

Prisca 5.1.0.17
Date of report: 19/09/25

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Patient data			
Name	Mrs. B.MANASA		Patient ID
Birth day	20/07/92		Sample ID
Age at sample date	33.2		Sample Date
Gestational age	12 + 5		
Correction factors			
Fetuses	1	IVF	no
Weight	64	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	3.2 mIU/mL	0.81	Method
fb-hCG	40.42 ng/mL	1.03	Scan date
Risks at sampling date			Crown rump length in mm
Age risk	1:398		Nuchal translucency MoM
Biochemical T21 risk	1:1437		Nasal bone
Combined trisomy 21 risk	1:7062		Sonographer
Trisomy 13/18 + NT	<1:10000		Qualifications in measuring NT
			MD
Risk			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 7062 women with the same data, there is one woman with a trisomy 21 pregnancy and 7061 women with not affected pregnancies.</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