

JITM Diagnostics

Patient data			
Name	MRS. HEMA NEGI(A)		Patient ID
Birthday	8/06/1991	Sample ID	2505045310/NOD(A)
Age at sample date	34.0	Sample Date	23/05/2025
Gestational age	12 + 6		
Correction factors			
Fetuses	2	IVF	no
Weight	88	diabetes	no
Smoker	no	Origin	Asian
Previous trisomy 21 pregnancies	no		
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	10.1 mIU/ml	1.89	12 + 4
fb-hCG	245 ng/ml	2.93	Method
			CRL Robinson
			Scan date
			21/05/2025
Risks at sampling date			Crown rump length in mm
Age risk	1:343		64.1
Biochemical T21 risk	1:492		Nuchal translucency MoM
Combined trisomy 21 risk	1:2387		0.82
Trisomy 13/18 + NT	<1:10000		Nasal bone
			present
			Sonographer
			..
			Qualifications in measuring NT
			..
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 2387 women with the same data, there is one woman with a trisomy 21 pregnancy and 2386 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!
Trisomy 13/18 + NT			
The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.			

Sign of Physician