

Prisca 5.1.0.17
Date of report: 02-10-2023

Patient data			
Name	Mrs. SMITA MOHANTY		Patient ID
Birthday	23-06-1985		Sample ID
Age at sample date	38.3		Sample Date
Gestational age	12 + 3		
Correction factors			
Fetuses	1	IVF	no
Weight	65	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	5.86 mIU/mL	1.69	Method
fb-hCG	47.12 ng/mL	1.13	Scan date
Risks at sampling date			Crown rump length in mm
Age risk	1:124		Nuchal translucency MoM
Biochemical T21 risk	1:1621		Nasal bone
Combined trisomy 21 risk	1:7212		Sonographer
Trisomy 13/18 + NT	<1:10000		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 7212 women with the same data, there is one woman with a trisomy 21 pregnancy and 7211 women with not affected pregnancies. 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