

*Free Home Sample Collection **9999 778 778** Download "MOLQ" App on

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				Date of Report PRISCA	17-04-2024 5.2.0.13
Patient Data					
Name	MRS. AN	NJU BALA F2	Patient ID		12404160126
Birthday		15-07-1990	Sample ID		11841116
Age at Sample date		33.8	Sample Date		16-04-1993
Gestational age		12+6			
Correction factors				Γ	
Fetuses	2 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	57.8 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	2	12+1
PAPP-A	6.1 mIU/ml	0.54	Method		CRL (<>Robinson)
fb-hCG	31.5 ng/ml	0.4	Scan date		12-04-2024
Risks at sampling date			Crown rump length in mm 55.5		
Age Risk		1:355	Nuchal translu	cency MoM	0.89
Biochemical T21 risk		1:3172	Nasal bone		Present
Combined trisomy 21 risk		<1:10000	Sonographer		DR.
Trisomy 13/18 + NT		<1:10000	Qualifications	in measuring NT	MBBS
Risk 1:10			Down's Syndr	ome Risk (Trisomy 21	Screening)
1:100 1:250 Cut off 1:1000 1:1000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 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 (with NT) it is expected that among 10000 women with the same data, there is one woman with a trisomy 21 pregnancy. 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 patient combined risk presumes that NT measurement was done according to accepted guidelines (Prenat Diagn 18:511-523; 1998). The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values		