



Book a Test Online www.molq.in

Date of Report	28/9/2022
PRISCA	5.1.0.17

Patient Data					
Name	Name MRS. SANJU(F1)		Patient ID		012209260209
Birthday		30/07/1992	Sample ID		11559523
Age at term	30.08		Sample Date		26/9/2022
Gestational age		13+1			
Correction factors					
Fetuses	2 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	55 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	2	13+0
PAPP-A	$6.98~\mathrm{mIU/ml}$	0.5	Method		CRL (<>Robinson)
fb-hCG	79.2 ng/ml	1.13	Scan date		26/9/2022
Risks at sampling date			Nuchal translu	cency	1.2
Age Risk		1:647	Nuchal translu	cency MoM	0.71
Biochemical T21 risk		1:565	Nasal bone		Present
Combined trisomy 21 ris	k	1:3437			
Trisomy 13/18 + NT		<1:10000			
Risk			Down's Syndro	ome Risk (Trisomy 2	1 Screening)
1:1000 1:1000 1:1000 1:10000 1:10000 1:10000 1:10000 1:10000 13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49 Age  Trisomy 13/18+NT The calculated risk for Trisomy 13/18 (with NT) is <1:10000, which indicates a low risk			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 3437 women with the same data, there is one woman with a trisomy 21 pregnancy and 3436 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		