

Prisca 5.2.0.13
Date of report: 12/02/22

Patient data					
Name	MITA PRAMANICK	Patient ID			
Birthday	15/08/03	Sample ID	10127598		
Age at sample date	18.5	Sample Date	09/02/22		
Gestational age	12 + 3				
Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight	45	diabetes	unknown		
Smoker	unknown	Origin	Asian		
Biochemical data					
Parameter	Value	Corr. MoM		Ultrasound data	
PAPP-A	11.85 mIU/mL	1.90		Gestational age	12 + 4
fb-hCG	27.44 ng/mL	0.67		Method	CRL Robinson
Risks at sampling date					
Age risk	1:1101			Scan date	10/02/22
Biochemical T21 risk	<1:10000			Crown rump length in mm	64
Combined trisomy 21 risk	<1:10000			Nuchal translucency MoM	0.97
Trisomy 13/18 + NT	<1:10000			Nasal bone	unknown
Risk					
1:10				Sonographer	DR.ARINDAM BHATTACHARJEE
1:100				Qualifications in measuring NT	M.D.
1:250					
1:1000					
1:10000					
13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49					
Age					
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 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 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