

Prisca 5.1.0.17
Date of report: 12-01-2026

Patient data			
Name	Mrs. AARTI MHASKE		Patient ID
Birthday	15-06-1991		Sample ID
Age at sample date	34.6		Sample Date
Gestational age	13 + 0		
Correction factors			
Fetuses	1	IVF	no
Weight	79	diabetes	no
Smoker	no	Origin	Asian
			Previous trisomy 21 pregnancies
			unknown
Biochemical data			Ultrasound data
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	3.59 mIU/mL	1.05	12 + 6
fb-hCG	36.55 ng/mL	1.07	Method
			CRL Robinson
			Scan date
			09-01-2026
Risks at sampling date			
Age risk			Crown rump length in mm
1:303			66.9
Biochemical T21 risk			Nuchal translucency MoM
1:1815			1.18
Combined trisomy 21 risk			Nasal bone
1:4332			present
Trisomy 13/18 + NT			Sonographer
<1:10000			NA
			Qualifications in measuring NT
			Sonographer
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 4332 women with the same data, there is one woman with a trisomy 21 pregnancy and 4331 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 held 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