



Book a Test Online www.molq.in

Date of Report 1/4/2022PRISCA 5.1.0.17

Patient Data					
Name MRS. SAIMA			Patient ID		042203300037
Birthday		15-04-1993	Sample ID		522007
Age at term		29.06	Sample Date		30/3/2022
Gestational age		12+2			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	59 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	2	11+6
PAPP-A	$2.82~\mathrm{mIU/ml}$	0.67	Method		CRL (<>Robinson)
fb-hCG	25.8 ng/ml	0.54	Scan date		28/03/2022
Risks at sampling date			Nuchal Translucency 1.3		
Age Risk		1:720	Nuchal Translucency MoM 0.		0.94
Biochemical T21 risk		1:6720	Nasal Bone Prese		Present
Combined T21 risk		<1:10000			
Trisomy 13/18+NT		<1:10000			
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
Risk 1:10			The calculated risk for Trisomy 21 is below the cut off, which represents a low risk.		
1:100 1:250 Cut off 1:1000 1:10000 13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49 Age			After the result of the Trisomy 21 test it is expected that among more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy 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!		
Trisomy 13/18 The calculated risk for indicates a low risk	or Trisomy 13/18 is <1	:10000 , which	The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values		
R	Risk Above Cut Off		Risk above Ago	e Risk	isk below A ge risk