

Patient data			
Name	MRS. SATGURU PREET KAUR	Patient ID	2105246466/DHJ
Birthday	07-07-1988	Sample ID	2105246466/DHJ
Age at sample date	32.9	Sample Date	25-05-2021
Gestational age	12 + 1		
Correction factors			
Fetuses	1	IVF	no
Weight	80.9	diabetes	no
Smoker	no	Origin	Asian
Biochemical data			
Parameter	Value	Corr. MoM	Ultrasound data
PAPP-A	3.63 mIU/ml	1.55	Gestational age 12 + 1
fb-hCG	18.7 ng/ml	0.44	Method CRL Robinson
Risks at sampling date			
Age risk	1:410		Scan date 25-05-2021
Biochemical T21 risk	<1:10000		Crown rump length in mm 58
Combined trisomy 21 risk	<1:10000		Nuchal translucency MoM 0.92
Trisomy 13/18 + NT	<1:10000		Nasal bone present
			Sonographer DR. RAJEEV MAHAJAN
			Qualifications in measuring NT M.D
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 held 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