

Prisca 5.1.0.17
Date of report: 19/04/25

N A

Patient data			
Name	Mrs. JAYA DHAVAK		Patient ID 0372504180113
Birthday	11/03/92		Sample ID A1555833
Age at sample date	33.1		Sample Date 18/04/25
Gestational age	12 + 6		
Correction factors			
Fetuses	1	IVF	no
Weight	64	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age 12 + 4
PAPP-A	2.1 mIU/mL	0.50	Method CRL Robinson
fb-hCG	36.2 ng/mL	0.96	Scan date 16/04/25
Risks at sampling date			Crown rump length in mm 64
Age risk	1:404		Nuchal translucency MoM 1.22
Biochemical T21 risk	1:526		Nasal bone present
Combined trisomy 21 risk	1:1199		Sonographer N A
Trisomy 13/18 + NT	<1:10000		Qualifications in measuring NT MD
Risk			Trisomy 21
The graph shows the risk of Trisomy 21 as a function of maternal age. The y-axis represents risk from 1:10 to 1:10000. The x-axis represents age from 13 to 49. A red shaded area above the curve represents the risk above the cut-off. A yellow shaded area below the curve represents the risk below the cut-off but above the age risk. A green shaded area below the curve represents the risk below the cut-off. A vertical line at age 33.1 indicates the patient's age risk.			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 1199 women with the same data, there is one woman with a trisomy 21 pregnancy and 1198 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