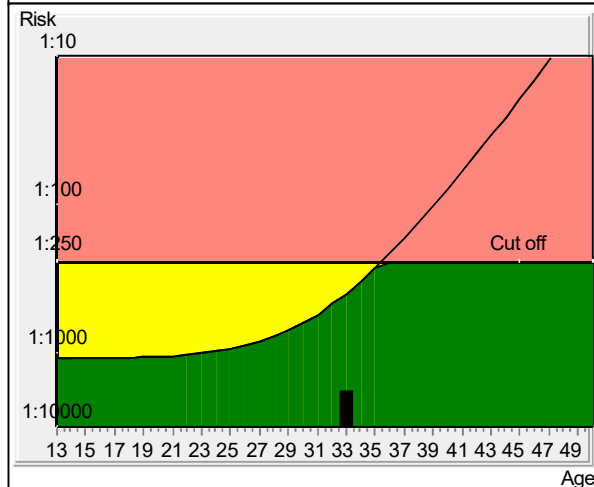


Patient data				
Name	MRS. NEHA KAUSHAL		Patient ID	2412036925/NOD 21/12/2024
Birthday	1/01/1992		Sample ID	
Age at sample date	33.0		Sample Date	
Gestational age	12 + 1			
Correction factors				
Fetuses	1	IVF	no	Previous trisomy 21
Weight	67	diabetes	no	pregnancies
Smoker	no	Origin	Asian	
Biochemical data			Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age	12 + 1
PAPP-A	1.62 mIU/ml	0.55	Method	CRL Robinson
fb-hCG	47.8 ng/ml	1.06	Scan date	21/12/2024
Risks at sampling date			Crown rump length in mm	58.62
Age risk		1:403	Nuchal translucency MoM	0.73
Biochemical T21 risk		1:533	Nasal bone	present
Combined trisomy 21 risk		1:3126	Sonographer	DR. PREETY SHARMA AGNIHOTRI
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 3126 women with the same data, there is one woman with a trisomy 21 pregnancy and 3125 women with not affected pregnancies.

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