

Prisca 5.2.0.13
Date of report: 09/01/22

Patient data			
Name	ARATI JENA	Patient ID	
Birthday	02/11/89	Sample ID	10095743
Age at sample date	32.2	Sample Date	07/01/22
Gestational age	12 + 5		
Correction factors			
Fetuses	1	IVF	unknown
Weight	52	diabetes	unknown
Smoker	unknown	Origin	Asian
		Previous trisomy 21	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	10.09 mIU/mL	1.71	
fb-hCG	11.02 ng/mL	0.31	
Risks at sampling date			Gestational age
Age risk	1:473		12 + 5
Biochemical T21 risk	<1:10000		Method
Combined trisomy 21 risk	<1:10000		CRL Robinson
Trisomy 13/18 + NT	<1:10000		Scan date
			07/01/22
			Crown rump length in mm
			66.7
			Nuchal translucency MoM
			0.88
			Nasal bone
			unknown
			Sonographer
			DR.G.SARKAR.
			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 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