

Prisca 5.1.0.17
Date of report: 17/10/25

N A

Patient data			
Name	Mrs. AARTI		Patient ID 0382510100125
Birthday	07/05/96		Sample ID B3574717
Age at sample date	29.4		Sample Date 10/10/25
Gestational age	12 + 5		
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 12 + 0
PAPP-A	3.38 mIU/mL	0.53	Method CRL Robinson
fb-hCG	40.52 ng/mL	0.89	Scan date 05/10/25
Risks at sampling date			Crown rump length in mm 56.8
Age risk	1:696		Nuchal translucency MoM 0.94
Biochemical T21 risk	1:1202		Nasal bone present
Combined trisomy 21 risk	1:5979		Sonographer N A
Trisomy 13/18 + NT	<1:10000		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 5979 women with the same data, there is one woman with a trisomy 21 pregnancy and 5978 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