MOLQ Express Lab
*Free Home Sample Collection 9999 778 778
Download "MOLQ" App on
Book a Test Online www.molq.in

Date of Report	27-02-19
PRISCA	5.0.2.37

Patient Data			
Name	**Mrs Pooja**	Patient ID	011902260102
Birthday	15-10-88	Sample ID	10418911
Age at delivery	30.4	Sample Date	26/02/19
Gestational age by	10+5		

Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	62	Diabetes	no	Pregnancies	
Smoker	no	Origin	Asian		

Biochemical Data

Parameter	Value		Corr Mom
PAPP-A	1.06	mIU/ml	0.53
fb-hCG	35.99	ng/ml	0.63

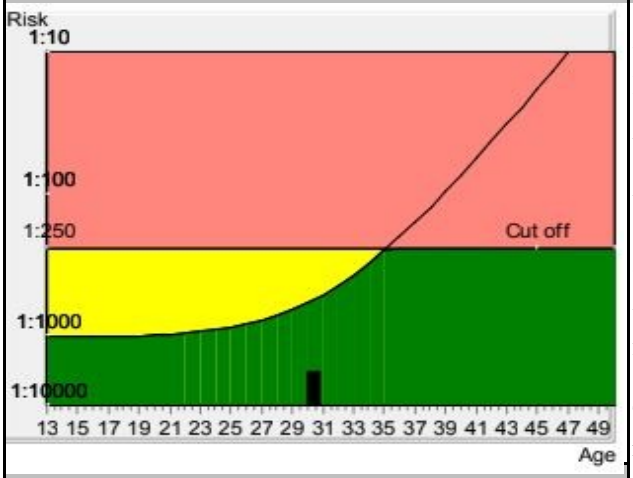
Ultrasound Data

Gestational age	10+5
Method	CRL (<>Hadlock)
Scan Date	26-02-19
CRL measurments	39.9
Nuchal translucency MoM	0.97
Nasal bone	Present
Sonographer	DR.VIKAS CHAUDHARY
Qualifications in measuring NT	MBBS,DMRD

Risks at sampling date

Age Risk	1:572
Biochemical T21 Risk	1:2097
Combined Trisomy 21 Risk	1:9567
Trisomy 13/18	<1:10000

Risk



Trisomy 13/18 + NT

The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.

Down's Syndrome Risk (Trisomy 21 Screening)

The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.

After the result of the Trisomy 21 test (with NT) it is expected that among 9567 women with the same data, there is one woman with a trisomy 21 pregnancy and 9566 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 risk calculations are statistical approach and have no diagnostic values!

The patient combined risk presumes the NT measurement was done according to accepted guidelines

Risk Above Cut Off | Risk above Age Risk | Risk below Age risk