

Patient data			
Name	MRS. GURPREET KAUR	Patient ID	
Birthday	18/02/1987	Sample ID	2412220542/AMB
Age at sample date	37.8	Sample Date	24/12/2024
Gestational age	13 + 1		
Correction factors			
Fetuses	1	IVF	no
Weight	68	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	8.2 mIU/ml	1.85	Gestational age
fb-hCG	20.5 ng/ml	0.51	Method
			CRL Robinson
			Scan date
			24/12/2024
Risks at sampling date			Crown rump length in mm
Age risk		1:142	71
Biochemical T21 risk		<1:10000	Nuchal translucency MoM
Combined trisomy 21 risk		<1:10000	0.62
Trisomy 13/18 + NT		<1:10000	Nasal bone
			present
			Sonographer
			..
			Qualifications in measuring NT
			..
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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy.</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