

Patient data			
Name	MRS. KOMAL	Patient ID	
Birthday	4/11/1995	Sample ID	2409003427/NOD
Age at sample date	28.8	Sample Date	2/09/2024
Gestational age	13 + 3		
Correction factors			
Fetuses	1	IVF	no
Weight	63	diabetes	no
Smoker	no	Origin	Asian
Previous trisomy 21 pregnancies			no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	6.5 mIU/ml	1.20	13 + 3
fb-hCG	95.8 ng/ml	2.41	Method
			CRL Robinson
			Scan date
			2/09/2024
Risks at sampling date			Crown rump length in mm
Age risk		1:759	75.4
Biochemical T21 risk		1:813	Nuchal translucency MoM
Combined trisomy 21 risk		1:4364	0.70
Trisomy 13/18 + NT		<1:10000	Nasal bone
			present
			Sonographer
			..
			Qualifications in measuring NT
			..
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 4364 women with the same data, there is one woman with a trisomy 21 pregnancy and 4363 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><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**