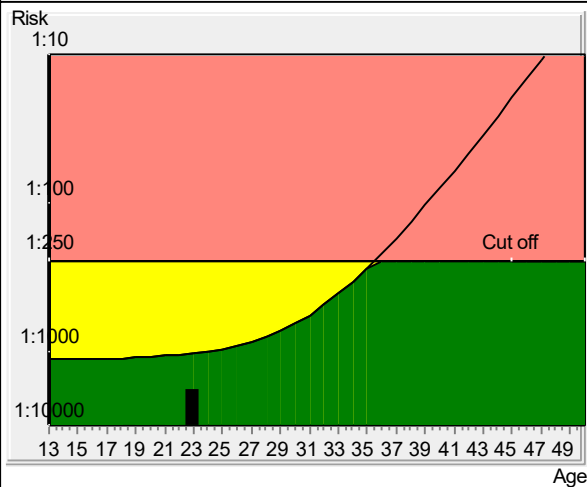


JITM Diagnostics

Patient data			
Name	MRS. MEENU KORI	Patient ID	
Birthday	26/02/2002	Sample ID	2501001449/NOD
Age at sample date	22.9	Sample Date	2/01/2025
Gestational age	12 + 6		
Correction factors			
Fetuses	1	IVF	no
Weight	55	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	1.45 mIU/ml	0.28	12 + 5
fb-hCG	32.1 ng/ml	0.72	Method
			CRL Robinson
			Scan date
			1/01/2025
Risks at sampling date			
Age risk		1:1046	Crown rump length in mm
Biochemical T21 risk		1:481	66.3
Combined trisomy 21 risk		1:3194	Nuchal translucency MoM
Trisomy 13/18 + NT		1:8976	0.59
			Nasal bone
			present
			Sonographer
			DR. NARENDRA KUMAR SINHA
			Qualifications in measuring NT
			M.D



**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 3194 women with the same data, there is one woman with a trisomy 21 pregnancy and 3193 women with not affected pregnancies.  
 The PAPP-A level is low.  
 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 1:8976, which represents a low risk.**

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**Sign of Physician**