Date of report: 22/05/2025

JITM Diagnostics

Patient data				
Name	MRS. BHARTI			
Birthday	5/06/1995	Sample ID		2505042265/NOD
Age at sample date	30.0	Sample Date		20/05/2025
Gestational age	13 + 6			
Correction factors				
Fetuses 1	IVF	no	Previous trisomy 21	no
Weight 48	diabetes	no	pregnancies	
Smoker no	Origin	Asian		
Biochemical data		Ultrasound data		
Parameter Value	Corr. MoM	Gestational age		13 + 1
PAPP-A 10.1 mIU/m	l 1.16	Method		CRL Robinson
fb-hCG 88.1 ng/ml	2.11	Scan date		15/05/2025
Risks at sampling date		Crown rump length in mm		72.29
Age risk	1:679	Nuchal translucency MoM		0.71
Biochemical T21 risk	1:959	Nasal bone		present
Combined trisomy 21 risk	1:5098	Sonographer		DR. SHITIJ BUDHIRAJA
Trisomy 13/18 + NT	<1:10000	Qualifications in measuring NT		M.D
Risk 1:10		Trisomy 21	ated risk for Trisom	
1:1000 1:1000 1:10000 1:10000 1:10000 Age Trisomy 13/18 + NT The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.		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 5098 women with the same data, there is one woman with a trisomy 21 pregnancy and 5097 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!		