

\*Free Home Sample Collection 9999 778 778



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 Date of Report
 19/7/2021

 PRISCA
 5.1.0.17

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Patient Data					
Name		MRS. ANJALI	Patient ID		012107180212
Birthday		10/5/2001	Sample ID		11122602
Age at delivery		20.2	Sample Date		18/07/2021
Gestational age		13+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	13+0
PAPP-A	2.95 mIU/ml	0.63	Method		CRL(<>Robinson
fb-hCG	32.5 ng/ml	0.78	Scan date		18/07/2021
Risks at sampling date			Crown rump le	ength in mm	60.1
Age Risk		1:1108	Nuchal translu	cency MOM	0.96
Biochemical T21 risk		1:4018	Nasal bone		Present
Combined Trisomy 21 Ris	k	<1:10000	Sonographer		DR.
Trisomy 13/18 + NT		<1:10000	Qualification in	n measuring NT	MD
Risk			Down's Syndro	ome Risk (Trisomy 21 S	creening)
1:100 1:250 Cut off 1:1000 1:10000 1:10000 1:10000 1:10000			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 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!		

Trisomy 13/18 + NT

The calculated risk for Trisomy 13/18 (with NT) is <1:10000, which indicates a low risk

The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values

Risk Above Cut Off

Risk above Age Risk



Risk below Age risk