

Prisca 5.1.0.17
Date of report: 26/10/24

Patient data			
Name	Mrs. MEENAKSHI		Patient ID 0352410210057
Birthday	24/05/05		Sample ID A1287410
Age at sample date	19.4		Sample Date 21/10/24
Gestational age	12 + 4		
Correction factors			
Fetuses	1	IVF	no
Weight	35	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age 12 + 4
PAPP-A	4.15 mIU/mL	0.56	Method CRL Robinson
fb-hCG	45.85 ng/mL	0.90	Scan date 21/10/24
Risks at sampling date			Crown rump length in mm 64
Age risk	1:1097		Nuchal translucency MoM 0.97
Biochemical T21 risk	1:2168		Nasal bone present
Combined trisomy 21 risk	1:9902		Sonographer N A
Trisomy 13/18 + NT	<1:10000		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 9902 women with the same data, there is one woman with a trisomy 21 pregnancy and 9901 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