

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

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				Date of Report PRISCA	30/3/2023 5.2.0.13
Patient Data					
Name	MRS. HI	MANI GARG	Patient ID		012303290178
Birthday		23/04/1988	Sample ID		11639219
Age at term		35.3	Sample Date		29/3/2023
Gestational age		12+6			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	72 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom			12+6
PAPP-A	4.28 mIU/ml	0.87	Method		CRL (<>Robinson)
fb-hCG	21.7 ng/ml	0.67	Scan date		29/3/2023
Risks at sampling date			Nuchal translucency 1.76		
Age Risk 1:280		1:280	Nuchal translucency MoM 1.0		
Biochemical T21 risk	hemical T21 risk 1:29		Nasal bone Pres		Present
Combined trisomy 21 risk	Σ.	<1:10000			
Trisomy 13/18 + NT		<1:10000			
Risk			Down's Syndr	ome Risk (Trisomy 2	1 Screening)
Risk 1:10 1:250 1:250 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:100 1			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 with NT 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		
<1:10000 , which indicate	1 IN I) 18	values			
Risk A	Above Cut Off		Risk above Ag	e Risk	Risk below Age risk