

Patient data			
Name	MRS. GURJEET KAUR	Patient ID	1702220193/AMB
Birthday	29-07-1991	Sample ID	1702220193/AMB
Age at sample date	25.5	Sample Date	13-02-2017
Gestational age	13 + 1		
Correction factors			
Fetuses	1 IVF	no	Previous trisomy 21
Weight	56 diabetes	no	pregancies
Smoker	no	Origin	Asian
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	4.25 mIU/ml	0.90	13 + 1
fb-hCG	58.3 ng/ml	1.80	Method
Risks at sampling date			CRL Robinson
Age risk	1:957		Scan date
Biochemical T21 risk	1:1190		13-02-2017
Combined trisomy 21 risk	1:6602		Crown rump length in mm
Trisomy 13/18 + NT	<1:10000		72.2
			Nuchal translucency MoM
			0.50
			Nasal bone
			present
			Sonographer
			DR. POONAM LOOMBA
			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 6602 women with the same data, there is one woman with a trisomy 21 pregnancy and 6601 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