

Prisca 5.1.0.17
Date of report: 09-08-2025

NA

Patient data			
Name	Mrs. SANIYA HAWALDAR	Patient ID	0662508080232
Birthday	18-06-2005	Sample ID	B3299115
Age at sample date	20.1	Sample Date	08-08-2025
Gestational age	13 + 1		
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	
PAPP-A	3.08 mIU/mL	0.52	Gestational age 12 + 4
fb-hCG	35.86 ng/mL	0.95	Method CRL Robinson
Risks at sampling date			Scan date 04-08-2025
Age risk		1:1109	Crown rump length in mm 64.5
Biochemical T21 risk		1:1592	Nuchal translucency MoM 0.91
Combined trisomy 21 risk		1:8392	Nasal bone present
Trisomy 13/18 + NT		<1:10000	Sonographer NA
			Qualifications in measuring NT NA
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 8392 women with the same data, there is one woman with a trisomy 21 pregnancy and 8391 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

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