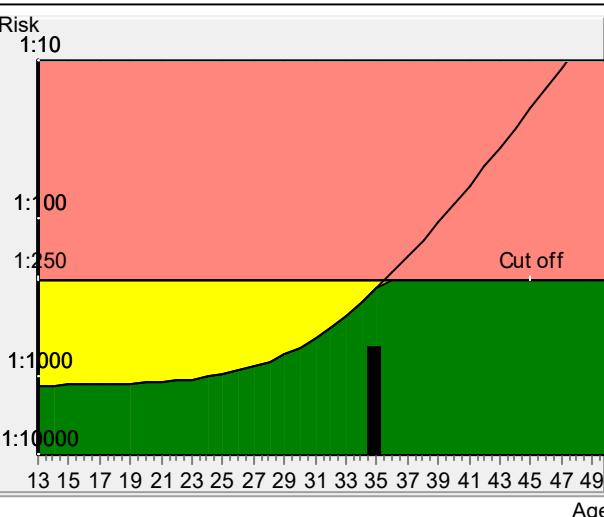


NA

Patient data								
Name	Mrs. POOJA MANI	Patient ID	0012510170356					
Birthday	10-12-1990	Sample ID	B3287472					
Age at sample date	34.9	Sample Date	17-10-2025					
Gestational age	13 + 2							
Correction factors								
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	unknown			
Weight	67.7	diabetes	no					
Smoker	no	Origin	Asian					
Biochemical data								
Parameter	Value	Corr. MoM	Ultrasound data					
PAPP-A	1.53 mIU/mL	0.33	Gestational age 13 + 1					
fb-hCG	34.9 ng/mL	1.05	Method CRL Robinson					
Risks at sampling date								
Age risk	1:289		Scan date 16-10-2025					
Biochemical T21 risk	1:97		Crown rump length in mm 72					
Combined trisomy 21 risk	1:649		Nuchal translucency MoM 0.72					
Trisomy 13/18 + NT	1:9571		Nasal bone present					
Sonographer NA								
Qualifications in measuring NT NA								
Trisomy 21								
The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk. <p>After the result of the Trisomy 21 test (with NT) it is expected that among 649 women with the same data, there is one woman with a trisomy 21 pregnancy and 648 women with not affected pregnancies.</p> <p>The PAPP-A level is low.</p> <p>The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician.</p> <p>Please note that risk calculations are statistical approaches and have no diagnostic value!</p> <p>The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).</p> <p>The laboratory can not be held responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!</p>								
								
Trisomy 13/18 + NT								
The calculated risk for Trisomy 13/18 (with nuchal translucency) is 1:9571, which represents a low risk.								

Sign of Physician