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

Prisca 5.0.2.37

Date of report: 12/05/19

Patient data				
Name	MRS. BALJEET		Patient ID	
Birthday	02-08-1992		Sample ID	
Age at sample date	26.0		Sample Date	
Gestational age	12 + 5			
Correction factors		<u> </u>		
Fetuses 1	IVF	no	Previous trisomy 21	no
Weight 47	diabetes	no	pregancies	
Smoker no	Origin	Asian		
Biochemical data		Ultrasound d	ata	
Parameter Value	Corr. Mol	Gestational a	age	12 + 5
PAPP-A 1.9 mIU/m	nl 0.64			
fb-hCG 59.6 ng/ml	1.78	Scan date 10-05-19		
Risks at sampling date	e		Crown rump length in mm	
Age risk	1:959	Nuchal translucency MoM 0.6		0.69
Biochemical T21 risk	1:546			present
Combined trisomy 21 risk 1:3312		Sonographe	er	
Trisomy 13/18 + NT <1:10000		3		
1:10 1:100 1:250 1:1000 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.	Trisomy 21 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 3312 women with the same data, there is one woman with a trisomy 21 pregnancy and 3311 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

below cut off Below Cut Off, but above Age Risk

above cut off