

Prisca 5.1.0.17
Date of report: 13-03-2025

N A

Patient data			
Name	Mrs. KOMAL MISAL	Patient ID	0662503120046
Birthday	17-03-2001	Sample ID	B2503259
Age at sample date	24.0	Sample Date	12-03-2025
Gestational age	13 + 3		
Correction factors			
Fetuses	1	IVF	no
Weight	65	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	2.18 mIU/mL	0.43	
fb-hCG	37.79 ng/mL	1.17	
Risks at sampling date			
Age risk		1:1030	
Biochemical T21 risk		1:565	
Combined trisomy 21 risk		1:3588	
Trisomy 13/18 + NT		<1:10000	
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 3588 women with the same data, there is one woman with a trisomy 21 pregnancy and 3587 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