

Patient data			
Name	MRS. RAJNI PATWAL	Patient ID	
Birthday	19/12/93	Sample ID	2305062229/NOD
Age at sample date	29.4	Sample Date	31/05/23
Gestational age	12 + 6		
Correction factors			
Fetuses	1	IVF	no
Weight	46	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	1.75 mIU/ml	0.28	Gestational age 12 + 3
fb-hCG	70.1 ng/ml	1.48	Method CRL Robinson
Risks at sampling date			
Age risk		1:698	Scan date 28/05/23
Biochemical T21 risk		1:63	Crown rump length in mm 60.8
Combined trisomy 21 risk		1:360	Nuchal translucency MoM 0.95
Trisomy 13/18 + NT		1:9324	Nasal bone present
			Sonographer DR. PRABHAT KUMAR DAS
			Qualifications in measuring NT M.D
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 360 women with the same data, there is one woman with a trisomy 21 pregnancy and 359 women with not affected pregnancies.</p> <p>The PAPP-A level is low.</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 1:9324, which represents a low risk.</p>			

Sign of Physician

