MOLQ Express Lab

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Date of Report	18-02-19
PRISCA	5.0.2.37

Patient Data

Name	**Mrs Rosy**	Patient ID	011902160262
Birthday	19-07-85	Sample ID	10433685
Age at delivery	33.6	Sample Date	16/02/19
Gestational age by	13+0		

Correction factors

Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	68.2	Diabetes	no	Pregnancies	
Smoker	no	Origin	Asian		

Biochemical Data

Parameter	Value		Corr Mom
PAPP-A	5.79	mIU/ml	0.77
fb-hCG	282.3	ng/ml	6.92

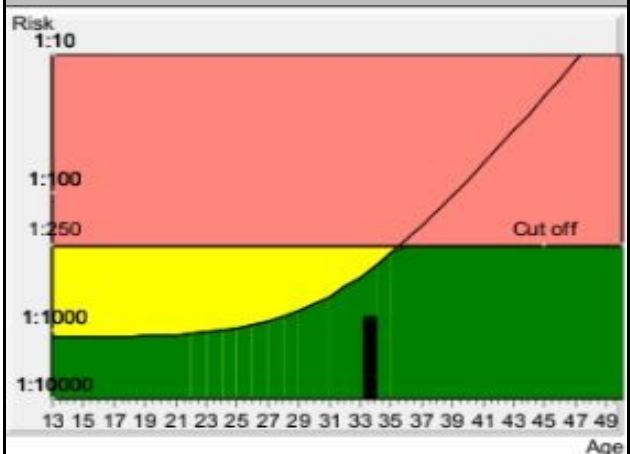
Ultrasound Data

Gestational age	12+6
Method	CRL (<>Hadlock)
Scan Date	16-02-19
CRL measurments	66.2
Nuchal translucency MoM	0.53
Nasal bone	Present
Sonographer	Dr. Sahil Loomba
Qualifications in measuring NT	MBBS.DNB

Risks at sampling date

Age Risk	1:371
Biochemical T21 Risk	1:148
Combined Trisomy 21 Risk	1:806
Trisomy 13/18	<1:10000

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 806 women with the same data, there is one woman with a trisomy 21 pregnancy and 805 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

Trisomy 13/18 + NT

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

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