

Patient data				
Name	MRS. MANPREET KAUR		Patient ID	2102245476/DHJ
Birthday	15-01-1985		Sample ID	2102245476/DHJ
Age at sample date	36.1		Sample Date	12-02-2021
Gestational age	12 + 5			
Correction factors				
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies
Weight	65	diabetes	no	
Smoker	no	Origin	Asian	
Biochemical data			Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age	12 + 3
PAPP-A	3.19 mIU/ml	0.81	Method	CRL Robinson
fb-hCG	54.3 ng/ml	1.27	Scan date	10-02-2021
Risks at sampling date			Crown rump length in mm	62.1
Age risk	1:215		Nuchal translucency MoM	0.62
Biochemical T21 risk	1:483		Nasal bone	present
Combined trisomy 21 risk	1:2628		Sonographer	..
Trisomy 13/18 + NT	<1:10000		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 2628 women with the same data, there is one woman with a trisomy 21 pregnancy and 2627 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

below cut off
 Below Cut Off, but above Age Risk
 above cut off