

Prisca 5.1.0.17
Date of report: 08/11/25

N A

Patient data					
Name	Mrs. BHARATHI		Patient ID		
Birthday	05/11/01	Sample ID			
Age at sample date	24.0	Sample Date			
Gestational age	13 + 1				
Correction factors					
Fetuses	1	IVF	no		
Weight	40.8	diabetes	no		
Smoker	no	Origin	Asian		
		Previous trisomy 21 pregnancies	unknown		
Biochemical data		Ultrasound data			
Parameter	Value	Corr. MoM			
PAPP-A	7 mIU/mL	0.90			
fb-hCG	36.85 ng/mL	0.89			
Risks at sampling date					
Age risk	1:1020	Gestational age	13 + 1		
Biochemical T21 risk	1:6429	Method	CRL Robinson		
Combined trisomy 21 risk	<1:10000	Scan date	05/11/25		
Trisomy 13/18 + NT	<1:10000	Crown rump length in mm	72		
		Nuchal translucency MoM	0.89		
		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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy. 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