

*Free Home			
Sample Collection			
9999 778 778			

Download "MOLQ" App on

Book a Test Online www.molq.in

				Date of Report PRISCA	5/2/2022 5.2.0.13
Patient Data					
Name	MRS. AN	NKITA PASSI	Patient ID		012202040087
Birthday		9/4/1983	Sample ID		11220004
Age at delivery		38.8	Sample Date		4/2/2022
Gestational age		12+0			
Correction factors				Γ	
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	77 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	e	12+0
PAPP-A	3.45  mIU/ml	1.27	Method		CRL (<>Robinson)
fb-hCG	52.4 ng/ml	1.14	Scan date		4/2/2022
Risks at sampling date			Nuchal translucency 0.9		
Age Risk		1:105	Nuchal translucency MoM		0.64
Biochemical T21 risk		1:796	Nasal Bone		Present
Combined T21 risk		1:3802			
Trisomy 13/18+NT		<1:10000			
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
Risk 1:10 1:00 1:250 1:1000 1:1000 13 15 17 19 21 23 25 2 Trisomy 13/18 (with N The calculated risk for <1:10000, which indicat	Г) Trisomy 13/18 with	Age	<ul> <li>The calculated risk for Trisomy 21 (with NT) is below the cut off, which represents a low risk.</li> <li>After the result of the Trisomy 21 test it is expected that among 3802 women with the same data, there is one woman with a trisomy 21 pregnancy and 3801 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!</li> <li>The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values</li> </ul>		
Risk	x Above Cut Off		Risk above Ag	e Risk	Risk below Age risk