

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

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				Date of Report PRISCA	29-07-2024 5.2.0.13
Patient Data					
Name	MRS. AKANKSH	IA DHAWAN	Patient ID		12407280185
Birthday		31-03-1989	Sample ID 11896367		
Age at Sample date	ge at Sample date 35.3				28-07-2024
Gestational age 12+2					
Correction factors				-	
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	95 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian	L	
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational ag	e	12+2
PAPP-A	4.5 mIU/ml	1.61	Method		CRL (<>Robinson)
fb-hCG	41.3 ng/ml	1.18	Scan date		28-07-2024
Risks at sampling date			Crown rump length in mm 58.		
Age Risk 1:251			Nuchal translucency MoM 0.85		
Biochemical T21 risk	isk1:2715		Nasal bone		present
Combined trisomy 21 risk <1:10000		Sonographer DR. RAHU			
Trisomy 13/18 + NT		<1:10000	Qualifications	in measuring NT	MBBS
Risk 1:10			Down's Syndrome Risk (Trisomy 21 Screening)		
1:100 1:250 Cut off 1:1000 1:10000 13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49 Trisomy 13/18+NT The calculated risk for Trisomy 13/18 (with NT) is <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. 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		
which indicates a low risk	····· 10/10 (with 111)		risk assessment	! Calculated risks have no e	diagnostic values



Risk Above Cut Off

