



Book a Test Online www.molq.in

25/8/2022 Date of Report PRISCA 5.1.0.17

Patient Data					
Name MRS. KM NEELAM			Patient ID 01220823025		
Birthday	12/11/1997		Sample ID		11245943
Age at term 25.02		2 Sample Date 23/8/202			
Gestational age		12+6			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	49 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age		12+5
PAPP-A	4.72 mIU/ml	0.61	Method		CRL (<>Robinson)
fb-hCG	65.2 ng/ml	1.78	Scan date		23/8/2022
Risks at sampling date			Nuchal translucency 1.4		
Age Risk 1:98		1:982	Nuchal translucency MoM 0		0.87
Biochemical T21 risk		1:492	Nasal bone		Presen
Combined trisomy 21 risk 1:2817					
Trisomy 13/18 + NT <1:10000					
Risk			Down's Syndro	ome Risk (Trisomy 21	Screening)
1:100 1:250 1:1000 1:1000 1:100000 1:10000 1:100000 1:100000 1:100000 1:100000 1:100000 1:100000 1:100000000			The calculated risk for Trisomy 21(with NT) is below the cut off, which represents a low risk. After the result of the Trisomy 21with NT test it is expected that among 2817 women with the same data, there is one woman with a trisomy 21 pregnancy and 2816 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		