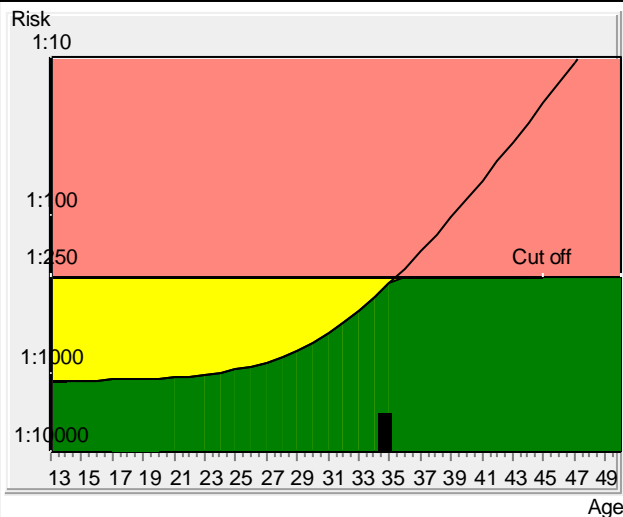


JITM Diagnostics

Patient data				
Name		MRS. SONAM	Patient ID	
Birthday		24/08/1990	Sample ID	
Age at sample date		34.7	2505044150/NOD	
Gestational age		12 + 4	Sample Date	
			22/05/2025	
Correction factors				
Fetuses	1	IVF	no	Previous trisomy 21
Weight	75	diabetes	no	pregnancies
Smoker	no	Origin	Asian	
Biochemical data			Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age	12 + 4
PAPP-A	3.66 mIU/ml	1.17	Method	CRL Robinson
fb-hCG	32.9 ng/ml	0.79	Scan date	22/05/2025
Risks at sampling date			Crown rump length in mm	64
Age risk		1:288	Nuchal translucency MoM	1.10
Biochemical T21 risk		1:4220	Nasal bone	present
Combined trisomy 21 risk		<1:10000	Sonographer	DR. ANKIT KHANDELWAL MBBS DNB
Trisomy 13/18 + NT		<1:10000	Qualifications in measuring NT	MD
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 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