

Patient data			
Name	Mrs. KHYATI SRIVASTAVA		Patient ID
Birthday	16-11-1986		Sample ID
Age at sample date	36.6		Sample Date
Gestational age	12 + 6		
Correction factors			
Fetuses	1	IVF	no
Weight	73	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	2.48 mIU/mL	0.70	Method
fb-hCG	39.78 ng/mL	1.10	Scan date
Risks at sampling date			Crown rump length in mm
Age risk	1:189		Nuchal translucency MoM
Biochemical T21 risk	1:419		Nasal bone
Combined trisomy 21 risk	1:2322		Sonographer
Trisomy 13/18 + NT	<1:10000		Qualifications in measuring NT
			MD
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 2322 women with the same data, there is one woman with a trisomy 21 pregnancy and 2321 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