KOS DIAGNOSTIC LAB 6349/1, NICHOLSON ROAD AMBALA CANTT

Patient data					
Name		MRS. MANDEEP (A)	Patient ID		2206220389/AMB (A)
Birthday		05-04-1999	Sample ID		2206220389/AMB (A)
Age at sample date		23.2	Sample Date		15-06-2022
Gestational age		12 + 1			
Correction factors					
Fetuses	2	IVF	no	Previous trisomy 21	no
Weight	45	diabetes	no	pregnancies	
Smoker	no	Origin	Asian		
Biochemical data			Ultrasound data	1	
Parameter	Value Corr. MoM		Gestational age 12 + 0		
PAPP-A	12.1 mIU/m	1.38	Method CRL Robinson		
fb-hCG	375 ng/ml	3.33	Scan date 14-06-2022		
Risks at sampling date	-		Crown rump length in mm		56.36
Age risk Biochemical	1:1011		Nuchal translucency MoM		0.65
T21 risk		1:594	Nasal bone		present
Combined trisomy 21 risk		1:3207	Sonographer		
Trisomy 13/18 + NT		<1:10000	Qualifications in measuring NT		MD
1:10 1:10 1:250 1:1000 1:1		Trisomy 21 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 3207 women with the same data, there is one woman with a trisomy 21 pregnancy and 3206women with not affected pregnancies. The free beta HCG level is high. The risk for this twin pregnancy has been calculated for asingletor pregnancy with corrected MoMs. The calculated risk by PRISCA depends on the accuracyof the information provided by the referring physician. Please note that risk calculations are statisticalapproaches and have no diagnostic value! The patient combined risk presumes the NT measurementwas done according to accepted guidelines (Prenat Diagn18: 511-523 (1998)). The laboratory can not be hold responsible for their impacton the risk assessment ! Calculated risks have no diagnostic value!			

Sign of Physician