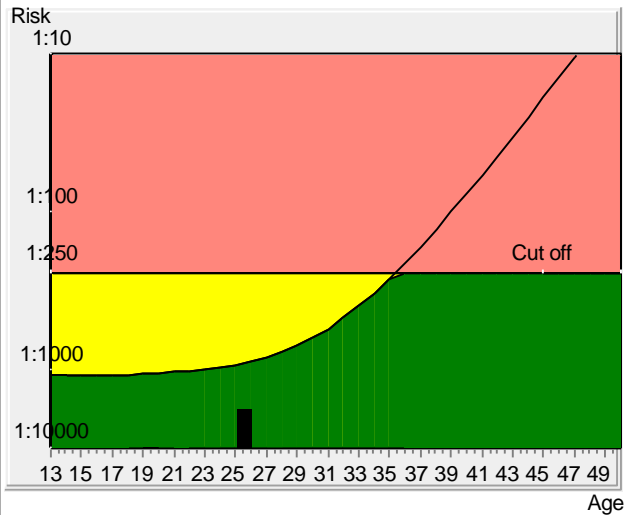


JITM DIAGNOSTICS

Prisca 5.2.0.13
Date of report: **15/01/2025**

J.I.T.M. DIAGNOSTICS

Patient data			
Name	MRS. MAMTA	Patient ID	
Birthdate	15/06/1999	Sample ID	2501021338/NOD
Age at sample date	25.6	Sample Date	14/01/2025
Gestational age	12 + 0		
Correction factors			
Fetuses	1	IVF	no
Weight	61	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	1 mIU/ml	0.32	Gestational age 11 + 4
fb-hCG	34 ng/ml	0.72	Method CRL Robinson
Risks at sampling date			Scan date 11/01/2025
Age risk		1:918	Crown rump length in mm 50
Biochemical T21 risk		1:634	Nuchal translucency MoM 0.67
Combined trisomy 21 risk		1:4053	Nasal bone present
Trisomy 13/18 + NT		<1:10000	Sonographer DR. RAHUL CHOUDHARY
			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 4053 women with the same data, there is one woman with a trisomy 21 pregnancy and 4052 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 < 1:10000, which represents a low risk.

Sign of Physician