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

Prisca 5.0.2.37

Date of report: 14-12-2016

Patient data				
Name	MRS. BABLI		Patient ID	
Birthday	22-12-1981		Sample ID 1612220151/AMB	
Age at sample date	35.0	Sample Date 13-12-2016		
Gestational age	11 + 6			
Correction factors				
Fetuses 1	IVF	yes	Previous trisomy 21 no	
Weight 61	diabetes	unknown	pregancies	
Smoker no	Origin	Asian		
		Ultrasound data		
Parameter Value	Corr. MoM	Gestational	age 11 + 6	
PAPP-A 4.1 mIU/m		Method CRL Robinson		
fb-hCG 22.1 ng/ml	0.56	Scan date 13-12-2016		
Risks at sampling date		Crown rump length in mm 54.8		
Age risk	1:266	Nuchal translucency MoM 0.70		
Biochemical T21 risk	<1:10000	Nasal bone present		
Combined trisomy 21 risk	,		Sonographer .	
Trisomy 13/18 + NT			Qualifications in measuring NT MD	
I INDIN		Trisomy 21 The calculated risk for Trisomy 21 (with nuchal		
1:1000 1:10000 1:10000 1315 1719 21 23 25 27 29 31 33 33 Trisomy 13/18 + NT The calculated risk for trisomy 13 translucency) is < 1:10000, which risk.	Age //18 (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 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