

Prisca 5.1.0.17
Date of report: 09-12-2024

N A

Patient data			
Name	Mrs. TABASSUM.	Patient ID	0382412060152.
Birthday	01-05-1984	Sample ID	A1419543.
Age at sample date	40.6	Sample Date	06-12-2024
Gestational age	13 + 4		
Correction factors			
Fetuses	1	IVF	no
Weight	72	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	8.34 mIU/mL	1.78	
fb-hCG	37.58 ng/mL	1.25	
Risks at sampling date		Gestational age	
Age risk		1:70	
Biochemical T21 risk		1:779	
Combined trisomy 21 risk		1:3384	
Trisomy 13/18 + NT		<1:10000	
		Method	
		CRL Robinson	
		Scan date	
		04-12-2024	
		Crown rump length in mm	
		74	
		Nuchal translucency MoM	
		0.87	
		Nasal bone	
		present	
		Sonographer	
		N A	
		Qualifications in measuring NT	
		MD	
Risk		Trisomy 21	
1:10		The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.	
1:100		After the result of the Trisomy 21 test (with NT) it is expected that among 3384 women with the same data, there is one woman with a trisomy 21 pregnancy and 3383 women with not affected pregnancies.	
1:250		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!	
1:1000		The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).	
1:10000		The laboratory can not be held responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!	
13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49			
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