Date of report: 15/04/25

JITM Diagnostics

Patient data			
Name MRS	MRS. SABIHA PARVEEN		
Birthday	16/08/96	Sample ID 2504029881/NOD	
Age at sample date	28.7	Sample Date	e 15/04/25
Gestational age	11 + 3		
Correction factors			
Fetuses 1	IVF	no	Previous trisomy 21 no
Weight 63	diabetes	no	pregnancies
Smoker no	Origin	Asian	
Biochemical data		Ultrasound data	
Parameter Value	Corr. MoM	Gestational age 11 + 2	
PAPP-A 2.1 mIU/m	l 0.94	Method CRL Robinson	
fb-hCG 45.5 ng/ml	0.91	Scan date 14/04/25	
Risks at sampling date		Crown rump length in mm 46.8	
Age risk	1:718	Nuchal translucency MoM 1.33	
Biochemical T21 risk	1:4859	Nasal bone present	
Combined trisomy 21 risk	1:7021	Sonographer DR. SANDEEP DUA	
Trisomy 13/18 + NT	<1:10000	Qualifications in measuring NT M.D	
Risk 1:10		Trisomy 21	
1:1000 1:10000 1:10000 1:10000 13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49 Age Trisomy 13/18 + NT The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.		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 7021 women with the same data, there is one woman with a trisomy 21 pregnancy and 7020 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!	