

*Free Home Sample Collection 9999 778 778



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 Date of Report
 2/8/2021

 PRISCA
 5.1.0.17

Patient Data				
Name MRS. KOMAL TANWAR		Patient ID		012107210160
Birthday	29/11/1995	Sample ID		11122552
Age at delivery 25.6		Sample Date		22/07/2021
Gestational age	13+1			
Correction factors				
Fetuses 1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg 51	Diabetes	NO	Pregnancies	unknown
Smoker NO	Origin	Asian		
Biochemical Data		Ultrasound Data		
Parameter Value	Corr Mom	Gestational age	e	12+6
PAPP-A 3.28	mIU/ml 0.49	Method		CRL(<>Robinson
fb-hCG 37.2	ng/ml 0.81	Scan date		21/07/2021
Risks at sampling date		Crown rump length in mm 64		
Age Risk	1:952	Nuchal translucency MOM		1.04
Biochemical T21 risk	1:1632	Nasal bone		Present
Combined Trisomy 21 Risk	1:1:6595	Sonographer		DR.
Trisomy 13/18 + NT <1:10000		Qualification in measuring NT		MD
Risk	Down's Syndrome Risk (Trisomy 21 Screening)			
1:100 1:250 1:1000 1:10000 1:1		The calculated risk for Trisomy 21 (with NT) is below the cut off, which represents a low risk. After the result of the Trisomy 21 test (with NT) it is expected that among 6595 women with the same data, there is one woman with a trisomy 21 pregnancy and 6594 women with not affected pregnancies. The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician. Please note that the risk calculations are statistical aapproaches and have no diagnostic value! The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values Risk above Age Risk Risk below Age risk		