MOLQ Express Lab
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Date of Report	10-01-19
PRISCA	5.0.2.37

Patient Data			
Name	**Mrs Madhuri Kumari**	Patient ID	011901090097
Birthday	08-04-95	Sample ID	10420386
Age at delivery	24.2	Sample Date	09/01/19
Gestational age	16+2		

Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	51	Diabetes	unknown	Pregnancies	
Smoker	unknown	Origin	Asian		

Biochemical Data

Parameter	Value		Corr Mom
AFP	39.71	ng/ml	0.88
uE3	1.02	mIU/ml	1.47
hCG	32996.5	ng/ml	1.01

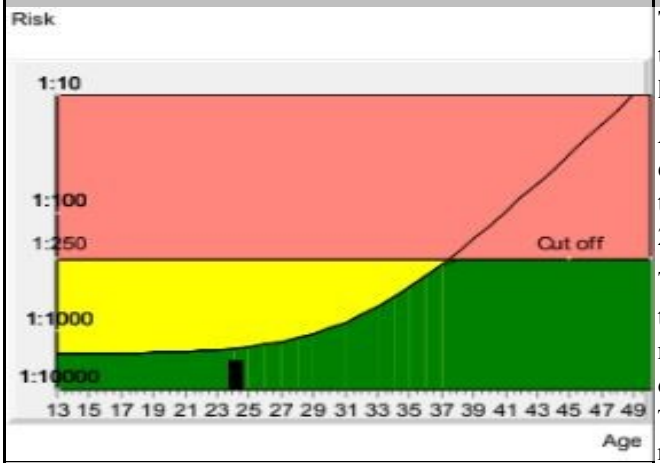
Risks at sampling date

Age Risk	1:1417
Biochemical T21 Risk	1:4472
Combined T21 Risk	1:2327
Trisomy 18	<1:10000

Ultrasound Data

Gestational age	12+4
Method	CRL (<>Hadlock)
Scan Date	14-12-18
NT Translucency	2.60 mm
NT Mom	1.57
Nasal Bone	Present
Sonographer	DR. A. JUNEJA
Qualifications in measuring NT	MD

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 2327 women with the same data, there is one woman with a trisomy 21 pregnancy and 2326 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