

Prisca 5.1.0.17
Date of report: 09-12-2024

N A

Patient data			
Name	Mrs. S MEHARAJ BANU	Patient ID	0042412080005
Birthday	01-01-1998	Sample ID	23125664
Age at sample date	26.9	Sample Date	08-12-2024
Gestational age	13 + 2		
Correction factors			
Fetuses	1	IVF	no
Weight	44	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	4.3 mIU/mL	0.57	
fb-hCG	73.36 ng/mL	1.90	
Risks at sampling date			
Age risk	1:886	Gestational age	13 + 2
Biochemical T21 risk	1:320	Method	CRL Robinson
Combined trisomy 21 risk	>1:50	Scan date	08-12-2024
Trisomy 13/18 + NT	1:664	Crown rump length in mm	74
		Nuchal translucency MoM	2.24
		Nasal bone	unknown
		Sonographer	N A
		Qualifications in measuring NT	MD
Risk		Trisomy 21	
		The calculated risk for Trisomy 21 (with nuchal translucency) is above the cut off, which indicates an increased risk. After the result of the Trisomy 21 Test (with nuchal translucency), it is expected that among less than 50 pregnancies with the same data, there is one 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 1:664, which represents a low risk.			

Sign of Physician