

Prisca 5.1.0.17
Date of report: 28/05/25

N A

Patient data			
Name	Mrs. HARSHA PARDESHI	Patient ID	0372505270083
Birthday	06/06/91	Sample ID	A1556269
Age at sample date	34.0	Sample Date	27/05/25
Gestational age	12 + 4		
Correction factors			
Fetuses	1	IVF	no
Weight	74	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	2.3 mIU/mL	0.73	12 + 3
fb-hCG	39.76 ng/mL	1.02	Method
			CRL Robinson
			Scan date
			26/05/25
Risks at sampling date			
Age risk		1:338	Crown rump length in mm
Biochemical T21 risk		1:988	61.3
Combined trisomy 21 risk		1:5346	Nuchal translucency MoM
Trisomy 13/18 + NT		<1:10000	0.57
			Nasal bone
			present
			Sonographer
			N A
			Qualifications in measuring NT
			MD
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 5346 women with the same data, there is one woman with a trisomy 21 pregnancy and 5345 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