

Prisca 5.1.0.17
Date of report: 25/12/25

PRIYANKA

Patient data			
Name	Mrs. MAHADEVI	Patient ID	0352512240036
Birthday	01/01/05	Sample ID	b3800379
Age at sample date	21.0	Sample Date	24/12/25
Gestational age	12 + 5		
Correction factors			
Fetuses	1	IVF	no
Weight	39	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	7.76 mIU/mL	1.11	12 + 1
fb-hCG	37.4 ng/mL	0.80	Method
			CRL Robinson
			Scan date
			20/12/25
Risks at sampling date		Trisomy 21	
Age risk	1:1081	Crown rump length in mm	57
Biochemical T21 risk	<1:10000	Nuchal translucency MoM	1.00
Combined trisomy 21 risk	<1:10000	Nasal bone	present
Trisomy 13/18 + NT	<1:10000	Sonographer	NA
		Qualifications in measuring NT	Sonographer
Age		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