.54	Method CRL Robinson		
fb-hCG 56.4 ng/ml	1.13	Scan date 18-02-2022		
Risks at sampling date		Crown rump length in mm 49.		
Age risk	1:398	Nuchal translucency MoM		0.75
Biochemical T21 risk	1:4375			present
Combined trisomy 21 risk <1:10000		Sonographer		
Trisomy 13/18 + NT	<1:10000	Qualifications in measuring NT Trisomy 21		MD
1:10 1:10 1:100 1:250 1:1000 1:100000 1:10000 1:100000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000	The calculated risk for Trisomy 21 (with nuchal 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