

Prisca 5.1.0.17
Date of report: 25-04-2025

N A

Patient data			
Name	Mrs. SUSHMITHA	Patient ID	0842504240044
Birthday	05-06-1999	Sample ID	B2678141
Age at sample date	25.9	Sample Date	24-04-2025
Gestational age	11 + 6		
Correction factors			
Fetuses	1	IVF	no
Weight	45	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	2 mIU/mL	0.48	
fb-hCG	46.67 ng/mL	0.87	
Risks at sampling date		Gestational age	
Age risk		11 + 6	
Biochemical T21 risk		Method	
1:898		CRL Robinson	
Biochemical T21 risk		Scan date	
1:1305		24-04-2025	
Combined trisomy 21 risk		Crown rump length in mm	
1:7702		55	
Trisomy 13/18 + NT		Nuchal translucency MoM	
<1:10000		0.48	
		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 7702 women with the same data, there is one woman with a trisomy 21 pregnancy and 7701 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 hold 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