

*Free Home Sample Collection 9999 778 778



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Date of Report 28/11/2021PRISCA 5.1.0.17

Patient Data					
Name	MRS. MEGHNA		Patient ID		012111270008
Birthday		21/08/1990	Sample ID		11056812
Age at delivery		31.3	Sample Date		27/11/2021
Gestational age		12+1			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	70 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	2	11+5
PAPP-A	$4.3~\mathrm{mIU/ml}$	1.33	Method		CRL(<>Robinson
fb-hCG	17.8 ng/ml	0.38	Scan date		24/11/2021
Risks at sampling date			Nuchal translucency 1.1		
Age Risk		1:535	Nuchal translucency MoM 0.2		0.79
Biochemical T21 risk		<1:10000	Nasal bone Present		
Combined T21 risk		<1:10000			
Trisomy 13/18+NT		<1:10000			
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
1:100 1:250 1:1000 1:1000 1:10000 1:3 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49 Age Trisomy 13/18 + NT The calculated risk for Trisomy 13/18 (with NT) is <1:10000, which indicates a low risk Risk Above Cut Off			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 it is expected that among more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy and 9999 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		