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

5.1.0.17

Date of report: 14-04-2021

Prisca

Patient data				
Name	MRS. MOHINI DEVI	Patient ID		2104220595/AMB
Birthday	26-01-1994	Sample ID		2104220595/AMB
Age at sample date	27.2	Sample Date		13-04-2021
Gestational age	13 + 2			
Correction factors				
Fetuses 1	IVF	no	Previous trisomy 21	no
Weight 57	diabetes	no	pregnancies	
Smoker no	Origin	Asian		
Biochemical data		Ultrasound da	ata	
Parameter Value	Corr. MoM	Gestational age 12 + 6		
PAPP-A 7.49 mIU/m	l 1.29	Method CRL Robinson		
fb-hCG 87.6 ng/ml	2.09	Scan date 10-04-2021		
Risks at sampling date		Crown rump length in mm 67.4		
Age risk	1:869	Nuchal translucency MoM		0.59
Biochemical T21 risk	1:1545	Nasal bone		present
Combined trisomy 21 risk	1:7986	Sonographe	er	
Trisomy 13/18 + NT	<1:10000	Qualifications in measuring NT MI		MD
1:100 1:250 Cut off 1:1000 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 7986 women with the same data, there is one woman with a trisomy 21 pregnancy and 7985 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