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

Prisca 5.0.2.37

Date of report: 08-05-2019

Patient data					
Name NEELAM			Patient ID		
Birthday	08-03-1993		Sample ID		211905070006
Age at sample date	at sample date 26		Sample Date 07-05-2019		
Gestational age 13 + 2					
Correction factors					
Fetuses 1	IVF		no	Previous trisomy 21	no
Weight 50	diabetes		no	pregancies	
Smoker no	Origin		Asian		
Biochemical data	-		Ultrasound da	ata	
Parameter Value		Corr. MoM	Gestational	age	13 + 2
PAPP-A 8.59 mIU/m	ıl 1.57 M		Method		CRL Robinson
fb-hCG 47.3 ng/ml	1.33 S		Scan date		06-05-2019
Risks at sampling date		Crown rump length in mm		69.69	
Age risk	1:970		Nuchal translucency MoM		0.66
Biochemical T21 risk			Nasal bone		present
•			Sonographer .		
			Qualifications in measuring NT MD Trisomy 21		
1:100 1:250 Cut off 1:10000 1:10000 13 15 17 19 21 23 2 27 29 31 33 35 37 39 41 43 45 47 49 Age Trisomy 13/18 + NT The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low 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 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