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

Prisca 5.0.2.37

Date of report: 13-07-2017

Patient data				
Name	MRS. PARVINDER KAUR		Patient ID	1707220362/AMB
Birthday	21-09-1982		Sample ID	1707220362/AMB
Age at sample date	34.8		Sample Date	e 12-07-2017
Gestational age		11 + 6		
Correction factors				
Fetuses	1	IVF	no	Previous trisomy 21 no
Weight	58	diabetes	no	pregancies
Smoker	no	Origin	Asian	
Biochemical data			Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age 11 + 6	
PAPP-A	4.79 mIU/ml	1.75	Method CRL Robinson	
fb-hCG	122.5 ng/ml	3.05	Scan date	12-07-2017
Risks at sampling date			Crown rump length in mm 54.7	
Age risk		1:276	Nuchal tran	slucency MoM 0.77
Biochemical T21 risk		1:310	Nasal bone	present
Combined trisomy 21	risk	1:1565	, ,	
Trisomy 13/18 + NT		<1:10000		
Risk 1:10			Trisomy 21	
1:1000 1:250 1:1000 1:1000 13 15 17 19 21 23 25 Trisomy 13/18 + NT The calculated risk for translucency) is < 1: 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 1565 women with the same data, there is one woman with a trisomy 21 pregnancy and 1564 women with not affected pregnancies. The free beta HCG level is high. 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!	
			<u> </u>	Sign of Physician

Sign of Physician

below cut off Below Cut Off, but above Age Risk above cut off