

Prisca

5.1.0.17

Date of report: 09-05-2023

Patient data			
Name	Mrs. M MAMATHA	Patient ID	0012305090113
Birthday	01-01-1990	Sample ID	24041997
Age at sample date	33.3	Sample Date	09-05-2023
Gestational age	12 + 2		
Correction factors			
Fetuses	1	IVF	unknown
Weight	58	diabetes	unknown
Smoker	unknown	Origin	Caucasian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	2.62 mIU/mL	0.82	12 + 1
fb-hCG	42.5 ng/mL	0.99	Method
			CRL Robinson
			Scan date
			08-05-2023
Risks at sampling date		Crown rump length in mm	
Age risk	1:378	58	
Biochemical T21 risk	1:1554	Nuchal translucency MoM	
Combined trisomy 21 risk	1:8127	0.72	
Trisomy 13/18 + NT	<1:10000	Nasal bone	
		present	
		Sonographer	
		NA	
		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 8127 women with the same data, there is one woman with a trisomy 21 pregnancy and 8126 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