

Prisca 5.1.0.17
Date of report: 28-02-2025

PRIYANKA

Patient data			
Name	Mrs. GAYATHRI		Patient ID
Birthday	27-02-2002	Sample ID	a1176105
Age at sample date	23.0	Sample Date	27-02-2025
Gestational age	13 + 0		
Correction factors			
Fetuses	1	IVF	no
Weight	58	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	7.72 mIU/mL	1.56	Method
fb-hCG	35.74 ng/mL	0.95	Scan date
Risks at sampling date			Crown rump length in mm 63 Nuchal translucency MoM 0.74 Nasal bone present Sonographer N A Qualifications in measuring NT MD
Age risk	1:1047		
Biochemical T21 risk	<1:10000		
Combined trisomy 21 risk	<1:10000		
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 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