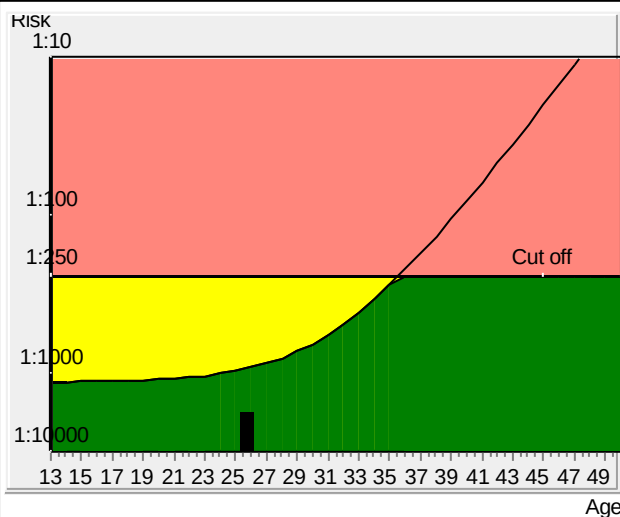


JITM Diagnostics

Patient data			
Name	MRS. SONAM KUMARI		Patient ID
Birthday	15/05/1999	Sample ID	2502024841/NOD
Age at sample date	25.8	Sample Date	13/02/2025
Gestational age	13 + 2		
Correction factors			
Fetuses	1	IVF	no
Weight	57	diabetes	no
Smoker	no	Origin	Asian
Previous trisomy 21 pregnancies	no		
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	5.8 mIU/ml	1.00	Gestational age 12 + 4
fb-hCG	30.8 ng/ml	0.74	Method CRL Robinson
Risks at sampling date			Scan date 8/02/2025
Age risk	1:951		Crown rump length in mm 64.1
Biochemical T21 risk	<1:10000		Nuchal translucency MoM 0.79
Combined trisomy 21 risk	<1:10000		Nasal bone present
Trisomy 13/18 + NT	<1:10000		Sonographer DR. S.K. SINGHAL
			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 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