

Prisca 5.1.0.17
Date of report: 16-10-2025

N A

Patient data			
Name	Mrs. R.MADHAVI		Patient ID
Birthday	05-05-2001		Sample ID
Age at sample date	24.4		Sample Date
Gestational age	12 + 4		
Correction factors			
Fetuses	1	IVF	no
Weight	42	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	5.94 mIU/mL	0.98	12 + 4
fb-hCG	40.2 ng/mL	0.85	Method
			CRL Robinson
			Scan date
			15-10-2025
Risks at sampling date			Crown rump length in mm
Age risk			64
1:985			Nuchal translucency MoM
Biochemical T21 risk			0.73
1:8303			Nasal bone
Combined trisomy 21 risk			present
<1:10000			Sonographer
Trisomy 13/18 + NT			N A
<1:10000			Qualifications in measuring NT
			MD
Risk			Trisomy 21
1:10			The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.
1:100			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.
1:250			The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician.
1:1000			Please note that risk calculations are statistical approaches and have no diagnostic value!
1:10000			The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).
Age			The laboratory can not be hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!
13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49			
Cut off			
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