

Prisca

5.1.0.17

Date of report:

08/09/25

NA

Patient data					
Name	Mrs. RITU MAHALLE	Patient ID	0372509070110		
Birthday	10/10/04	Sample ID	B3128643 TWIN B		
Age at sample date	20.9	Sample Date	06/09/25		
Gestational age	12 + 3				
Correction factors					
Fetuses	2	IVF	no		
Weight	44	diabetes	no		
Smoker	no	Origin	Asian		
Biochemical data		Ultrasound data			
Parameter	Value	Corr. MoM			
PAPP-A	6.8 mIU/mL	0.61	Gestational age		
fb-hCG	42 ng/ml	0.43	Method		
Risks at sampling date		CRL Robinson			
Age risk	1:1072	Scan date			
Biochemical T21 risk	<1:10000	Crown rump length in mm			
Combined trisomy 21 risk	<1:10000	Nuchal translucency MoM			
Trisomy 13/18 + NT	<1:10000	0.65			
Trisomy 21		Nasal bone			
Risk		present			
1:10		Sonographer			
1:100		NA			
1:250		Qualifications in measuring NT			
1:1000		MD			
1:10000		Trisomy 21			
13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49		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 risk for this twin pregnancy has been calculated for a singleton pregnancy with corrected MoMs.			
		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 held 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