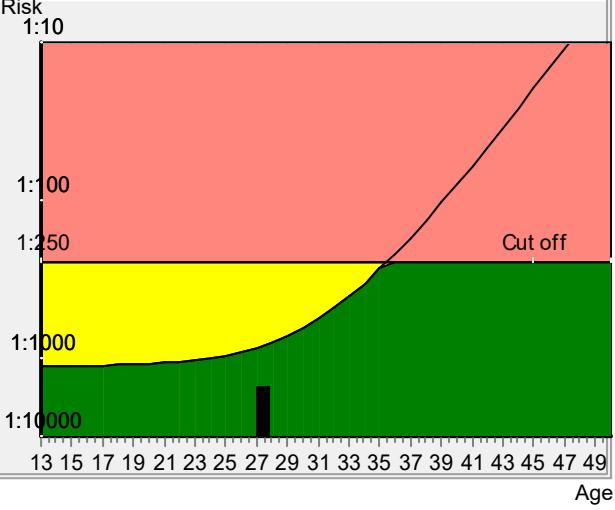


Patient data			
Name	Mrs. NANDANI	Patient ID	0442511220022
Birthday	22-06-1998	Sample ID	a1804986
Age at sample date	27.4	Sample Date	21-11-2025
Gestational age	12 + 5		
Correction factors			
Fetuses	1	IVF	no
Weight	66.5	diabetes	no
Smoker	no	Origin	Asian
Biochemical data			
Parameter	Value	Corr. MoM	Ultrasound data
PAPP-A	1.16 mIU/mL	0.31	Gestational age 12 + 5
fb-hCG	40.55 ng/mL	1.05	Method CRL Robinson
Risks at sampling date			
Age risk	1:839		Scan date 21-11-2025
Biochemical T21 risk	1:220		Crown rump length in mm 66.3
Combined trisomy 21 risk	1:1511		Nuchal translucency MoM 0.71
Trisomy 13/18 + NT	<1:10000		Nasal bone present
			Sonographer NA
			Qualifications in measuring NT Sonographer
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 1511 women with the same data, there is one woman with a trisomy 21 pregnancy and 1510 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 < 1:10000, which represents a low risk.			

Sign of Physician