

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

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				Date of Report	06-12-2024
Patient Data				PRISCA	5.2.0.13
Name	MR	S. JAI SHREE	Patient ID		12412040086
Birthday		-	Sample ID		11996222
Age at Sample date			Sample Date		04-12-2024
Gestational age		13+4	<u>^</u>		
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	69 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian	3	
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age		13+4
PAPP-A	7.5 mIU/ml	1.12	Method		CRL (<>Robinson)
fb-hCG	19.8 ng/ml	0.76	Scan date		04-12-2024
Risks at sampling date	15.6 19/11	0.70	Crown rump l	ength in mm	70.4
Age Risk		1:545	Nuchal translu	-	1.13
Biochemical T21 risk		1:8028	Nasal bone		
Combined trisomy 21 risk		<1:10000	Sonographer		present
		<1:10000	Ŭ .	in manageming NT	DR.
Trisomy 13/18 + NT		<1:10000		in measuring NT	MBBS
Risk 1:10			Down's Syndrome Risk (Trisomy 21 Screening) 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,		
1:100	/	/	-	nong more than 10000 wo nan with a trisomy 21 preg	
1:250		Cutoff			
				isk by PRISCA depends o	•
1 000			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		
1:1000		done according to accepted guidelines (Prenat Diagn 18:511-523;			
13 15 17 19 21 23 25 27 2 Trisomy 13/18+NT	293133353/39	4143454749	1990).	cannot be hold responsible	e for their impact on the
The calculated risk for Trisomy $13/18$ (with NT) is $<1:10000$, which indicates a low risk			risk assessment! Calculated risks have no diagnostic values		
winen mulcaus a 10w 115K			1		



Risk Above Cut Off

