

Patient data			
Name	MRS. AMARJEET KAUR	Patient ID	
Birthday	30/08/88	Sample ID	10009299
Age at sample date	32.8	Sample Date	04/07/21
Gestational age	12 + 6		
Correction factors			
Fetuses	1	IVF	no
Weight	63	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data			Ultrasound data
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	2.5 mIU/ml	0.58	12 + 4
fb-hCG	23.1 ng/ml	0.54	Method
			CRL Robinson
Risks at sampling date			Scan date
Age risk		1:424	02/07/21
Biochemical T21 risk		1:2582	Crown rump length in mm
Combined trisomy 21 risk		1:5634	63
Trisomy 13/18 + NT		<1:10000	Nuchal translucency MoM
			1.23
			Nasal bone
			present
			Sonographer
			DR. JASWINDER SINGH
			Qualifications in measuring NT
			M.D
Risk			Trisomy 21
1:10			The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.
1:100			After the result of the Trisomy 21 test (with NT) it is expected that among 5634 women with the same data, there is one woman with a trisomy 21 pregnancy and 5633 women with not affected pregnancies.
1:250			The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician.
1:1000			Please note that risk calculations are statistical approaches and have no diagnostic value!
1:10000			The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).
			The laboratory can not be hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!
	Age		
	13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49		
Trisomy 13/18 + NT			
The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.			

Sign of Physician

