

Patient data			
Name	MRS. VARSHA	Patient ID	
Birthday	11/01/1998	Sample ID	2412027827/NOD
Age at sample date	26.9	Sample Date	16/12/2024
Gestational age	13 + 0		
Correction factors			
Fetuses	1	IVF	no
Weight	60	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	2.12 mIU/ml	0.43	Gestational age 12 + 3
fb-hCG	21.9 ng/ml	0.52	Method CRL Robinson
Risks at sampling date			Scan date 12/12/2024
Age risk		1:878	Crown rump length in mm 61.7
Biochemical T21 risk		1:2780	Nuchal translucency MoM 0.69
Combined trisomy 21 risk		<1:10000	Nasal bone present
Trisomy 13/18 + NT		<1:10000	Sonographer ..
			Qualifications in measuring NT ..
Risk			Trisomy 21
			<p><b>The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.</b></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><b>The calculated risk for trisomy 13/18 (with nuchal translucency) is &lt; 1:10000, which represents a low risk.</b></p>			

Sign of Physician