

Prisca 5.1.0.17
 Date of report: 09/10/25

N A

Patient data			
Name	Mrs. MADHU KUMARI		Patient ID
Birth day	15/02/99	Sample ID	B2884008
Age at sample date	26.6	Sample Date	08/10/25
Gestational age	12 + 3		
Correction factors			
Fetuses	1	IVF	no
Weight	54	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	4.58 mIU/mL	1.06	
fb-hCG	44.52 ng/mL	1.00	
Risks at sampling date			
Age risk	1:877		Gestational age
Biochemical T21 risk	1:6153		Method
Combined trisomy 21 risk	1:7915		Scan date
Trisomy 13/18 + NT	<1:10000		Crown rump length in mm
			Nuchal translucency MoM
			Nasal bone
			Sonographer
			Qualifications in measuring NT
			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 7915 women with the same data, there is one woman with a trisomy 21 pregnancy and 7914 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