

Prisca 5.1.0.17
Date of report: 29/04/25

N A

Patient data			
Name	Mrs. KRISHNAVENI	Patient ID	0012504280205
Birthday	02/04/95	Sample ID	A1743320
Age at sample date	30.1	Sample Date	28/04/25
Gestational age	13 + 1		
Correction factors			
Fetuses	1	IVF	no
Weight	53	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	3.67 mIU/mL	0.63	
fb-hCG	37.01 ng/mL	0.99	
Risks at sampling date			
Age risk		1:654	
Biochemical T21 risk		1:1432	
Combined trisomy 21 risk		1:8020	
Trisomy 13/18 + NT		<1:10000	
			Gestational age 12 + 3
			Method CRL Robinson
			Scan date 23/04/25
			Crown rump length in mm 61
			Nuchal translucency MoM 0.70
			Nasal bone present
			Sonographer N A
			Qualifications in measuring NT MD
Risk		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 8020 women with the same data, there is one woman with a trisomy 21 pregnancy and 8019 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