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

Prisca 5.0.2.37

Date of report: 17-12-2018

Patient data				
Name	MRS. RENU	Patient ID		211812170001
Birthday	22-09-1989	Sample ID		211812170001
Age at sample date	29	Sample Date	Э	14-06-2018
Gestational age	13+ 6			
Correction factors				
Fetuses 1	IVF	no	Previous trisomy 21	no
Weight 51.6	diabetes	no pregancies		
Smoker no	Origin	Asian		
Biochemical data		Ultrasound data		
Parameter Value	Corr. MoM	Gestational age 13 + 6		
PAPP-A 2.34 mIU/m	nl 0.63	Method CRL Robinson		
fb-hCG 64.2 ng/ml	1.96	Scan date 14-12-2018		
Risks at sampling date	at sampling date		Crown rump length in mm 78	
Age risk	1:900	Nuchal translucency MoM		1.04
Biochemical T21 risk	1:396	Nasal bone		present
Combined trisomy 21 risk	1:1545	Sonographe	er	
Trisomy 13/18 + NT	<1:10000	Qualifications in measuring NT MD		
Risk 1:10	Trisomy 21 The calculated risk for Trisomy 21 (with nuchal			
1:100 1:1000 1:1000 1:10000 1315 1719 212325 2729 31333 Trisomy 13/18 + NT The calculated risk for trisomy 13 translucency) is < 1:10000, which risk.	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 1545 women with the same data, there is one woman with a trisomy 21 pregnancy and 1544 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