

Prisca 5.1.0.17
Date of report: 19-11-2025

N A

Patient data			
Name	Mrs. SUNITA PANDHARE		Patient ID
Birthday	08-06-1992		Sample ID
Age at sample date	33.4		Sample Date
Gestational age	13 + 0		
Correction factors			
Fetuses	1	IVF	no
Weight	71	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	2.86 mIU/mL	0.73	13 + 0
fb-hCG	36.97 ng/mL	1.05	Method
			CRL Robinson
			Scan date
			18-11-2025
Risks at sampling date			Crown rump length in mm
Age risk			70
Biochemical T21 risk			Nuchal translucency MoM
1:1063			0.57
Combined trisomy 21 risk			Nasal bone
1:5766			present
Trisomy 13/18 + NT			Sonographer
<1:10000			N A
			Qualifications in measuring NT
			MD
Risk		Trisomy 21	
		<p>The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.</p> <p>After the result of the Trisomy 21 test (with NT) it is expected that among 5766 women with the same data, there is one woman with a trisomy 21 pregnancy and 5765 women with not affected pregnancies.</p> <p>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!</p> <p>The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).</p> <p>The laboratory can not be hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!</p>	
Trisomy 13/18 + NT			
<p>The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.</p>			

Sign of Physician

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