

Prisca 5.2.0.13
Date of report: 16/12/21

Patient data			
Name	MOUPRIYA GHOSH	Patient ID	
Birthday	29/04/96	Sample ID	10077097
Age at sample date	25.6	Sample Date	16/12/21
Gestational age	13 + 3		
Correction factors			
Fetuses	1	IVF	unknown
Weight	87	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	6.52 mIU/mL	1.60	
fb-hCG	15.13 ng/mL	0.61	
Risks at sampling date			Gestational age
Age risk	1:961		13 + 1
Biochemical T21 risk	<1:10000		Method
Combined trisomy 21 risk	<1:10000		CRL Robinson
Trisomy 13/18 + NT	<1:10000		Scan date
			14/12/21
			Crown rump length in mm
			72
			Nuchal translucency MoM
			0.67
			Nasal bone
			unknown
			Sonographer
			DR.RAJU SINGHA.
			Qualifications in measuring NT
			MD.
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 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