

Prisca 5.1.0.17
Date of report: 02/05/25

N A

Patient data			
Name	Miss. PUJA NAYAK		Patient ID 0652504280220
Birth day	01/03/03		Sample ID A0958434
Age at sample date	22.2		Sample Date 28/04/25
Gestational age	11 + 5		
Correction factors			
Fetuses	1	IVF	no
Weight	60	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age 11 + 5
PAPP-A	1 mIU/mL	0.36	Method CRL Robinson
fb-hCG	48.08 ng/mL	0.96	Scan date 28/04/25
Risks at sampling date			Crown rump length in mm 51.6
Age risk	1:1021		Nuchal translucency MoM 0.72
Biochemical T21 risk	1:516		Nasal bone present
Combined trisomy 21 risk	1:3347		Sonographer N A
Trisomy 13/18 + NT	<1:10000		Qualifications in measuring NT MD
Risk			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 3347 women with the same data, there is one woman with a trisomy 21 pregnancy and 3346 women with not affected pregnancies. The PAPP-A level is low. 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 hold 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

below cut off
 Below Cut Off, but above Age Risk
 above cut off