

\*Free Home Sample Collection 9999 778 778



Book a Test Online www.molq.in

Date of Report	10-04-19
PRISCA	5.0.2.37

				PRISCA	5.0.2.37
Jame Mrs Sucharita Swain		Patient ID		011904080144	
	18-06-97		Sample ID		10419366
		21.8	Sample Date		08-04-2019
		12+3			
1	IVF		unknown	Previous trisomy 21	unknown
78.2	Diabetes		no	Pregnancies	
no	Origin		Asian		
			Ultrasound D	ata	
Value		Corr Mom	Gestational ag	e	12+3
3.65	mIU/ml	1.25	Method		LMP
29.7	ng/ml	0.70	LMP Date		12-01-19
			CRL measurm	ents	
			Nuchal translu	cency MoM	
		1:1055	Nasal bone		
		1:600	Sonographer		
		<1:10000	Qualifications in measuring NT		
		<1:10000			
			Down's Syndi	rome Risk (Trisomy 21 Sc	reening)
1:100 1:250 Out off		The calculated risk for Trisomy 21 (is below the cut off, which indicates a low risk.  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.  The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician. Please note that risk calculations are statistical approaches and have no diagnostic value!			
	1 78.2 no Value 3.65 29.7	1 IVF 78.2 Diabetes no Origin  Value  3.65 mIU/ml 29.7 ng/ml	18-06-97 21.8 12+3  1 IVF 78.2 Diabetes no Origin  Value Corr Mom 3.65 mIU/ml 1.25 29.7 ng/ml 0.70  1:1055 1:600 <1:10000 <1:10000	78.2 Diabetes no Origin Asian  Value Corr Mom Gestational age 3.65 mIU/ml 1.25 Method 29.7 ng/ml 0.70 LMP Date CRL measurm Nuchal translutions 1:1055 Nasal bone Sonographer <1:10000 Qualifications <1:10000  Down's Syndia Asian  Cut off The calculated which indicat among more that there is one would not be a simple of the information of the informat	Mrs Sucharita Swain 18-06-97 21.8 Sample ID Sample Date 12+3  IVF unknown Previous trisomy 21 Pregnancies  Origin Asian  Ultrasound Data  Value Corr Mom 3.65 mIU/ml 1.25 29.7 ng/ml 0.70 LMP Date CRL measurments Nuchal translucency MoM 1:1055 1:600 Sonographer <1:10000 <1:10000  Down's Syndrome Risk (Trisomy 21 See which indicates a low risk.  After the result of the Trisomy 21 (is by which indicates a low risk.  After the result of the Trisomy 21 pregnancies of the information provided by the referring Please note that risk calculations are statis approaches and have no diagnostic value!