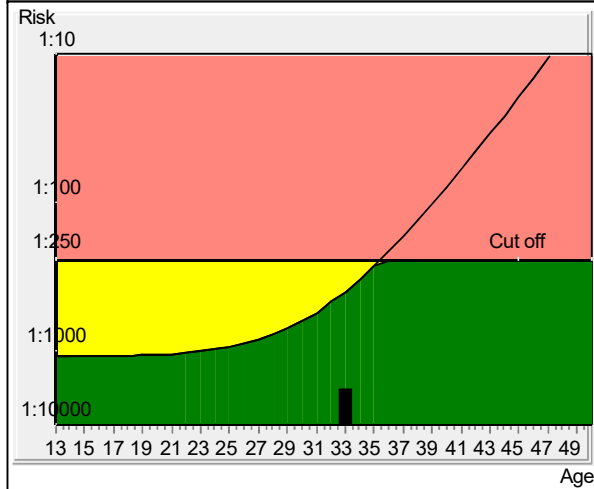


Patient data			
Name	MRS. NEHA KAUSHAL	Patient ID	
Birthday	1/01/1992	Sample ID	2412036925/NOD
Age at sample date	33.0	Sample Date	21/12/2024
Gestational age	12 + 1		
Correction factors			
Fetuses	1	IVF	no
Weight	67	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	1.62 mlU/ml	0.55	12 + 1
fb-hCG	47.8 ng/ml	1.06	Method
			CRL Robinson
			Scan date
			21/12/2024
Risks at sampling date			Crown rump length in mm
Age risk		1:403	58.62
Biochemical T21 risk		1:533	Nuchal translucency MoM
Combined trisomy 21 risk		1:3126	0.73
Trisomy 13/18 + NT		<1:10000	Nasal bone
			present
			Sonographer
			DR. PREETY SHARMA AGNIHOTRI
			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 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!

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