MOLQ Express Lab
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Date of Report	17/12/19
PRISCA	5.0.2.37

Patient Data

Name	**Mrs Lalita Fetus A**	Patient ID	011912140199
Birthday	4/3/1991	Sample ID	10303062
Age at delivery	28.8	Sample Date	14/12/19
Gestational age	11+6		

Correction factors

Fetuses	2	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	75	Diabetes	no	Pregnancies	
Smoker	no	Origin	Asian		

Biochemical Data

Parameter	Value	Corr Mom
PAPP-A	3.54 mIU/ml	0.72
fb-hCG	112.5 ng/ml	1.10

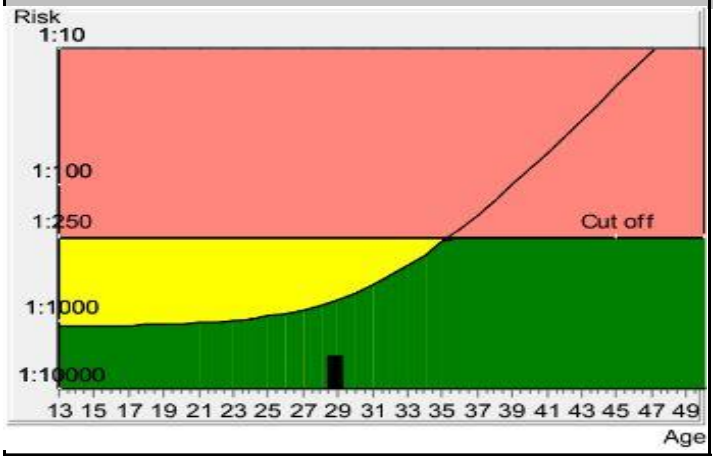
Ultrasound Data

Gestational age	11+5
Method	CRL (<>Robinson)
Scan Date	14/12/19
Crown Rump Length (mm)	49.1
	0.90
Nasal Bone	present
Sonographer	DR. VIKASH GOYAL
Qualification in measuring NT	DMRD

Risks at sampling date

Age Risk	1:721
Overall Population Risk	1:1699
Trisomy 21 Risk	1:8523
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 8523 women with the same data, there is one woman with a trisomy 21 pregnancy and 8522 women with not affected pregnancies.
The risk for this twin pregnancy has been calculated for a singleton pregnancy with corrected MoMs.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

Trisomy 13/18 + NT

The calculated risk for trisomy 13/18 is <1:10000, which represents a low risk.

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