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

Prisca 5.1.0.17

Date of report: 25-09-2021

Patient data				
Name	MRS. DIPA			2109220776/AMB
Birthday	30-04-1992			2109220776/AMB
Age at sample date	e date 29.4		Sample Date	
Gestational age 12 + 6				
Correction factors				
Fetuses 1	IVF	no	Previous trisomy 21	no
Weight 52	diabetes	no	pregnancies	
Smoker no	Origin	Asian		
Biochemical data		Ultrasound data		
Parameter Value	Corr. MoM	M Gestational age 12 + 3		
PAPP-A 3.48 mIU/m	nl 0.64	Method CRL Robinson		
fb-hCG 110.2 ng/ml	2.43	Scan date 21-09-2021		
Risks at sampling date	ks at sampling date		Crown rump length in mm 62	
Age risk	1:701	Nuchal trans	Nuchal translucency MoM 0.	
Biochemical T21 risk	1:184			present
Combined trisomy 21 risk 1:1138		Sonographer .		
Trisomy 13/18 + NT	<1:10000	Qualification	ns in measuring NT	MD
1:10 1:250 1:1000 1:10000 1:3 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 risk.	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 1138 women with the same data, there is one woman with a trisomy 21 pregnancy and 1137 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

above cut off