

Prisca 5.1.0.17
Date of report: 24-12-2025

Patient data			
Name	Mrs. SHITAL KHANDARE	Patient ID	0662512230162
Birthday	28-02-2003	Sample ID	B4372158
Age at sample date	22.8	Sample Date	23-12-2025
Gestational age	12 + 5		
Correction factors			
Fetuses	1	IVF	no
Weight	46	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	5.38 mIU/mL	0.93	Gestational age 12 + 5
fb-hCG	41.07 ng/mL	0.93	Method CRL Robinson
Risks at sampling date			Scan date 23-12-2025
Age risk		1:1042	Crown rump length in mm 65.1
Biochemical T21 risk		1:6381	Nuchal translucency MoM 0.66
Combined trisomy 21 risk		<1:10000	Nasal bone present
Trisomy 13/18 + NT		<1:10000	Sonographer NA
			Qualifications in measuring NT Sonographer
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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy. 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