

Patient data			
Name	MRS. GURJEET	Patient ID	17040220611/AMB
Birth day	05-08-1991	Sample ID	17040220611/AMB
Age at sample date	25.7	Sample Date	24-04-2017
Gestational age	13 + 1		
Correction factors			
Fetuses	1 IVF	no	Previous trisomy 21
Weight	48 diabetes	no	pregancies
Smoker	no	Origin	Asian
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	5.4 mIU/ml	0.96	12 + 5
fb-hCG	67.1 ng/ml	1.96	Method
Risks at sampling date			CRL Robinson
Age risk		1:948	Scan date
Biochemical T21 risk		1:1091	21-04-2017
Combined trisomy 21 risk		1:6024	Crown rump length in mm
Trisomy 13/18 + NT		<1:10000	65.2
			Nuchal translucency MoM
			0.68
			Nasal bone
			present
			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 6024 women with the same data, there is one woman with a trisomy 21 pregnancy and 6023 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