

Prisca 5.1.0.17
Date of report: 02-02-2025

NA

Patient data			
Name	Mrs. J SANGEETHA	Patient ID	0012502010211
Birthday	15-02-1992	Sample ID	24858564
Age at sample date	33.0	Sample Date	01-02-2025
Gestational age	11 + 1		
Correction factors			
Fetuses	1	IVF	no
Weight	48	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	1.81 mIU/mL	0.66	11 + 1
fb-hCG	94.99 ng/mL	1.57	Method
			CRL Robinson
			Scan date
			01-02-2025
Risks at sampling date			Trisomy 21
Age risk		1:387	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 1940 women with the same data, there is one woman with a trisomy 21 pregnancy and 1939 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		1:330	
Combined trisomy 21 risk		1:1940	
Trisomy 13/18 + NT		<1:10000	
Age			
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