MOLQ Express Lab

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Date of Report	11/6/2019
PRISCA	5.0.2.37

Patient Data

Name	**Mrs Monika**	Patient ID	011906090279
Birthday	7/11/1994	Sample ID	10303232
Age at delivery	24.6	Sample Date	09/06/19
Gestational age	11+3		

Correction factors

Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	39	Diabetes	no	Pregnancies	unknown
Smoker	no	Origin	Asian		

Biochemical Data

Parameter	Value	Corr Mom
PAPP-A	3.41 mIU/ml	0.72
fb-hCG	77.56 ng/ml	1.25

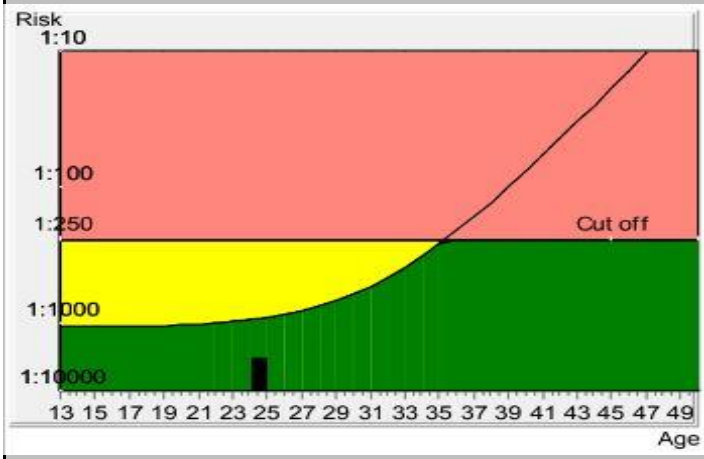
Risks at sampling date

Age Risk	1:939
Biochemical Trisomy 21 Risk	1:1689
Combined Trisomy 21 Risk	1:6452
Trisomy 13/18 + NT	<1:10000

Ultrasound Data

Gestational age	11+2
Method	CRL (<>Robinson)
Scan Date	9/6/2019
Crown Rump Length (mm)	45.2
Nuchal translucency MoM	1.04
Nasal Bone	present
Sonographer	DR.RAJESH ARORA
Qualification in measuring NT	HMC

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)

TThe 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 6452 women with the same data, there is one woman with a trisomy 21 pregnancy and 6451 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 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

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