

Prisca 5.1.0.17
Date of report: 19/04/25

N A

Patient data			
Name	Mrs. MINAKSHI RAJBONGSHI		Patient ID
Birth day	16/09/90		Sample ID
Age at sample date	34.6		Sample Date
Gestational age	11 + 6		
Correction factors			
Fetuses	1	IVF	no
Weight	66	diabetes	no
Smoker	no	Origin	Asian
			Previous trisomy 21 pregnancies
			unknown
Biochemical data			Ultrasound data
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	2.34 mIU/mL	0.88	11 + 5
fb-hCG	46.67 ng/mL	0.99	Method
			CRL Robinson
			Scan date
			17/04/25
			Crown rump length in mm
			53
			Nuchal translucency MoM
			0.71
			Nasal bone
			unknown
			Sonographer
			N A
			Qualifications in measuring NT
			MD
Risks at sampling date			Trisomy 21
Age risk	1:289		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 7150 women with the same data, there is one woman with a trisomy 21 pregnancy and 7149 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!
Biochemical T21 risk	1:1392		
Combined trisomy 21 risk	1:7150		
Trisomy 13/18 + NT	<1:10000		
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