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

Prisca 5.1.0.17

Date of report: 04-12-2021

Patient data				
Name	MRS. NIRU SHARMA			2112220050/AMB
Birthday	02-12-1988		Sample ID	
Age at sample date	date 33.0		Sample Date	
Gestational age	12 + 6			
Correction factors				
Fetuses 1	IVF	no	Previous trisomy 21	no
Weight 67.1	diabetes	no	pregnancies	
Smoker no	Origin	Asian		
Biochemical data		Ultrasound da	nta	
Parameter Value	Corr. MoM	Gestational age 12 + 2		
PAPP-A 2.12 mIU/m	nl 0.53	Method CRL Robinson		
fb-hCG 47.2 ng/ml	1.13	Scan date 29-11-2021		
Risks at sampling date			Crown rump length in mm	
Age risk			Nuchal translucency MoM  Nasal bone  pr	
Biochemical T21 risk	· · · · · · · · · · · · · · · · · · ·			preser
Combined trisomy 21 risk 1:2510		Sonographe		
Trisomy 13/18 + NT			Qualifications in measuring NT ME Trisomy 21	
1:1000 1:10000 1:10000 13 15 17 19 21 23 25 27 29 31 33 3 Trisomy 13/18 + NT The calculated risk for trisomy 13 translucency) is < 1:10000, which	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 2510 women with the same data, there is one woman with a trisomy 21 pregnancy and 2509 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

Below Cut Off, but above Age Risk

above cut off