Date of report: 21/08/2024

JITM Diagnostics

Patient data				
Name	MRS. SWATI			
Birthday	20/07/1990			2408035404/NOD
Age at sample date	34.1	Sample Date		20/08/2024
Gestational age	13 + 5			
Correction factors				
Fetuses 1	IVF	no	Previous trisomy 21	no
Weight 80	diabetes	no	pregnancies	
Smoker no	Origin	Asian		
Biochemical data Ultrasound data				
Parameter Value	Value Corr. MoM Gestational age		13 + 5	
PAPP-A 10.1 mIU/m	1 2.25	Method		CRL Robinson
fb-hCG 24.1 ng/ml	0.67	Scan date		20/08/2024
Risks at sampling date		Crown rump length in mm		79.9
Age risk	1:344		slucency MoM	0.68
Biochemical T21 risk	<1:10000	Nasal bone		present
Combined trisomy 21 risk	/ 21 risk <1:10000		r	DR. NEERJA CHOPRA
Trisomy 13/18 + NT	<1:10000	Qualifications in measuring NT		M.D
1:1000 1:10000 1:10000 1:10000 1:10000 1:10000 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.		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 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!		