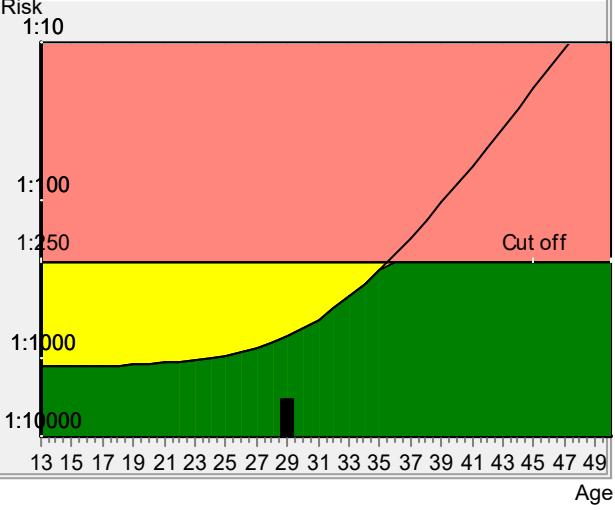


NA

Patient data								
Name	Mrs. SARITA NANNAWARE	Patient ID	0372510130140					
Birthday	05-10-1996	Sample ID	b3640439					
Age at sample date	29.0	Sample Date	13-10-2025					
Gestational age	12 + 6							
Correction factors								
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	unknown			
Weight	38	diabetes	no					
Smoker	no	Origin	Asian					
Biochemical data								
Parameter	Value	Corr. MoM	Ultrasound data					
PAPP-A	6.46 mIU/mL	0.85	Gestational age 12 + 5					
fb-hCG	38.84 ng/mL	0.85	Method CRL Robinson					
Risks at sampling date								
Age risk	1:730		Scan date 12-10-2025					
Biochemical T21 risk		1:4579	Crown rump length in mm 65					
Combined trisomy 21 risk		<1:10000	Nuchal translucency MoM 1.14					
Trisomy 13/18 + NT		<1:10000	Nasal bone present					
Sonographer NA								
Qualifications in measuring NT NA								
Trisomy 21								
<b>The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.</b> <p>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.</p> <p>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!</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								
<b>The calculated risk for trisomy 13/18 (with nuchal translucency) is &lt; 1:10000, which represents a low risk.</b>								

Sign of Physician

 below cut off

 Below Cut Off, but above Age Risk

 above cut off