Date of report: 3/07/2025

JITM Diagnostics

Patient data				
lame MRS. JYOTI		Patient ID		
Birthday	4/08/1995		Sample ID 2507004620/NOD	
Age at sample date	29.9	Sample Date	e 3/07/2025	
Gestational age	12 + 2			
Correction factors				
Fetuses 1	IVF	no	Previous trisomy 21 no	
Weight 62.7	diabetes	no	pregnancies	
Smoker no	Origin	Asian		
Biochemical data		Ultrasound da	ata	
Parameter Value	Corr. MoM	Gestational age 12 + 1		
PAPP-A 2.5 mIU/n	nl 0.73	Method CRL Robinson		
fb-hCG 42.1 ng/ml	0.93	Scan date 2/07/2025		
Risks at sampling date		Crown rump length in mm 58		
Age risk	1:647	Nuchal translucency MoM 1.32		
Biochemical T21 risk	1:2351	Nasal bone present		
Combined trisomy 21 risk	1:3607	Sonographer DR. ANKIT KHANDELWAL MBBS DNE		
Trisomy 13/18 + NT	<1:10000	Qualifications in measuring NT MD		
1:100 1:250 1:1000 1:10000 13 15 17 19 21 23 25 27 29 31 33 3 Trisomy 13/18 + NT The calculated risk for trisomy 13 translucency) is < 1:10000, which risk.	Age 8/18 (with nuchal	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 3607 women with the same data, there is one woman with a trisomy 21 pregnancy and 3606 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!		