MOLQ Express Lab
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Date of Report	22-10-18
PRISCA	5.0.2.37

Patient Data			
Name	**Mrs Aradhana**	Patient ID	011810210065
Birthday	14-05-91	Sample ID	10362535
Age at delivery	27.4	Sample Date	21-10-18
Gestational age	12+0		

Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	61	Diabetes	unknown	Pregnancies	unknown
Smoker	Unknown	Origin	Asian		

Biochemical Data

Parameter	Value		Corr Mom
PAPP-A	2.36	mIU/ml	0.77
fb-hCG	43.6	ng/ml	0.92

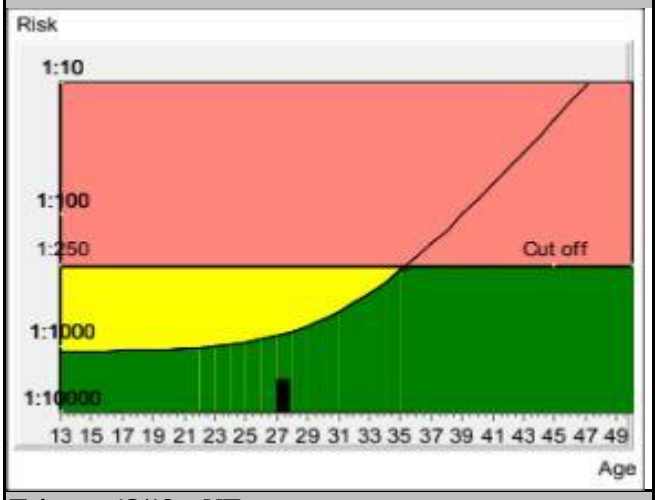
Risks at sampling date

Age Risk	1:816
Biochemical T21 Risk	1:3365
Combined Trisomy 21 Risk	1:600
Trisomy 13/18	<1:10000

Ultrasound Data

Gestational age	10+6
Method	CRL (<>Robinson)
Scan Date	13-10-18
Crown rump length (mm)	40.2
Nuchal translucency MoM	
Nasal Bone	
Sonographer	Dr. Sanjeev Kumar
Qualifications in measuring NT	MBBS,PGDUS, DMRD

Risk



Trisomy 13/18 + NT

The calculated risk for Trisomy 13/18 is <1:10000, which indicates a low risk

Down's Syndrome Risk (Trisomy 21 Screening)

The calculated risk for Trisomy 21 Is below the cut off, which represents a low risk.

After the result of the Trisomy 21 test (with NT) it is expected that among 3365 women with the same data, there is one women with a trisomy 21 pregnancy and 3364 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 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