

Prisca 5.1.0.17
Date of report: 17/10/25

NA

Patient data			
Name	Mrs. KOMAL	Patient ID	0772510170056
Birthday	01/01/04	Sample ID	B3658610
Age at sample date	21.8	Sample Date	16/10/25
Gestational age	10 + 6		
Correction factors			
Fetuses	1	IVF	no
Weight	83	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	0.69 mIU/mL	0.57	10 + 4
fb-hCG	54.66 ng/mL	1.02	Method
			CRL Robinson
			Scan date
			14/10/25
Risks at sampling date			Trisomy 21
Age risk	1:993		Crown rump length in mm
Biochemical T21 risk	1:1547		39
Combined trisomy 21 risk	1:2156		Nuchal translucency MoM
Trisomy 13/18 + NT	<1:10000		1.35
			Nasal bone
			unknown
			Sonographer
			NA
			Qualifications in measuring NT
			NA
			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 2156 women with the same data, there is one woman with a trisomy 21 pregnancy and 2155 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