

Prisca 5.2.0.13
Date of report: 12/02/22

Patient data			
Name	MITA PRAMANICK		Patient ID
Birthday	15/08/03		Sample ID
Age at sample date	18.5		Sample Date
Gestational age	12 + 3		
Correction factors			
Fetuses	1	IVF	unknown
Weight	45	diabetes	unknown
Smoker	unknown	Origin	Asian
		Previous trisomy 21	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	11.85 mIU/mL	1.90	Method
fb-hCG	27.44 ng/mL	0.67	Scan date
Risks at sampling date			Crown rump length in mm
Age risk			1:1101
Biochemical T21 risk			<1:10000
Combined trisomy 21 risk			<1:10000
Trisomy 13/18 + NT			<1:10000
			Nuchal translucency MoM
			0.97
			Nasal bone
			unknown
			Sonographer
			DR.ARINDAM BHATTACHARJEE.
			Qualifications in measuring NT
			M.D.
Trisomy 21			
<p>The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.</p> <p>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.</p> <p>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!</p> <p>The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).</p> <p>The laboratory can not be hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!</p>			
Trisomy 13/18 + NT			
<p>The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.</p>			

Sign of Physician

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