

Patient data			
Name	MRS. POOJA		Patient ID
Birthday	01/01/96		Sample ID
Age at sample date	29.2		Sample Date
Gestational age	12 + 0		
Correction factors			
Fetuses	1	IVF	no
Weight	40	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	2.7 mIU/ml	0.54	Gestational age
fb-hCG	43.1 ng/ml	0.78	Method
Risks at sampling date			13/03/25
Age risk	1:694		Crown rump length in mm
Biochemical T21 risk	1:1708		51.8
Combined trisomy 21 risk	1:4832		Nuchal translucency MoM
Trisomy 13/18 + NT	<1:10000		1.15
			Nasal bone
			present
			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 4832 women with the same data, there is one woman with a trisomy 21 pregnancy and 4831 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