

Prisca

5.1.0.17

Date of report:

23-07-2023

Patient data			
Name	Mrs. S THARUNI	Patient ID	0012307230209
Birthday	17-12-2004	Sample ID	24660351
Age at sample date	18.6	Sample Date	23-07-2023
Gestational age	12 + 4		
Correction factors			
Fetuses	1	IVF	no
Weight	86	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	1.26 mIU/mL	0.49	
fb-hCG	39.76 ng/mL	1.07	
Risks at sampling date			
Age risk		1:1106	
Biochemical T21 risk		1:1037	
Combined trisomy 21 risk		1:5886	
Trisomy 13/18 + NT		<1:10000	
			Gestational age 12 + 3
			Method CRL Robinson
			Scan date 22-07-2023
			Crown rump length in mm 62
			Nuchal translucency MoM 0.87
			Nasal bone present
			Sonographer R.UMA
			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 5886 women with the same data, there is one woman with a trisomy 21 pregnancy and 5885 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!	
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