

Patient data			
Name	MRS. RAMANDEEP KAUR	Patient ID	
Birthday	26-01-1992	Sample ID	2305220274/AMB
Age at sample date	31.3	Sample Date	09-05-2023
Gestational age	13 + 1		
Correction factors			
Fetuses	1	IVF	no
Weight	72	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	5.2 mIU/ml	1.26	Gestational age 12 + 5
fb-hCG	29.4 ng/ml	0.74	Method CRL Robinson
Risks at sampling date			Scan date 06-05-2023
Age risk		1:554	Crown rump length in mm 66.6
Biochemical T21 risk		<1:10000	Nuchal translucency MoM 1.48
Combined trisomy 21 risk		1:8685	Nasal bone present
Trisomy 13/18 + NT		<1:10000	Sonographer ..
			Qualifications in measuring NT MD
Risk			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 8685 women with the same data, there is one woman with a trisomy 21 pregnancy and 8684 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>The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.</p>			

Sign of Physician