

Prisca 5.1.0.17
Date of report: 29-01-2025

NA

Patient data			
Name	Mrs. MAYURI HAIGALE		Patient ID 0372501280125
Birthday	25-04-1994		Sample ID A1541252
Age at sample date	30.8		Sample Date 28-01-2025
Gestational age	13 + 5		
Correction factors			
Fetuses	1	IVF	no
Weight	52	diabetes	no
Smoker	no	Origin	Asian
			Previous trisomy 21 pregnancies unknown
Biochemical data			Ultrasound data
Parameter	Value	Corr. MoM	Gestational age 13 + 4
PAPP-A	4.93 mIU/mL	0.68	Method CRL Robinson
fb-hCG	62.2 ng/mL	1.95	Scan date 27-01-2025
Risks at sampling date			Crown rump length in mm 78
Age risk	1:609		Nuchal translucency MoM 0.79
Biochemical T21 risk	1:324		Nasal bone present
Combined trisomy 21 risk	1:1948		Sonographer NA
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 1948 women with the same data, there is one woman with a trisomy 21 pregnancy and 1947 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 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