Date of report: 23/05/2025

JITM Diagnostics

Patient data				
Name	MRS. HEMA NEGI(B)		Patient ID	
Birthday	8/06/1991	Sample ID 2505045310/NOD(B)		
Age at sample date	34.0	Sample Date 23/05/2025		
Gestational age	13 + 0			
Correction factors				
Fetuses 2	IVF	no	Previous trisomy 21 no	
Weight 88	diabetes	no	pregnancies	
Smoker no	Origin	Asian		
Biochemical data		Ultrasound data		
Parameter Value	Corr. MoM	1 Gestational age 12 + 5		
PAPP-A 10.1 mIU/m	l 1.78	Method CRL Robinson		
fb-hCG 245 ng/ml	2.97	Scan date 21/05/2025		
Risks at sampling date		Crown rump length in mm 65.2		
Age risk	1:344	Nuchal translucency MoM 0.83		
Biochemical T21 risk	1:431	Nasal bone present		
Combined trisomy 21 risk	1:2111	Sonographer		
Trisomy 13/18 + NT	<1:10000	Qualifications in measuring NT		
Risk 1:10		Trisomy 21	ated risk for Trisomy 21 (with nuchal	
1:100 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	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 2111 women with the same data, there is one woman with a trisomy 21 pregnancy and 2110 women with not affected pregnancies. The free beta HCG level is high. The risk for this twin pregnancy has been calculated for a singleton pregnancy with corrected MoMs. 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!		