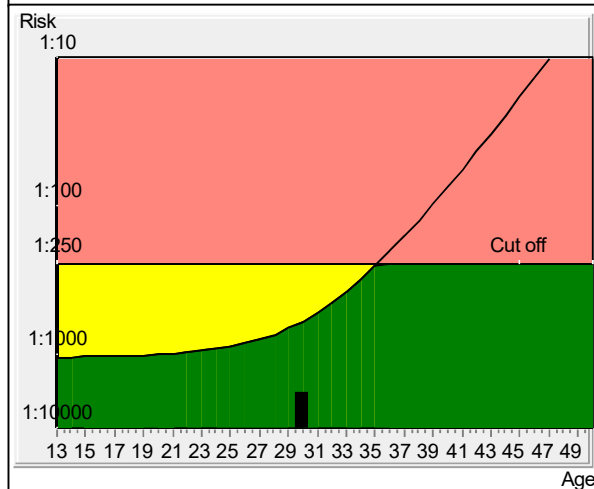


Patient data				
Name		MRS. GUNJAN MISHRA		Patient ID
Birthday		3/03/1995		Sample ID
Age at sample date		29.9		Sample Date
Gestational age		10 + 6		2501050550/NOD
				30/01/2025
Correction factors				
Fetuses	1	IVF	no	Previous trisomy 21
Weight	87	diabetes	no	pregnancies
Smoker	no	Origin	Asian	
Biochemical data			Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age	10 + 6
PAPP-A	3.6 mIU/ml	3.33	Method	CRL Robinson
fb-hCG	156 ng/ml	3.20	Scan date	30/01/2025
Risks at sampling date			Crown rump length in mm	42.3
Age risk		1:610	Nuchal translucency MoM	0.76
Biochemical T21 risk		1:749	Nasal bone	present
Combined trisomy 21 risk		1:3680	Sonographer	DR. GAURAV KUMAR
Trisomy 13/18 + NT		<1:10000	Qualifications in measuring NT	M.D
Risk			Trisomy 21	



Trisomy 13/18 + NT

The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low 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 3680 women with the same data, there is one woman with a trisomy 21 pregnancy and 3679 women with not affected pregnancies.

The free beta HCG level is high.

The PAPP-A level is high.

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!

Sign of Physician