

Date of report: 06-06-2023

Patient data			
Name	MRS. SANDEEP KAUR (B)		Patient ID
Birthday	22-08-1989		Sample ID
Age at sample date	33.8		Sample Date
Gestational age	12 + 0		05-06-2023
Correction factors			
Fetuses	2	IVF	yes
Weight	58	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	7.56 mIU/ml	1.24	11 + 5
fb-hCG	375 ng/ml	3.60	Method
			CRL Robinson
			Scan date
			03-06-2023
Risks at sampling date			Crown rump length in mm
			51.58
Age risk			Nuchal translucency MoM
1:343			0.58
Biochemical T21 risk			Nasal bone
1:133			present
Combined trisomy 21 risk			Sonographer
1:744			DR. SAMEER GARG
Trisomy 13/18 + NT			Qualifications in measuring NT
<1:10000			M.D
Risk			Trisomy 21
			<p><b>The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.</b></p> <p>After the result of the Trisomy 21 test (with NT) it is expected that among 744 women with the same data, there is one woman with a trisomy 21 pregnancy and 743 women with not affected pregnancies.</p> <p>The free beta HCG level is high.</p> <p>The risk for this twin pregnancy has been calculated for a singleton pregnancy with corrected MoMs.</p> <p>The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician.</p> <p>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><b>The calculated risk for trisomy 13/18 (with nuchal translucency) is &lt; 1:10000, which represents a low risk.</b></p>			

Sign of Physician

