

Prisca 5.2.0.13
Date of report: 01/12/21

Patient data			
Name	MOUMITA DAS	Patient ID	
Birthday	31/03/95	Sample ID	10063298
Age at sample date	26.7	Sample Date	01/12/21
Gestational age	13 + 1		
Correction factors			
Fetuses	1	IVF	unknown
Weight	52	diabetes	no
Smoker	no	Origin	Asian
Previous trisomy 21		unknown	
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	9.7 mIU/mL	1.41	
fb-hCG	83.18 ng/mL	2.63	
Risks at sampling date		Gestational age	
Age risk		12 + 6	
Biochemical T21 risk		Method	
1:1037		CRL Robinson	
Combined trisomy 21 risk		Scan date	
1:5313		29/11/21	
Trisomy 13/18 + NT		Crown rump length in mm	
<1:10000		67	
		Nuchal translucency MoM	
		0.82	
		Nasal bone	
		unknown	
		Sonographer	
		DR.B.K.BISWAS.	
		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 5313 women with the same data, there is one woman with a trisomy 21 pregnancy and 5312 women with not affected pregnancies. The free beta HCG level is high. 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