

Prisca 5.1.0.17
Date of report: 10-05-2025

N A

Patient data			
Name	Mrs. KAMINI MANHAR		Patient ID
Birthday	15-11-1995		Sample ID
Age at sample date	29.5		Sample Date
Gestational age	13 + 1		
Correction factors			
Fetuses	1	IVF	no
Weight	55	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	3.2 mIU/mL	0.58	12 + 4
fb-hCG	36.52 ng/mL	0.99	Method
			CRL Robinson
			Scan date
			03-05-2025
Risks at sampling date			Crown rump length in mm
			64
Age risk	1:702		Nuchal translucency MoM
Biochemical T21 risk	1:1213		0.67
Combined trisomy 21 risk	1:6961		Nasal bone
Trisomy 13/18 + NT	<1:10000		present
			Sonographer
			N A
			Qualifications in measuring NT
			MD
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 6961 women with the same data, there is one woman with a trisomy 21 pregnancy and 6960 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