KOS DIAGNOSTIC LAB

AMBALA CANTT

Prisca 5.2.0.13

Date of report: 25/10/2024

Patient data				
Name	MRS. MARIYAM			
Birthday	1/11/1993	Sample ID		2410220542/AMB
Age at sample date	31.0	Sample Date		24/10/2024
Gestational age	13 + 2			
Correction factors				
Fetuses 1	IVF	no	Previous trisomy 21	no
Weight 64	diabetes	no	pregnancies	
Smoker no	Origin	Asian		
Biochemical data		Ultrasound data		
Parameter Value	Corr. MoM	Gestational age 11 +		
PAPP-A 2.4 mIU/r	nl 0.48	Method CRL Robin		CRL Robinson
fb-hCG 14.8 ng/m	0.37	Scan date		10/10/2024
ks at sampling date		Crown rump length in mm		47.8
Age risk	1:582	Nuchal translucency MoM		1.00
Biochemical T21 risk	1:4466	Nasal bone		present
Combined trisomy 21 risk	<1:10000	• '		DR. INDERJIT SINGH
Trisomy 13/18 + NT	1:7613	Qualifications in measuring NT Trisomy 21		M.D
1:10 1:1000 1:100000 1:100000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:100	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 free beta HCG level is low. 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!			

Below Cut Off, but above Age Risk

below cut off

Sign of Physician

above cut off