

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

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1/1999	[ Patient ID Sample ID Sample Date		012303140100 11507608 14/3/2023	
1/1999 24.8	) Sample ID 3 Sample Date		11507608	
24.8	Sample Date			
	•		14/3/2023	
12+6	5			
	unknown	Previous trisomy 21	unknown	
	NO	Pregnancies	unknown	
	Asian			
	Ultrasound Da	ata		
ſom			12+5	
43	Method		CRL (<>Robinson)	
51	Scan date		13/03/2023	
	Nuchal translu	cency	1.6	
	Nuchal translu	cency MoM	0.98	
	Nasal bone Preser		Present	
00				
00				
	Down's Syndro	ome Risk (Trisomy 2	1 Screening)	
1:100 1:250 1:1000			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 values	
		47 49 Age The laboratory on the risk assovalues	<ul> <li>the information provided by the reference of the info</li></ul>	