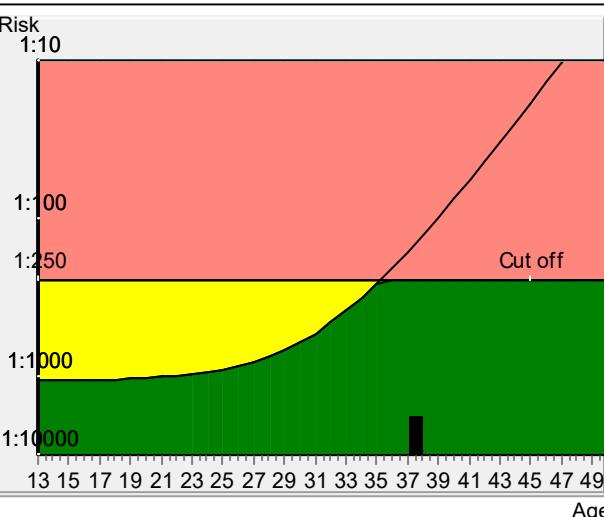


N A

Patient data			
Name	Mrs. MINARVA NAYAK	Patient ID	0652501240230
Birthday	25-06-1987	Sample ID	A1320699
Age at sample date	37.6	Sample Date	24-01-2025
Gestational age	11 + 2		
Correction factors			
Fetuses	1	IVF	no
Weight	70	diabetes	no
Smoker	no	Origin	Asian
Biochemical data			
Parameter	Value	Corr. MoM	Ultrasound data
PAPP-A	1.84 mIU/mL	0.98	Gestational age 11 + 1
fb-hCG	29.48 ng/mL	0.57	Method CRL Robinson
Risks at sampling date			
Age risk	1:140		Scan date 23-01-2025
Biochemical T21 risk	1:2781		Crown rump length in mm 45
Combined trisomy 21 risk	<1:10000		Nuchal translucency MoM 0.81
Trisomy 13/18 + NT	<1:10000		Nasal bone present
Sonographer N A			
Qualifications in measuring NT MD			
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 held 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