MOLQ Express Lab
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Date of Report	01-02-19
PRISCA	5.0.2.37

Patient Data			
Name	**Mrs Varsha**	Patient ID	011901270140
Birthday	06-08-98	Sample ID	10428977
Age at delivery	20.5	Sample Date	27/01/2019
Gestational age	13+6		

Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	52.6	Diabetes	no	Pregnancies	
Smoker	no	Origin	Asian		

Biochemical Data

Parameter	Value	Corr Mom
PAPP-A	5.12 mIU/ml	0.62
fb-hCG	41.36 ng/ml	0.99

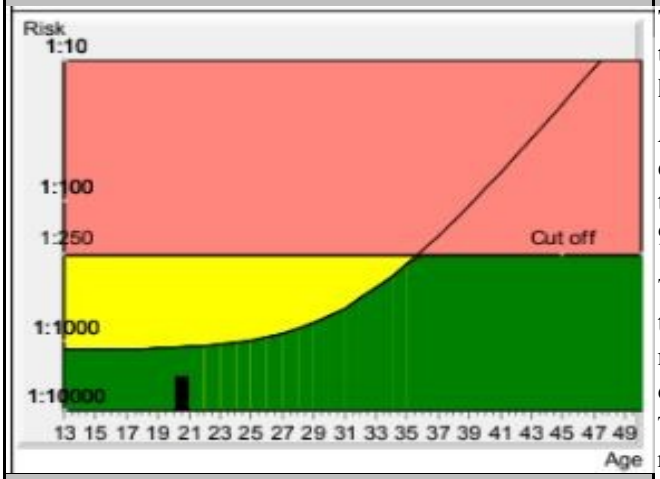
Risks at sampling date

Age Risk	1:1129
Biochemical T21 Risk	1:2425
Combined Trisomy 21 Risk	<1:10000
Trisomy 13/18 + NT	<1:10000

Ultrasound Data

Gestational age	13+6
Method	CRL (<>Hadlock)
Scan Date	27-01-19
CRL	77.4
Nuchal translucency MoM	1.00
Nasal Bone	Present
Sonographer	DR.S.MAHATO
Qualifications in measuring NT	MD

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 10000 women with the same data, there is one woman with a trisomy 21 pregnancy and 9999 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

The laboratory can not be held responsible for their impact on the risk assessment! Calculated value has no diagnostic value!

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