

Prisca 5.1.0.17
Date of report: 27-09-2024

N A

Patient data			
Name	Mrs. NAMEERA MEHVISH	Patient ID	0012409260484
Birthday	11-01-2001	Sample ID	A1108607
Age at sample date	23.7	Sample Date	26-09-2024
Gestational age	13 + 1		
Correction factors			
Fetuses	1	IVF	no
Weight	59	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data			Ultrasound data
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	4.3 mIU/mL	0.84	13 + 1
fb-hCG	57.1 ng/mL	1.59	Method
			CRL Robinson
			Scan date
			26-09-2024
			Crown rump length in mm
			71
			Nuchal translucency MoM
			0.73
			Nasal bone
			present
			Sonographer
			N A
			Qualifications in measuring NT
			MD
Risks at sampling date			Trisomy 21
Age risk		1:1030	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 8240 women with the same data, there is one woman with a trisomy 21 pregnancy and 8239 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!
Biochemical T21 risk		1:1483	
Combined trisomy 21 risk		1:8240	
Trisomy 13/18 + NT		<1:10000	
Risk Age			
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