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

Prisca 5.0.2.37

Date of report: 21-12-2018

Patient data				
Name	MRS. SEEMA			1812220279/AMB
Birthday	12-10-1992	Sample ID		1812220279/AMB
Age at sample date	26.2	Sample Date	9	20-12-2018
Gestational age	13 + 2			
Correction factors				
Fetuses 1	IVF	no Previous trisomy 21 no		
Weight 59.8	diabetes	no	pregancies	
Smoker no	Origin	Asian		
Biochemical data	T.		Ultrasound data	
Parameter Value	Value Corr. MoM (Gestational age 13 + 0	
PAPP-A 4.85 mIU/m	I 1.06	Method CRL Robinson		
fb-hCG 48.2 ng/ml	1.56	Scan date 18-12-2018		
Risks at sampling date	e		Crown rump length in mm 7	
Age risk	1:929	Nuchal translucency MoM 0.75		
Biochemical T21 risk	1:2310	Nasal bone present		
Combined trisomy 21 risk	ned trisomy 21 risk <1:10000		Sonographer	
Trisomy 13/18 + NT	<1:10000	Qualifications in measuring NT MD		
Risk 1:10	1	Trisomy 21	ated risk for Trisomy 21	
1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy. 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

Below Cut Off, but above Age Risk

above cut off