Date of report: 18/02/2025

JITM Diagnostics

Patient data				
Name	MRS. KRISHNA	Patient ID		
Birthday	7/09/1985			2502031864/NOD
Age at sample date	39.4	Sample Date	•	17/02/2025
Gestational age	12 + 3			
Correction factors				
Fetuses 1	IVF	no	Previous trisomy 21	no
Weight 83	diabetes	no pregnancies		
Smoker no	Origin	Asian		
Biochemical data	mical data		Ultrasound data	
Parameter Value	Corr. MoM	MoM Gestational age		12 + 3
PAPP-A 1.38 mIU/m	0.53	Method		CRL Robinson
fb-hCG 47.1 ng/ml	1.14	Scan date 17/02/2025		
Risks at sampling date	Crown rump length in mm		61	
Age risk	1:91	Nuchal translucency MoM		0.70
Biochemical T21 risk	1:95	Nasal bone Sonographer		present
Combined trisomy 21 risk	risk 1:565		r	DR. SANJEEV SHARMA
Trisomy 13/18 + NT	<1:10000	Qualifications in measuring NT		M.D
1:100 1:250 Cut off 1:100000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:100000 1:1000		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 565 women with the same data, there is one woman with a trisomy 21 pregnancy and 564 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!		
The calculated risk for trisomy 13 translucency) is < 1:10000, which risk.				

Sign of Physician