KOS DIAGNOSTIC LAB

6349/1 NICHOLSON ROAD, AMBALA CANTT

5.1.0.17

Date of report: 28/08/22

Prisca

Patient data						
Name		MRS.	POONAM	Patient ID		2208220794/AMB
Birthday	01/11/85			Sample ID		2208220794/AMB
Age at sample date	date 36.8			Sample Date 27/08/2		
Gestational age			12 + 0			
Correction factors						
Fetuses	1	IVF		yes	Previous trisomy 21	nc
Weight	87.8	diabetes		no	pregnancies	
Smoker	no	Origin		Asian		
Biochemical data				Ultrasound data		
Parameter	Value	Value Corr. MoM (age	11 + 6
PAPP-A	1.82 mIU/m	nl	0.92	Method		CRL Robinsor
fb-hCG	29.2 ng/ml		0.68	Scan date 26/08/22		
Risks at sampling date			Crown rump length in mm 55			
Age risk	je risk 1:175			Nuchal translucency MoM 1.17		
Biochemical T21 risk				Nasal bone		present
Combined trisomy 21 risk 1:5264			Sonographer			
Trisomy 13/18 + NT <1:10000						
Risk 1:10				Trisomy 21 The calculated risk for Trisomy 21 (with nuchal		
1: 00 1: 50 Cut off 1: 1000 1: 1000 1: 1000 1: 10000 1: 10000 1: 10000 1: 10000 1: 10000 1: 10000 1: 1000 Age				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 5264 women with the same data, there is one woman with a trisomy 21 pregnancy and 5263 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!		
Trisomy 13/18 + NT The calculated risk translucency) is < 1 risk.						

above cut off