

Patient data			
Name	MRS. AMITA (A)	Patient ID	2204220189/AMB (A)
Birthday	31/03/93	Sample ID	2204220189/AMB (A)
Age at sample date	29.0	Sample Date	08/04/22
Gestational age	13 + 4		
Correction factors			
Fetuses	2	IVF	no
Weight	60	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	13.2 mIU/ml	1.17	Gestational age 13 + 3
fb-hCG	375 ng/ml	4.35	Method CRL Robinson
Risks at sampling date		Scan date 07/04/22	
Age risk	1:748	Crown rump length in mm 75.4	
Biochemical T21 risk	1:193	Nuchal translucency MoM 0.77	
Combined trisomy 21 risk	1:1113	Nasal bone present	
Trisomy 13/18 + NT	<1:10000	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 1113 women with the same data, there is one woman with a trisomy 21 pregnancy and 1112 women with not affected pregnancies. The free beta HCG level is high. The risk for this twin pregnancy has been calculated for a singleton pregnancy with corrected MoMs. 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! The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)). The laboratory can not be hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!</p>	
		<p>The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.</p>	
Trisomy 13/18 + NT			

Sign of Physician