MOLQ Express Lab

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Date of Report	10-01-19
PRISCA	5.0.2.37

Patient Data			
Name	**Mrs Chagni**	Patient ID	011901090096
Birthday	01-01-89	Sample ID	10420387
Age at delivery	30.4	Sample Date	09/01/19
Gestational age	19+3		

Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	66	Diabetes	no	Pregnancies	
Smoker	no	Origin	Asian		

Biochemical Data

Parameter	Value		Corr Mom
AFP	49.02	ng/ml	0.82
uE3	1.85	mIU/ml	1.38
hCG	9410.5	ng/ml	0.55

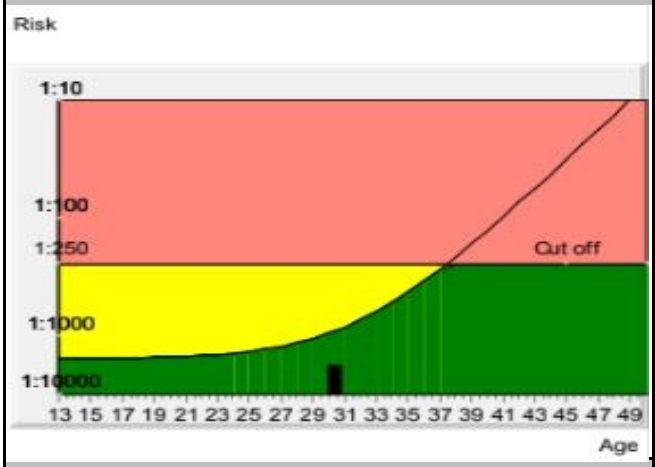
Risks at sampling date

Age Risk	1:919
Biochemical T21 Risk	1:6056
Combined T21 Risk	<1:10000
Trisomy 18	<1:10000

Ultrasound Data

Gestational age	12+2
Method	CRL (<>Hadlock)
Scan Date	20-11-19
CRL measurments	57.4 mm
NT Translucency	0.66
NT Mom	1.00
Nasal Bone	Present
Sonographer	DR.RITU JAIN
Qualifications in measuring NT	

Risk



Trisomy 13/18 + NT

The calculated risk for trisomy 18 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 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 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