

Prisca 5.1.0.17
Date of report: 19/01/25

NA

Patient data			
Name	Mrs. HARSHAVATI BORADE		Patient ID 0372501180105
Birth day	01/01/94		Sample ID A0319971
Age at sample date	31.0		Sample Date 18/01/25
Gestational age	13 + 6		
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 + 5
PAPP-A	5.27 mIU/mL	0.69	Method CRL Robinson
fb-hCG	51.05 ng/mL	1.67	Scan date 17/01/25
Risks at sampling date			Crown rump length in mm 79.54
Age risk 1:587			Nuchal translucency MoM 0.62
Biochemical T21 risk 1:478			Nasal bone present
Combined trisomy 21 risk 1:2809			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 2809 women with the same data, there is one woman with a trisomy 21 pregnancy and 2808 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