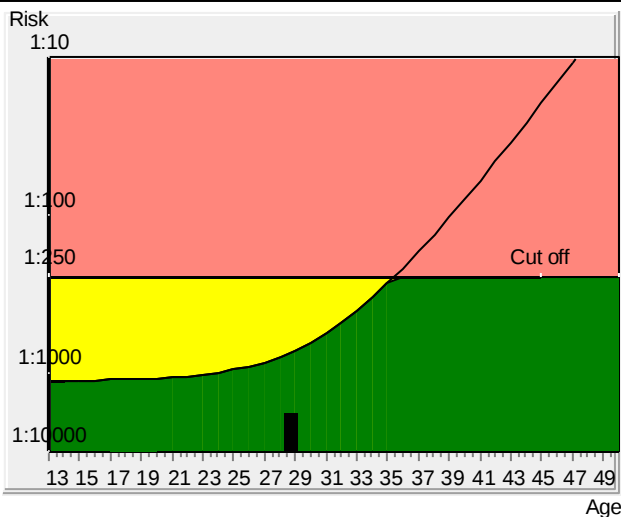


JITM Diagnostics

Patient data			
Name	MRS. SHEETAL	Patient ID	
Birthday	23/09/1996	Sample ID	2506012445/NOD
Age at sample date	28.7	Sample Date	8/06/2025
Gestational age	12 + 4		
Correction factors			
Fetuses	1	IVF	no
Weight	66	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	6.38 mIU/ml	1.75	
fb-hCG	40.6 ng/ml	0.94	
Risks at sampling date		Gestational age	12 + 3
Age risk	1:746	Method	CRL Robinson
Biochemical T21 risk	<1:10000	Scan date	7/06/2025
Combined trisomy 21 risk	<1:10000	Crown rump length in mm	62.7
Trisomy 13/18 + NT	<1:10000	Nuchal translucency MoM	1.18
		Nasal bone	present
		Sonographer	DR. ANKIT KHANDELWAL MBBS DNB
		Qualifications in measuring NT	MD
		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!

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