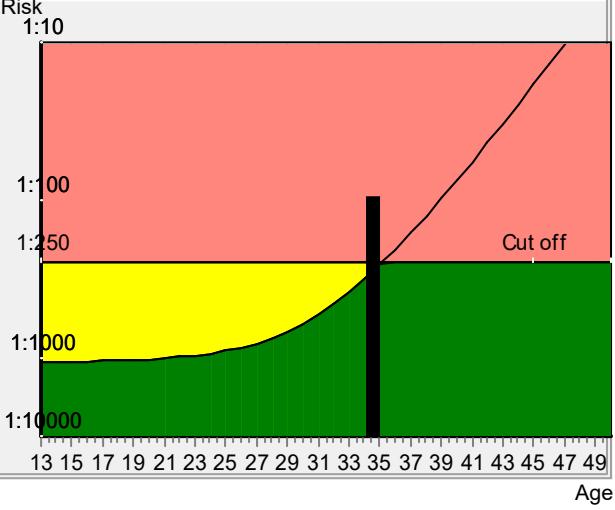


N A

Patient data								
Name	Mrs. RUMA KUMARI	Patient ID	0772510190055					
Birthday	02-04-1991	Sample ID	B3659198					
Age at sample date	34.5	Sample Date	19-10-2025					
Gestational age	11 + 0							
Correction factors								
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	unknown			
Weight	70	diabetes	no					
Smoker	no	Origin	Asian					
Biochemical data								
Parameter	Value	Corr. MoM	Ultrasound data					
PAPP-A	1.53 mIU/mL	0.94	Gestational age 10 + 6					
fb-hCG	52.01 ng/mL	0.95	Method CRL Robinson					
Risks at sampling date								
Age risk	1:281		Scan date 18-10-2025					
Biochemical T21 risk	1:1723		Crown rump length in mm 42.6					
Combined trisomy 21 risk	1:94		Nuchal translucency MoM 2.10					
Trisomy 13/18 + NT	1:1331		Nasal bone present					
Sonographer N A								
Qualifications in measuring NT MD								
Trisomy 21								
The calculated risk for Trisomy 21 (with nuchal translucency) is above the cut off, which indicates an increased risk.								
After the result of the Trisomy 21 test (with NT) it is expected that among 94 women with the same data, there is one woman with a trisomy 21 pregnancy and 93 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 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 1:1331, which represents a low risk.								

Sign of Physician