

Prisca 5.2.0.13
Date of report: 12/01/22

Patient data			
Name	ASMA KHATOON	Patient ID	
Birthday	22/11/90	Sample ID	10097796
Age at sample date	31.1	Sample Date	08/01/22
Gestational age	13 + 6		
Correction factors			
Fetuses	1	IVF	unknown
Weight	51	diabetes	unknown
Smoker	unknown	Origin	Asian
		Previous trisomy 21	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	7.3 mIU/mL	0.82	
fb-hCG	9.76 ng/mL	0.39	
Risks at sampling date			
Age risk	1:580		
Biochemical T21 risk	<1:10000		
Combined trisomy 21 risk	<1:10000		
Trisomy 13/18 + NT	<1:10000		
			Gestational age 13 + 4
			Method CRL Robinson
			Scan date 06/01/22
			Crown rump length in mm 78.61
			Nuchal translucency MoM 0.84
			Nasal bone unknown
			Sonographer DR.TULIKA DE.
			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 < 1:10000, which represents a low risk.			

Sign of Physician

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