

Prisca 5.2.0.13
Date of report: 26/01/22

Patient data			
Name	ARIFUN NAHAR RAHAMAN	Patient ID	
Birthday	27/09/86	Sample ID	10111947
Age at sample date	35.3	Sample Date	25/01/22
Gestational age	13 + 6		
Correction factors			
Fetuses	1	IVF	unknown
Weight	65	diabetes	unknown
Smoker	unknown	Origin	Asian
		Previous trisomy 21	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	4.5 mIU/mL	0.67	
fb-hCG	7.01 ng/mL	0.30	
Risks at sampling date			Gestational age
Age risk	1:265		13 + 0
Biochemical T21 risk	1:6073		Method
Combined trisomy 21 risk	<1:10000		CRL Robinson
Trisomy 13/18 + NT	1:7009		Scan date
			19/01/22
			Crown rump length in mm
			69.5
			Nuchal translucency MoM
			1.03
			Nasal bone
			unknown
			Sonographer
			DR. SALAUDDIN MOLLA.
			Qualifications in measuring NT
			MBBS.
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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy. The free beta HCG level is low. 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 1:7009, which represents a low risk.			

Sign of Physician

below cut off
 Below Cut Off, but above Age Risk
 above cut off